

State of Iowa

Iowa
Administrative
Code
Supplement

Biweekly
March 22, 2023



Published by the
STATE OF IOWA
UNDER AUTHORITY OF IOWA CODE SECTION 17A.6

The Iowa Administrative Code (IAC) Supplement is published biweekly pursuant to Iowa Code sections 2B.5A and 17A.6. The Supplement is a compilation of updated Iowa Administrative Code chapters that reflect rule changes which have been adopted by agencies and filed with the Administrative Rules Coordinator as provided in Iowa Code sections 7.17, 17A.4, and 17A.5 and published in the Iowa Administrative Bulletin bearing the same publication date as the one for this Supplement. To determine the specific changes to the rules, refer to the Iowa Administrative Bulletin. To maintain a loose-leaf set of the IAC, insert the chapters according to the instructions included in the Supplement.

In addition to the rule changes adopted by agencies, the chapters may reflect objection to a rule or a portion of a rule filed by the Administrative Rules Review Committee (ARRC), the Governor, or the Attorney General pursuant to Iowa Code section 17A.4(6); an effective date delay or suspension imposed by the ARRC pursuant to section 17A.8(9) or 17A.8(10); rescission of a rule by the Governor pursuant to section 17A.4(8); nullification of a rule by the General Assembly pursuant to Article III, section 40, of the Constitution of the State of Iowa; other action relating to rules enacted by the General Assembly; updated chapters for the Uniform Rules on Agency Procedure; or an editorial change to a rule by the Administrative Code Editor pursuant to Iowa Code section 2B.13(2).

INSTRUCTIONS

FOR UPDATING THE

IOWA ADMINISTRATIVE CODE

Agency names and numbers in bold below correspond to the divider tabs in the IAC binders. New and replacement chapters included in this Supplement are listed below. Carefully remove and insert chapters accordingly.

Editor's telephone 515.281.3355 or 515.242.6873

Agriculture and Land Stewardship Department[21]

Replace Analysis
Replace Chapter 93

Soil Conservation and Water Quality Division[27]

Replace Chapter 2

Iowa Finance Authority[265]

Replace Analysis
Replace Chapter 31
Insert Chapter 47

Professional Licensure Division[645]

Replace Chapter 60
Replace Chapter 64

Pharmacy Board[657]

Replace Analysis
Replace Chapter 3
Replace Chapter 6
Replace Chapter 8
Replace Chapter 18
Replace Chapter 21
Replace Chapter 39

Revenue Department[701]

Replace Chapters 102 and 103
Replace Chapter 111

AGRICULTURE AND LAND STEWARDSHIP DEPARTMENT[21]

[Created by 1986 Iowa Acts, chapter 1245]
[Prior to 7/27/88, Agriculture Department[30]]
Rules under this Department “umbrella” also include
Agricultural Development Authority[25] and Soil Conservation Division[27]

CHAPTER 1 ADMINISTRATION

- 1.1(159) Organization
- 1.2(159) Consumer protection and industry services division
- 1.3(159) Administration division
- 1.4(159) Soil conservation division
- 1.5(159) Food safety and animal health

CHAPTER 2 CONTESTED CASE PROCEEDINGS AND PRACTICE (Uniform Rules)

- 2.1(17A,159) Scope and applicability
- 2.2(17A,159) Definitions
- 2.3(17A,159) Time requirements
- 2.4(17A,159) Requests for contested case proceeding
- 2.5(17A,159) Notice of hearing
- 2.6(17A,159) Presiding officer
- 2.12(17A,159) Service and filing of pleadings and other papers
- 2.15(17A,159) Motions
- 2.16(17A,159) Prehearing conference
- 2.17(17A,159) Continuances
- 2.22(17A,159) Default
- 2.23(17A,159) Ex parte communication
- 2.24(17A,159) Recording costs
- 2.25(17A,159) Interlocutory appeals
- 2.26(17A,159) Final decision
- 2.27(17A,159) Appeals and review
- 2.28(17A,159) Applications for rehearing
- 2.29(17A,159) Stays of agency action

CHAPTER 3 PETITIONS FOR RULE MAKING (Uniform Rules)

- 3.1(17A) Petition for rule making
- 3.3(17A) Inquiries
- 3.5(17A) Petitions for related entities

CHAPTER 4 DECLARATORY ORDERS (Uniform Rules)

- 4.1(17A,159) Petition for declaratory order
- 4.2(17A,159) Notice of petition
- 4.3(17A,159) Intervention
- 4.4(17A,159) Briefs
- 4.5(17A,159) Inquiries
- 4.6(17A,159) Service and filing of petitions and other papers
- 4.7(17A,159) Consideration

- 4.8(17A,159) Action on petition
- 4.9(17A,159) Refusal to issue order
- 4.12(17A,159) Effect of a declaratory order

CHAPTER 5
AGENCY PROCEDURE FOR RULE MAKING
(Uniform Rules)

- 5.1(17A,159) Applicability
- 5.3(17A,159) Public rule-making docket
- 5.4(17A,159) Notice of proposed rule making
- 5.5(17A,159) Public participation
- 5.6(17A,159) Regulatory analysis
- 5.10(17A,159) Exemptions from public rule-making procedures
- 5.11(17A,159) Concise statement of reasons
- 5.13(17A,159) Agency rule-making record

CHAPTER 6
PUBLIC RECORDS AND FAIR INFORMATION PRACTICES
(Uniform Rules)

- 6.1(17A,22) Definitions
- 6.3(17A,22) Requests for access to records
- 6.6(17A,22) Procedure by which additions, dissents, or objections may be entered into certain records
- 6.9(17A,22) Disclosures without the consent of the subject
- 6.10(17A,22) Routine use
- 6.11(17A,22) Consensual disclosure of confidential records
- 6.12(17A,22) Release to subject
- 6.13(17A,22) Availability of records
- 6.14(17A,22) Personally identifiable information
- 6.15(17A,22) Other groups of records
- 6.16(17A,22) Data processing systems
- 6.17(159,252J,272D) Release of confidential licensing information for collection purposes

CHAPTER 7
COLLECTION PROCEDURES

- 7.1(159,252J,272D) Licensing actions
- 7.2(159,252J,272D) Collection procedures

CHAPTER 8
WAIVER OF RULES

- 8.1(17A,159) Definition
- 8.2(17A,159) Scope of chapter
- 8.3(17A,159) Applicability
- 8.4(17A,159) Criteria for waiver
- 8.5(17A,159) Filing of petition
- 8.6(17A,159) Content of petition
- 8.7(17A,159) Additional information
- 8.8(17A,159) Notice
- 8.9(17A,159) Hearing procedures
- 8.10(17A,159) Ruling
- 8.11(17A,159) Public availability
- 8.12(17A,159) Submission of waiver information
- 8.13(17A,159) Cancellation of a waiver
- 8.14(17A,159) Violations

- 8.15(17A,159) Defense
- 8.16(17A,159) Judicial review

CHAPTERS 9 to 11

Reserved

CHAPTER 12

RENEWABLE FUELS AND COPRODUCTS

- 12.1(159A) Purpose
- 12.2(159A) Definitions
- 12.3(159A) General provisions
- 12.4(159A) Renewable fuels motor vehicle fuels decals

CHAPTER 13

RENEWABLE FUEL INFRASTRUCTURE BOARD—ORGANIZATION

- 13.1(159A) Definitions
- 13.2(159A) Renewable fuel infrastructure board

CHAPTER 14

RENEWABLE FUEL INFRASTRUCTURE PROGRAM FOR
RETAIL MOTOR FUEL SITES

- 14.1(159A) Purpose
- 14.2(159A) Eligible applicants

CHAPTER 15

RENEWABLE FUEL INFRASTRUCTURE PROGRAM FOR
BIODIESEL TERMINAL GRANTS

- 15.1(159A) Purpose
- 15.2(159A) Eligible applicants

CHAPTER 16

RENEWABLE FUEL INFRASTRUCTURE PROGRAM ADMINISTRATION

- 16.1(159A) Form of award available; award amount
- 16.2(159A) Application process
- 16.3(159A) Review process
- 16.4(159A) Contract administration

CHAPTERS 17 to 19

Reserved

CHAPTER 20

REFERENDUM

- 20.1(159) Purpose
- 20.2(159) Definitions
- 20.3(159) Voter eligibility
- 20.4(159) Referendum methods and procedures
- 20.5(159) Contesting referendum results
- 20.6(159) Official certification

CHAPTER 21

Reserved

CHAPTER 22

APIARY

22.1(160)	Diseases
22.2(160)	Parasites
22.3(160)	Requirement for the sale of bees
22.4(160)	Certificate of inspection required
22.5(160)	Certificate of inspection expiration
22.6(160)	American Foulbrood treatment
22.7(160)	Varroa mite treatment
22.8(160)	Undesirable subspecies of honeybees
22.9(160)	European honeybee certification
22.10(160)	Prohibit movement of bees from designated states
22.11(160)	Inspection required for the sale of bees, comb, or used equipment

CHAPTERS 23 to 35

Reserved

CHAPTER 36

EGG HANDLERS

36.1(196)	Definitions
36.2(196)	Licensing
36.3(196)	Minimum sanitation and operating requirements
36.4(196)	Egg grading or candling area
36.5(196)	Water supply
36.6(196)	Egg storage
36.7(196)	Eggs used in food preparation
36.8(196)	Labeling and packaging
36.9(196)	Restricted eggs
36.10(196)	Inspections and records
36.11(196)	Enforcement
36.12(196)	Health and hygiene of personnel
36.13(196)	Iowa grades

CHAPTERS 37 to 39

Reserved

CHAPTER 40

AGRICULTURAL SEEDS

40.1(199)	Agricultural seeds
40.2(199)	Seed testing
40.3(199)	Labeling
40.4 and 40.5	Reserved
40.6(199)	Classes and sources of certified seed
40.7(199)	Labeling of seeds with secondary noxious weeds
40.8(199)	Germination standards for vegetable seeds
40.9(199)	White sweet clover
40.10(199)	Labeling of conditioned seed distributed to wholesalers
40.11(199)	Seeds for sprouting
40.12(199)	Relabeling
40.13(199)	Hermetically sealed seed
40.14(199)	Certification of seed and potatoes
40.15(199)	Federal regulations adopted
40.16(199)	Seed libraries

CHAPTER 41
COMMERCIAL FEED

41.1(198)	Definitions and terms
41.2(198)	Label format
41.3(198)	Label information
41.4(198)	Expression of guarantees
41.5(198)	Suitability
41.6(198)	Ingredients
41.7(198)	Directions for use and precautionary statements
41.8(198)	Nonprotein nitrogen
41.9(198)	Drug and feed additives
41.10(198)	Adulterants
41.11(198)	Good manufacturing practices
41.12(198)	Cottonseed product control

CHAPTER 42
PET FOOD

42.1(198)	Definitions and terms
42.2(198)	Label format and labeling
42.3(198)	Brand and product names
42.4(198)	Expression of guarantees
42.5(198)	Ingredients
42.6(198)	Drugs and pet food additives
42.7(198)	Statements of calorie content
42.8(198)	Descriptive terms

CHAPTER 43
FERTILIZERS AND AGRICULTURAL LIME

43.1(200)	Additional plant food elements besides N, P and K
43.2(200)	Warning required
43.3(200)	Specialty fertilizer labels
43.4(200)	Pesticides in fertilizers
43.5(200)	Cancellation or suspension of registration or license
43.6(200)	Standard for the storage and handling of anhydrous ammonia
43.7(200)	Groundwater protection fee
43.8 to 43.19	Reserved
43.20(201)	Agricultural lime
43.21(200)	Minimum requirements for registration of fertilizer and soil conditioners
43.22(200)	Provisional product registration
43.23(200)	Review of product registrations
43.24(200)	Product claims
43.25 to 43.29	Reserved
43.30(201A)	Definitions
43.31(201A)	Determination of ECCE
43.32(201A)	Sample procedure
43.33(201A)	Sample analysis
43.34(201A)	Sample fee
43.35(201A)	Certification
43.36(201A)	Compliance with certification
43.37(201A)	Labeling
43.38(201A)	Toxic materials prohibited

- 43.39(201A) Added materials
- 43.40(201A) Egg shells

CHAPTER 44
ON-SITE CONTAINMENT
OF PESTICIDES, FERTILIZERS AND SOIL CONDITIONERS

PESTICIDES

- 44.1(206) Definitions
- 44.2(206) On-site containment of pesticides
- 44.3(206) Design plans and specifications
- 44.4(206) Certification of construction
- 44.5(206) New pesticide storage and mixing site location
- 44.6(206) Pesticide storage and mixing site
- 44.7(206) Secondary containment for nonmobile bulk pesticide storage and mixing
- 44.8(206) Pesticide storage and mixing site containers
- 44.9(206) Transportation of bulk pesticides
- 44.10(206) Mixing, repackaging and transfer of pesticides
- 44.11(206) Distribution of bulk pesticides
- 44.12(206) Secondary containment for aerial applicator aircraft
- 44.13 to 44.49 Reserved

FERTILIZERS AND SOIL CONDITIONERS

- 44.50(200) On-site containment of fertilizers and soil conditioners
- 44.51(200) Definitions
- 44.52(200) Design plans and specifications
- 44.53(200) New fertilizer or soil conditioner storage site location
- 44.54(200) Certification of construction
- 44.55(200) Secondary containment for liquid fertilizers and liquid soil conditioner storage
- 44.56(200) Secondary containment for nonliquid fertilizers and soil conditioners
- 44.57(200) Fertilizer loading, unloading, and mixing area
- 44.58(200) Wash water and rinsates

CHAPTER 45
PESTICIDES

DIVISION I

- 45.1(206) Definitions and standards
- 45.2(206) Methods of analysis
- 45.3(206) Registration required
- 45.4(206) Registration of products
- 45.5(206) Registration, general application of
- 45.6(206) Revocation, suspension or denial of registration
- 45.7(206) Changes in labeling or ingredient statement
- 45.8(206) Label requirements
- 45.9(206) Directions for use—when necessary
- 45.10(206) Other claims
- 45.11(206) Name of product
- 45.12(206) Brand names, duplication of, or infringement on
- 45.13(206) Ingredient statement
- 45.14(206) Net contents
- 45.15(206) Coloration of highly toxic materials
- 45.16(206) Illegal acts
- 45.17(206) Guarantee of pesticide

45.18(206)	Shipments for experimental use
45.19(206)	Enforcement
45.20(206)	Hazardous rodenticides
45.21(206)	Highly toxic
45.22(206)	License and certification standards for pesticide applicators
45.23(206)	Sale or possession of thallium
45.24(206)	Warning, caution and antidote statements
45.25(206)	Declaration of pests
45.26(206)	Record-keeping requirements
45.27(206)	Use of high volatile esters
45.28(206)	Emergency single purchase/single use of restricted pesticide
45.29(206)	Application of general use pesticide by nonlicensed commercial applicator
45.30(206)	Restricted use pesticides classified
45.31(206)	Application of pesticides toxic to bees
45.32(206)	Use of DDT and DDD
45.33(206)	Use of inorganic arsenic
45.34(206)	Use of heptachlor
45.35(206)	Use of lindane
45.36(206)	Reports of livestock poisoning
45.37(206)	Approval of use of inorganic arsenic formulation
45.38 to 45.44	Reserved
45.45(206)	Ethylene dibromide (EDB) residue levels in food
45.46(206)	Use of pesticide Command 6EC
45.47(206)	Reporting of pesticide sales
45.48(206)	Dealer license fees
45.49(206)	Pesticide use recommendations
45.50(206)	Notification requirements for urban pesticide applications
45.51(206)	Restrictions on the distribution and use of pesticides containing the active ingredient atrazine or any combination of active ingredients including atrazine
45.52(206)	Continuing instructional courses for pesticide applicator recertification

DIVISION II

45.53 to 45.99	Reserved
----------------	----------

DIVISION III
CIVIL PENALTIES

45.100(206)	Definitions
45.101(206)	Commercial pesticide applicator peer review panel
45.102(206)	Civil penalties—establishment, assessment, and collection
45.103(206)	Review period
45.104(206)	Review by peer review panel
45.105(206)	Response by peer review panel

CHAPTER 46
CROP PESTS

46.1(177A)	Nursery stock
46.2(177A)	Hardy
46.3(177A)	Person
46.4(177A)	Nursery growers
46.5(177A)	Nursery
46.6(177A)	Nursery dealer
46.7(177A)	Out-of-state nursery growers and nursery dealers
46.8(177A)	Nursery inspection
46.9(177A)	Nursery dealer certificate

46.10(177A)	Proper facilities
46.11(177A)	Storage and display
46.12(177A)	Nursery stock viability qualifications
46.13(177A)	Certificates
46.14(177A)	Miscellaneous and service inspections
46.15(177A)	Insect pests and diseases
46.16(177A)	Firewood labeling

CHAPTER 47

IOWA ORGANIC PROGRAM

47.1(190C)	Iowa organic program
47.2	Reserved
47.3(190C)	Drift
47.4	Reserved
47.5(190C)	Recognition
47.6(190C)	General requirements
47.7	Reserved
47.8(190C)	Certification agent

ADMINISTRATIVE

47.9(190C)	Fees
47.10(190C)	Compliance

CHAPTER 48

PESTICIDE ADVISORY COMMITTEE

48.1(206)	Function
48.2(206)	Staff
48.3(206)	Advisors
48.4(206)	Meetings
48.5(206)	Open records
48.6(206)	Budget
48.7(206)	Review of pesticide applicator instructional course and examination

CHAPTER 49

BULK DRY ANIMAL NUTRIENTS

49.1(200A)	Definitions
49.2(200A)	License
49.3(200A)	Registration
49.4(200A)	Additional plant elements
49.5(200A)	Distribution statement
49.6(200A)	Distribution reports
49.7(200A)	Storage of bulk dry animal nutrients
49.8(200A)	Manure management plans

CHAPTER 50

WOMEN, INFANTS, AND CHILDREN/FARMERS' MARKET NUTRITION PROGRAM
AND SENIOR FARMERS' MARKET NUTRITION PROGRAM

50.1(159,175B)	Authority and scope
50.2(159,175B)	Severability
50.3(159,175B)	Definitions
50.4(159,175B)	Program description and goals
50.5(159,175B)	Administration and agreements
50.6(159,175B)	Distribution of benefits
50.7(159,175B)	Recipient responsibilities

- 50.8(159,175B) Farmers' market, farmstand, and community supported agriculture (CSA) authorization and priority
- 50.9(159,175B) Vendor certification
- 50.10(159,175B) Certified vendor obligations
- 50.11(159,175B) Certified vendor noncompliance sanctions
- 50.12(159,175B) Appeal
- 50.13(159,175B) Deadlines
- 50.14(159,175B) Discrimination complaints

CHAPTER 51
FARM-TO-SCHOOL FUND

- 51.1(190A) Purpose
- 51.2(190A) Definitions
- 51.3(190A) Application to participate
- 51.4(190A) Eligible purchases
- 51.5(190A) Reimbursement for purchases

CHAPTER 52
MARKETING

CHOOSE IOWA PROMOTIONAL PROGRAM

- 52.1(159) Definitions
- 52.2(159) Product qualification
- 52.3(159) Application for membership
- 52.4(159) Fees
- 52.5(159) Approval for use of logo
- 52.6(159) Self-certification
- 52.7(159) Compliance
- 52.8(159) Violations
- 52.9 to 52.19 Reserved

VALUE-ADDED AGRICULTURE GRANT PROGRAM

- 52.20(159) Definitions
- 52.21(159) Eligibility
- 52.22(159) Application and review process
- 52.23(159) Scoring criteria
- 52.24(159) Disbursement of funds

CHAPTERS 53 to 57
Reserved

CHAPTER 58
NOXIOUS WEEDS

- 58.1(317) Definitions
- 58.2(317) Purple loosestrife
- 58.3(317) Records
- 58.4(317) Noxious weed lists

CHAPTER 59
Reserved

CHAPTER 60

POULTRY

- 60.1(168) Egg-type chickens, meat-type chickens, turkeys, domestic waterfowl, domestic game birds and exhibition poultry
- 60.2(168) License for dealers of baby chicks or domestic fowls
- 60.3(163) Turkeys
- 60.4(163) Registration of exhibitions involving poultry

CHAPTER 61

DEAD ANIMAL DISPOSAL

- 61.1(167) Dead animal disposal—license
- 61.2(167) Animal disposal—persons defined
- 61.3(167) Disposing of dead animals by cooking
- 61.4(167) License fee
- 61.5(167) Certificate issuance
- 61.6(167) Filing certificate
- 61.7(167) License renewal
- 61.8 to 61.10 Reserved
- 61.11(167) Disposal plant plans
- 61.12(167) Disposal plant specifications
- 61.13 and 61.14 Reserved
- 61.15(167) Conveyances requirements
- 61.16(167) Disposal plant trucks
- 61.17(167) Disposal employees
- 61.18(167) Tarpaulins
- 61.19(167) Disposal vehicles—disinfection
- 61.20 to 61.22 Reserved
- 61.23(167) Rendering plant committee
- 61.24(167) Rendering plant—spraying
- 61.25(167) Penalty
- 61.26 and 61.27 Reserved
- 61.28(167) Anthrax
- 61.29(167) Anthrax—disposal
- 61.30(167) Classical swine fever—carcasses
- 61.31(167) Noncommunicable diseases—carcasses
- 61.32(167) Carcass disposal—streams
- 61.33(167) Improper disposal

CHAPTER 62

REGISTRATION OF IOWA-FOALED
HORSES AND IOWA-WHELPED DOGS

- 62.1(99D) Definitions
- 62.2(99D) Iowa horse and dog breeders' fund and Iowa thoroughbred horse breeders' promotion fund
- 62.3(99D) Forms
- 62.4(99D) Disciplinary actions
- 62.5(99D) Access to premises and records
- 62.6(99D) Registration fees
- 62.7 to 62.9 Reserved

THOROUGHBRED DIVISION

- 62.10(99D) Iowa thoroughbred stallion requirements
- 62.11(99D) Notification requirements

62.12(99D)	Stallion qualification and application procedure
62.13(99D)	Application information
62.14(99D)	Breeding record—report of mares bred
62.15(99D)	Iowa-foaled horses and brood mares
62.16(99D)	Iowa-foaled horse status
62.17 to 62.19	Reserved

STANDARD BRED DIVISION

62.20(99D)	Iowa standardbred stallion requirements
62.21(99D)	Notification requirements
62.22(99D)	Stallion qualification and application procedure
62.23(99D)	Application information
62.24(99D)	Breeding record—report of mares bred
62.25(99D)	Iowa-foaled horses and brood mares
62.26(99D)	Iowa-foaled horse status
62.27 to 62.29	Reserved

QUARTER HORSE DIVISION

62.30(99D)	Iowa quarter horse stallion requirements
62.31(99D)	Notification requirements
62.32(99D)	Stallion qualification and application procedure
62.33(99D)	Application information
62.34(99D)	Breeding record—report of mares bred
62.35(99D)	Iowa-foaled horses and brood mares
62.36(99D)	Iowa-foaled horse status
62.37(99D)	Embryo transfer for Iowa-foaled status
62.38 and 62.39	Reserved

GREYHOUND DOG DIVISION

62.40(99D)	Iowa-whelped dog requirements
62.41(99D)	Procedures for registration

CHAPTER 63

BRANDING

63.1(169A)	Location of brands on livestock
63.2(169A)	Brands in conflict

CHAPTER 64

INFECTIOUS AND CONTAGIOUS DISEASES

64.1(163)	Reporting disease
64.2(163)	Disease prevention and suppression
64.3(163)	Duties of township trustees and health board
64.4(163)	“Exposed” defined
64.5(163)	Sale of vaccine
64.6(163)	“Quarantine” defined
64.7(163)	Chiefs of Iowa and U.S. animal industries to cooperate
64.8(163)	Animal blood sample collection
64.9	Reserved

GLANDERS AND FARCY CONTROL

64.10(163)	Preventing spread of glanders
64.11(163)	Disposal of diseased animal
64.12(163)	Glanders quarantine
64.13(163)	Tests for glanders and farcy
64.14	Reserved

BLACKLEG CONTROL

64.15(163) Blackleg
64.16 Reserved

DEPARTMENT NOTIFICATION OF DISEASES

64.17(163) Notification of chief of animal industry
64.18 to 64.22 Reserved

RABIES CONTROL

64.23(163) Rabies—exposed animals
64.24(163) Rabies quarantine
64.25(351) Control and prevention of rabies
64.26 to 64.29 Reserved

SCABIES OR MANGE CONTROL

64.30(163) Scabies or mange quarantine
64.31 Reserved

DISEASE CONTROL AT FAIRS AND EXHIBITS

64.32(163) State fairgrounds—disinfection of livestock quarters
64.33(163) County fairs—disinfection of livestock quarters
64.34(163) Health requirements for exhibition of livestock, poultry and birds at the state fair, district shows and exhibitions
64.35(163) Health requirements for exhibition of livestock, poultry and birds at exhibitions
64.36 and 64.37 Reserved

DISEASE CONTROL BY CONVEYANCES

64.38(163) Transportation companies—disinfecting livestock quarters
64.39(163) Livestock vehicles—disinfection
64.40 Reserved

INTRASTATE MOVEMENT OF LIVESTOCK

64.41(163) General
64.42(163) Veterinary inspection
64.43(163) Swine
64.44 to 64.46 Reserved

BRUCELLOSIS

64.47(163) Definitions as used in these rules
64.48 Reserved
64.49(163) Certified brucellosis-free herd
64.50(163) Restraining animals
64.51(163) Quarantines
64.52(163) Identification of bovine animals
64.53(163) Cleaning and disinfection
64.54(163) Disposal of reactors
64.55(163) Brucellosis tests and reports
64.56(163) Suspect animals designated as reactors
64.57(163) Indemnity not allowed
64.58(163) Area testing
64.59 to 64.62 Reserved

BOVINE BRUCELLOSIS

64.63(164) Back tagging in bovine brucellosis control
64.64(164) Fee schedule
64.65(163) Definitions
64.66 Reserved

ERADICATION OF SWINE BRUCELLOSIS

64.67(163A)	Brucellosis test
64.68(163A)	Veterinarians to test
64.69 and 64.70	Reserved
64.71(163A)	Fee schedule
64.72	Reserved

ERADICATION OF BOVINE TUBERCULOSIS

64.73(163)	Tuberculin tests classified
64.74(163)	Acceptance of intradermic test
64.75(163)	Adoption of intradermic test
64.76(163)	Ophthalmic test
64.77(163)	Tuberculin test deadline
64.78(163)	Health certificate
64.79(163)	Ear tags
64.80(163)	Cattle importation
64.81(163)	Tuberculin reactors
64.82(163)	Steers—testing
64.83(163)	Female cattle—testing
64.84(163)	Certificates and test charts
64.85(163)	Slaughtering reactors
64.86(163)	Agriculture tuberculin rules
64.87(163)	“Tuberculosis-free accredited herd” defined
64.88(163)	Retesting
64.89(163)	Accredited herd
64.90(163)	Selection of cattle for tuberculin tests
64.91(163)	Identification for test
64.92(163)	Removing cattle from herd
64.93(163)	Milk
64.94(163)	Sanitary measures
64.95(163)	Interstate shipment
64.96(163)	Reactors—removal
64.97(163)	Certificate
64.98(163)	Violation of certificate
64.99(163)	Tuberculin—administration
64.100(163)	Sale of tuberculin
64.101(165)	Fee schedule
64.102 and 64.103	Reserved

CHRONIC WASTING DISEASE (CWD)

64.104(163)	Definitions
64.105(163)	Supervision of the cervid CWD surveillance identification program
64.106(163)	Surveillance procedures
64.107(163)	Official cervid tests
64.108(163)	Investigation of CWD affected animals identified through surveillance
64.109(163)	Duration of quarantine
64.110(163)	Herd plan
64.111(163)	Identification and disposal requirements
64.112(163)	Cleaning and disinfecting
64.113(163)	Methods for obtaining certified CWD cervid herd status
64.114(163)	Recertification of CWD cervid herds
64.115(163)	Movement into a certified CWD cervid herd
64.116(163)	Movement into a monitored CWD cervid herd

- 64.117(163) Recognition of monitored CWD cervid herds
- 64.118(163) Recognition of certified CWD cervid herds
- 64.119 to 64.132 Reserved

ERADICATION OF SWINE TUBERCULOSIS

- 64.133(159) Indemnity
- 64.134(159) Fee schedule
- 64.135 to 64.146 Reserved

PSEUDORABIES DISEASE

- 64.147(163,166D) Definitions. As used in these rules:
- 64.148 to 64.150 Reserved
- 64.151(163,166D) Quarantines
- 64.152(163,166D) Nondifferentiable pseudorabies vaccine disapproved
- 64.153(166D) Pseudorabies disease program areas
- 64.154(163,166D) Identification
- 64.155(163,166D,172B) Certificates of inspection
- 64.156(166D) Noninfected herds
- 64.157(166D) Herd cleanup plan for infected herds (eradication plan)
- 64.158(166D) Feeder pig cooperator plan for infected herds
- 64.159(166D) Herds of unknown status
- 64.160(166D) Approved premises
- 64.161(166D) Sales to approved premises
- 64.162(166D) Certification of veterinarians to initiate approved herd cleanup plans and approved feeder pig cooperator plan agreements and fee basis
- 64.163(166D) Nondifferentiable pseudorabies vaccine disapproved
- 64.164 to 64.169 Reserved

PARATUBERCULOSIS (JOHNE'S) DISEASE

- 64.170(165A) Definitions
- 64.171(165A) Supervision of the Johne's disease program
- 64.172(165A) Official Johne's disease tests
- 64.173(165A) Vaccination allowed
- 64.174(165A) Herd plan
- 64.175(165A) Identification and disposal requirements
- 64.176(165A) Segregation, cleaning, and disinfecting
- 64.177(165A) Intrastate movement requirements
- 64.178(165A) Import requirements
- 64.179 to 64.184 Reserved

LOW PATHOGENIC AVIAN INFLUENZA (LPAI)

- 64.185(163) Definitions
- 64.186(163) Supervision of the low pathogenic avian influenza program
- 64.187(163) Surveillance procedures
- 64.188(163) Official LPAI tests
- 64.189(163) Investigation of LPAI affected poultry identified through surveillance
- 64.190(163) Duration of quarantine
- 64.191(163) Flock plan
- 64.192(163) Cleaning and disinfecting
- 64.193 to 64.199 Reserved

SCRAPIE DISEASE

- 64.200(163) Definitions
- 64.201(163) Supervision of the scrapie eradication program
- 64.202(163) Identification

64.203(163)	Restrictions on the removal of official identification
64.204(163)	Records
64.205(163)	Responsibility of persons handling animals in commerce to ensure the official identification of animals
64.206(163)	Veterinarian's responsibilities when identifying sheep or goats
64.207(163)	Flock plans
64.208(163)	Certificates of Veterinary Inspection
64.209(163)	Requirements for shows and sales
64.210(163)	Movement restrictions for animals and flocks
64.211(163)	Approved terminal feedlots

CHAPTER 65

ANIMAL AND LIVESTOCK IMPORTATION

65.1(163)	Definitions
65.2(163)	Pre-entry permits
65.3(163)	General requirements and limitations
65.4(163)	Cattle and bison
65.5(163,166D)	Swine
65.6(163)	Goats
65.7(163)	Sheep
65.8(163)	Equine
65.9(163)	Cervidae
65.10(163)	Dogs and cats
65.11(163)	Poultry, domestic fowl, and hatching eggs
65.12(163)	Swine production health plan (SPHP)
65.13(163)	Penalties

CHAPTER 66

LIVESTOCK MOVEMENT

66.1(163)	Definitions and permits
66.2(163)	Animal health sanitation and record-keeping requirements
66.3(163)	Duties and responsibilities of the livestock market management
66.4(163)	Duties and responsibilities of the livestock market veterinary inspector
66.5(163)	Classification of livestock markets and permit holders
66.6(163)	Requirements for state-federal (specifically) approved markets
66.7(163)	Requirements for sale of all bovine animals
66.8(163)	Testing
66.9(163)	Order of sale through auction markets
66.10(163)	Releasing cattle
66.11(163,172B)	Movement of livestock within the state
66.12(189,189A)	Movement of food-producing animals and their products into the state
66.13(163,202C)	Feeder pig dealer bonding/letter of credit requirement and claims procedures
66.14(163)	Intrastate movement requirements
66.15 to 66.19	Reserved
66.20(163)	Revocation or denial of permit

CHAPTER 67

ANIMAL WELFARE

67.1(162)	Definitions
67.2(162)	Animals included in rules
67.3(162)	Housing facilities and primary enclosures
67.4(162)	General care and husbandry standards
67.5(162)	Transportation

67.6(162)	Purchase, sale, trade and adoption
67.7(162)	Boarding kennels, commercial kennels, animal shelters, pounds and dealers
67.8(162)	Dog day cares
67.9(162)	In-home facilities
67.10(162)	Rescues
67.11(162)	Foster oversight organizations and foster care homes
67.12(162)	Public health
67.13(162)	Access, seizure and impoundment
67.14(162)	Loss of license or denial of license
67.15(162)	Applicability to commercial establishments with federal licenses
67.16(162)	Acceptable forms of euthanasia
67.17(162)	Greyhound breeder or farm fee
67.18(162)	Research facilities

CHAPTER 68 DAIRY

68.1(192,194)	Definitions
68.2(192)	Licenses and permits required
68.3	Reserved
68.4(192)	Certification of personnel
68.5(190,192,194)	Milk tests
68.6(190,192,194)	Test bottles
68.7 and 68.8	Reserved
68.9(192,194)	Tester's license
68.10(192,194)	Contaminating activities prohibited in milk plants
68.11(192,194)	Suspension of dairy farm permits

GRADE A MILK

68.12(192)	Milk standards
68.13(192,194)	Public health service requirements
68.14(190,192,194,195)	Laboratories

GRADE B MILK

68.15(192,194)	Milk standards
68.16(194)	Legal milk
68.17(194)	New producers
68.18(194)	Testing and exclusion of Class III milk
68.19(194)	Unlawful milk
68.20(194)	Price differential
68.21(194)	Penalties for plants and producers
68.22(192,194)	Farm requirements for milk for manufacturing
68.23 to 68.25	Reserved
68.26(190,192,194)	Tests for abnormal milk
68.27(192,194)	Standards for performing farm inspections

DAIRY FARM WATER

68.28 to 68.34	Reserved
68.35(192)	Dairy farm water supply
68.36(192)	Antibiotic testing
68.37(192,194)	Milk truck approaches
68.38 and 68.39	Reserved

MILK TANKER, MILK HAULER, MILK GRADER, CAN MILK TRUCK BODY

68.40(192)	Definitions
68.41(192)	Bulk milk tanker license required

68.42(192)	Bulk milk tanker construction
68.43(192)	Bulk milk tanker cleaning and maintenance
68.44(192)	Bulk tanker sanitization
68.45(192)	Bulk milk tanker cleaning facility
68.46(192)	Bulk milk tanker cleaning tag
68.47(192)	Dairy plant, receiving station or transfer station records
68.48(192)	Milk hauler license required
68.49	Reserved
68.50(192)	Supplies required for milk collection and sampling
68.51(192)	Milk hauler sanitization
68.52(192)	Examining milk by sight and smell
68.53(192)	Milk hauler hand washing
68.54(192)	Milk temperature
68.55(192)	Connecting the milk hose
68.56(192)	Measuring the milk in the bulk tank
68.57(192)	Milk sample for testing
68.58(192)	Milk collection record
68.59(192)	Loading the milk from the bulk tank to the milk tanker
68.60(192)	Milk samples required for testing
68.61(192)	Bulk milk sampling procedures
68.62(192)	Temperature control sample
68.63(192)	Producer sample identification
68.64(192)	Care and delivery of producer milk samples
68.65(192)	Milk sample carrying case
68.66(192)	Bulk milk delivery
68.67(192)	False samples or records
68.68(192)	Violations prompting immediate suspension
68.69(192)	Milk grader license required
68.70(192)	New milk grader license applicant
68.71(192,194)	Can milk truck body

CHAPTER 69

MILK ROOM AND BULK TANKS FOR MANUFACTURING MILK

69.1(192)	Milk room
69.2(192)	Drainage
69.3(192)	Walls and ceilings
69.4(192)	Milk room windows
69.5(192)	Doors
69.6(192)	Ventilation
69.7(192)	Bulk tank location
69.8(192)	Hose port
69.9(192)	Safety regulations
69.10(192)	Properly located tank

CHAPTER 70

Reserved

CHAPTER 71

STANDARDS FOR DAIRY PRODUCTS

71.1(190)	Dairy products
71.2(189,210)	Requirements for packaging and labeling
71.3(210)	Requirements for the method of sale of commodities
71.4(210)	Requirements for unit pricing

- 71.5(189,190) Flavors
71.6(190) Standard for light butter

CHAPTERS 72 to 75

Reserved

CHAPTER 76

MEAT AND POULTRY INSPECTION

- 76.1(189A) Federal Wholesome Meat Act regulations adopted
76.2(189A) Federal Wholesome Meat Act regulations adopted
76.3(189A) Federal Poultry Products Inspection Act regulations adopted
76.4(189A) Inspection required
76.5(189A) Custom/exempt facilities sanitation standard operating procedures
76.6(189A) Forms and marks
76.7(189A) Products to be marked with official marks
76.8(189A,167) Registration
76.9(189A,167) Dead, dying, disabled or diseased animals
76.10(189A) Denaturing and identification of livestock or poultry products not intended for use as human food
76.11(189A,167) Transportation of decharacterized inedible meat or carcass parts
76.12(189A) Records
76.13(189A) Voluntary inspections of exotic animals
76.14(189A) Federal Wholesome Meat Act regulations adopted for the regulation of farm deer

CHAPTER 77

DANGEROUS WILD ANIMALS

- 77.1(717F) Definitions
77.2(717F) Prohibitions
77.3(717F) Continued ownership—requirements of the individual
77.4(717F) Continued ownership—insurance required
77.5(717F) Continued ownership—electronic identification device
77.6(717F) Continued ownership—registration form
77.7(717F) Continued ownership—registration fee
77.8(717F) Continued ownership—records required
77.9(717F) Continued ownership—enclosure required
77.10(717F) Continued ownership—signs required
77.11(717F) Escape notification required
77.12(717F) Relinquishment
77.13(717F) Seizure, custody and disposal
77.14(717F) Exemptions

CHAPTERS 78 to 84

Reserved

CHAPTER 85

WEIGHTS AND MEASURES

WEIGHTS

- 85.1(215) “Sensibility reciprocal” defined
85.2 to 85.4 Reserved
85.5(215) “Counter scale” defined
85.6(215) “Spring and computing scales” defined
85.7(215) “Automatic grain scale” defined
85.8(215) “Motor truck scales” defined

85.9(215)	“Livestock scales” defined
85.10(215)	“Grain dump scales” defined
85.11(215)	Scale pit
85.12(215)	Pitless scales
85.13(215)	Master weights
85.14(215)	Scale design
85.15(215)	Weighbeams
85.16(215)	Beam box
85.17	Reserved
85.18(215)	Weight capacity
85.19(215)	Provision for sealing coin slot
85.20(215)	Stock racks
85.21(215)	Lengthening of platforms
85.22(215)	Accessibility for testing purposes
85.23(215)	Assistance in testing operations
85.24(215)	Beam scale
85.25(215)	Spring scale
85.26(215)	Weighbeam or beam
85.27(215)	Livestock scale

SCALES

85.28(215)	Wheel-load weighers and axle-load scales
85.29	Reserved

REGISTERED SERVICERS

85.30(215)	Servicer’s license fee
85.31 and 85.32	Reserved

MEASURES

85.33(214A,208A)	Motor fuel and antifreeze tests and standards
85.34(215)	Tolerances on petroleum products measuring devices
85.35(215)	Meter adjustment
85.36(215)	Recording elements
85.37(215)	Air eliminator
85.38(215)	Delivery outlets
85.39(189,215)	Weights and measures
85.40(215)	ILP inspection tag or mark
85.41(215)	Meter repair
85.42(215)	Security seal
85.43(215)	LP-gas meter repairs
85.44(215)	LP-gas delivery
85.45(215)	LP-gas meter registration
85.46(215)	Reporting new LP-gas meters
85.47	Reserved
85.48(214A,215)	Advertisement of the price of liquid petroleum products for retail use
85.49(214A,215)	Gallage determination for retail sales
85.50(214,214A,215)	Blender pumps
85.51	Reserved

MOISTURE-MEASURING DEVICES

85.52(215A)	Testing devices
85.53(215A)	Rejecting devices
85.54(215,215A)	Specifications and standards for moisture-measuring devices
85.55 and 85.56	Reserved

85.57(215) Testing high-moisture grain
 85.58 to 85.62 Reserved

HOPPER SCALES

85.63(215) Hopper scales

CHAPTER 86
 E-15 ACCESS STANDARD

86.1(214A) Definitions
 86.2(214A) E-15 access standard—retail dealer compliance
 86.3(214A) Incompatible infrastructure waivers
 86.4(214A) Small retail motor fuel site exemption
 86.5(214A) Terminable events

CHAPTERS 87 to 89
 Reserved

CHAPTER 90
 STATE LICENSED WAREHOUSES
 AND WAREHOUSE OPERATORS

90.1(203C) Application of rules
 90.2(203C) Definitions
 90.3(203C) Types of products to be warehoused
 90.4(203C,203D) Application for a warehouse operator license
 90.5(203C) Warehouse operator license
 90.6(203C) Posting of license
 90.7(203C) Renewal, expiration and reinstatement of license—payment of license fee
 90.8(203C) Financial statements
 90.9(203C) Bonds and irrevocable letters of credit
 90.10(203C) Insurance
 90.11(203C) Notice to the warehouse bureau
 90.12(203C) Issuance of warehouse receipts
 90.13(203C) Cancellation of warehouse receipts
 90.14(203C) Lost or destroyed receipt
 90.15(203C) Warehouse receipts
 90.16(203C) Tariffs
 90.17(203C) Records
 90.18(203C) Adjustment of records
 90.19(203C) Shrinkage due to moisture
 90.20(203C) Monthly grain statements
 90.21(203C) Grain stored in another warehouse
 90.22(203C) Warehouse operator's obligation and storage
 90.23(203C) Storing of products
 90.24(203C) Facilities
 90.25(203C) Maintenance of storage facilities
 90.26(203C) Temporary grain storage facilities
 90.27(203C) Emergency ground pile storage space
 90.28(203C) Polyethylene (polyvinyl) bag storage space
 90.29(203C) Prioritization of inspections of warehouse operators
 90.30(203C) Department of agriculture and land stewardship enforcement procedures
 90.31(203C) Review proceedings

CHAPTER 91
LICENSED GRAIN DEALERS

91.1(203)	Application of rules
91.2(203)	Definitions
91.3(203,203D)	Application for a grain dealer license
91.4(203)	Grain dealer license not transferable
91.5(203)	Posting of license
91.6(203)	Surrender of license
91.7(203)	Renewal, expiration and reinstatement of license—payment of license and indemnity fund fees
91.8(203)	Financial statements
91.9(203)	Bonds and irrevocable letters of credit
91.10(203)	Payment
91.11(203)	Books and records
91.12(203)	Assignment of contracts
91.13(203)	Filing of monthly grain statement and reports
91.14(203)	Notice to the warehouse bureau
91.15(203)	Shrinkage due to moisture
91.16(203)	Requirements for Class 2 licensees
91.17(203)	Requirements for licensees authorized to issue credit-sale contracts
91.18(203)	Department of agriculture and land stewardship enforcement procedures
91.19(203)	Review proceedings
91.20(203)	Prioritization of inspections of grain dealers
91.21(203)	Claims against credit-sale contract bond
91.22(203)	Electronic grain contracts
91.23(203)	Electronic grain contract providers and provider agreements
91.24(203)	Electronic grain contract users and user agreements
91.25(203)	Electronic grain contracts—issuance and form
91.26(203)	Security of a provider’s electronic central filing system or a licensee’s electronic database

CHAPTER 92
PARTICIPATION IN GRAIN INDEMNITY FUND

92.1(203D)	Mandatory participation in fund
92.2(203D)	Required fees
92.3(203D)	New license applicants
92.4(203D)	Due date for payment of the per-bushel and participation fees
92.5(203D)	Penalty for delinquent submission of per-bushel and participation fees
92.6(203D)	Penalty for delinquent payment of per-bushel fee discovered during examination

CHAPTER 93
GRAIN INDEMNITY FUND BOARD—ORGANIZATION AND OPERATIONS

93.1(203D)	Location
93.2(203D)	The board
93.3(203D)	Authority of the board
93.4(203D)	Meetings
93.5(203D)	Minutes
93.6(203D)	Board decisions
93.7(203D)	Records

CHAPTER 94
CLAIMS AGAINST THE GRAIN DEPOSITORS
AND SELLERS INDEMNITY FUND

94.1(203D)	Definitions
94.2(203D)	By whom claims can be made
94.3(203D)	Procedure for filing claims
94.4(203D)	Time limitations
94.5(203D)	Claims by depositors where bureau is receiver
94.6(203D)	Notice of claims
94.7(203D)	Report by bureau
94.8(203D)	Determination of claims
94.9(203D)	Appeal from determination
94.10(203D)	Payment of valid claims—conflicting interests

CHAPTER 95
CIVIL PENALTIES

95.1(203,203C)	Definitions
95.2(203,203C)	Grain industry peer review panel
95.3(203,203C)	Organization and location
95.4(203,203C)	Membership
95.5(203,203C)	Staff
95.6(203,203C)	Meetings
95.7(203,203C)	Criteria for assessing civil penalties
95.8(203,203C)	Notice of civil penalty assessment—informal settlement
95.9(203,203C)	Panel review
95.10(203,203C)	Scope of panel review
95.11(203,203C)	Panel response
95.12(203,203C)	Civil penalty assessment
95.13(203,203C)	Judicial assessment
95.14(203,203C)	Civil penalty payment

CHAPTER 96
HEMP

96.1(204)	Definitions
96.2(204)	Licensing
96.3(204)	National criminal history record check
96.4(204)	Licensee reports
96.5(204)	Fees
96.6(204)	Annual review of licensees to ensure licensure compliance
96.7(204)	Sampling procedures for official testing of hemp for THC content
96.8(204)	Approved testing methods of hemp for THC content
96.9(204)	Harvesting timing
96.10(204)	Order of destruction
96.11(204)	Negligent violations
96.12(204)	Negligent violation program
96.13(204)	State plan

CHAPTER 93
GRAIN INDEMNITY FUND BOARD—ORGANIZATION AND OPERATIONS

[Prior to 7/27/88, 21—Ch 63]

21—93.1(203D) Location. The office of the grain indemnity fund board is located in the Wallace State Office Building, Des Moines, Iowa; telephone (515)281-5321; mailing address: Grain Indemnity Fund Board, c/o Grain Warehouse Bureau, Iowa Department of Agriculture and Land Stewardship, Henry A. Wallace Building, Des Moines, Iowa 50319.

This rule is intended to implement Iowa Code section 203D.4.

21—93.2(203D) The board. The grain indemnity fund board consists of seven members: the secretary of agriculture or the secretary's designee who shall serve as chairperson, the state treasurer or the state treasurer's designee who shall serve as treasurer, a representative of the banking industry and four representatives of the grain industry. Grain industry representatives shall consist of two grain producers, one representative of warehouse operators licensed in accordance with Iowa Code section 203C.6 and one representative of grain dealers licensed in accordance with Iowa Code section 203.3. Each industry representative shall be appointed by the governor from a list of three nominees made by the secretary of agriculture.

This rule is intended to implement Iowa Code section 203D.4.
[ARC 9388B, IAB 2/23/11, effective 3/30/11]

21—93.3(203D) Authority of the board. The board has authority to determine the amount and validity of claims made against the fund, to review and adjust the per-bushel fee and the grain dealer and warehouse operator participation fee, and to approve costs of administering the fund. In addition, the board has the authority to act as an advisor to the secretary of agriculture on administrative matters affecting the fund, and as a result the board will make only policy recommendations in regard to the areas of administration delegated to the department in Iowa Code chapter 203D.

This rule is intended to implement Iowa Code section 203D.4.
[ARC 9388B, IAB 2/23/11, effective 3/30/11]

21—93.4(203D) Meetings. Unless otherwise determined by the chairperson, the board will meet at 2 p.m. on the third Thursday of each month. In-person board meetings will generally be held in a conference room in the Henry A. Wallace building. Telephone conference call meetings may be permitted and will generally be hosted from the offices of the grain warehouse bureau of the Iowa department of agriculture and land stewardship in the Henry A. Wallace building. The establishment and public notice of meeting dates and locations are the responsibility of the chairperson, unless the majority of the members of the board eligible to vote request a meeting. In addition, the board will schedule meetings when circumstances require the board to address claims made against the fund and, for these meetings, establishment and public notice of meeting dates and locations are the responsibility of the chairperson.

93.4(1) Agenda. The tentative agenda is prepared by the chairperson in advance of the board meeting and will be mailed to board members in advance of the meeting date. A copy of the agenda will be mailed to those members of the public who request it and will be prominently posted at the board's office at least 24 hours before the meeting. Members of the public wishing to be scheduled on the board's agenda should notify the chairperson ten days in advance of the meeting and provide written materials explaining their reasons for wishing to address the board. In the case of a board meeting held to deal with claims against the fund, the filing of a written appeal under rule 21—94.9(203D) will satisfy the requirements of the preceding sentence. The chairperson shall have the authority to make all final decisions on the content and length of agenda items.

93.4(2) General conduct of meetings. The chairperson presides at all board meetings. Only individuals recognized by the presiding officer may address the board; in general, Robert's Rules of Order will govern the meeting unless otherwise stated in this chapter or by special action of the board.

In all discussions before the board, members of the public shall address any questions for the board to the presiding officer. Individual questioning of board members will not be allowed without the explicit consent of the presiding officer and the board members in question.

93.4(3) Voting. The board consists of seven members who are all eligible to vote on issues. A majority of board members shall constitute a quorum. The affirmative vote of four board members is necessary to carry an action.

93.4(4) Public participation. All meetings are open to the public in accordance with the open meetings law, Iowa Code chapter 21, except that portions of a meeting may be closed in accordance with the open meetings law. In the chairperson's discretion, a 15-minute public forum may be scheduled on each agenda of regularly scheduled meetings to allow the public, if necessary, an opportunity to address the board on any issue that may have arisen after the agenda was posted.

This rule is intended to implement Iowa Code sections 203D.4, 203D.5, 203D.5A and 203D.6.
[ARC 9388B, IAB 2/23/11, effective 3/30/11]

21—93.5(203D) Minutes. The minutes of all board meetings are recorded and kept by the grain warehouse bureau in the board's office.

This rule is intended to implement Iowa Code section 203D.4.
[ARC 9388B, IAB 2/23/11, effective 3/30/11]

21—93.6(203D) Board decisions. The actions of the board will be authoritatively recorded in the minutes of the board meeting at which the actions were taken. The board may adopt, amend, or repeal rules subject to Iowa Code chapter 17A to govern the operations of the board, to adjust or waive the per-bushel fee and the annual dealer-warehouse fee, and to govern the process of making claims against the fund. These rules shall be published by the department in the Iowa Administrative Code. The board may also recommend the adoption of other rules by the department relating to the fund. The content of any rules will be authoritatively established when they are published by the department in the Iowa Administrative Code.

This rule is intended to implement Iowa Code sections 203D.4, 203D.5, 203D.5A and 203D.6.

21—93.7(203D) Records. The records of all the business transacted and other information with respect to the activities of the board are public records and are on file in the board's office. All records including board minutes are available for inspection during regular business hours. Copies may be obtained at a cost of 25 cents per page.

This rule is intended to implement Iowa Code section 203D.4.

21—93.8(203D) Waiver of per-bushel and participation fees. Rescinded ARC 6954C, IAB 3/22/23, effective 4/26/23.

[Filed 2/20/87, Notice 12/17/86—published 3/11/87, effective 4/15/87]

[Filed 11/24/87, Notice 7/1/87—published 12/16/87, effective 3/16/88]

[Filed without Notice 4/29/88—published 5/18/88, effective 7/1/88]

[Filed emergency 7/8/88 after Notice 6/1/88—published 7/27/88, effective 7/8/88]

[Filed 10/21/05, Notice 9/14/05—published 11/23/05, effective 12/28/05]

[Filed ARC 9388B (Notice ARC 9165B, IAB 10/20/10), IAB 2/23/11, effective 3/30/11]

[Filed ARC 6954C (Notice ARC 6803C, IAB 1/11/23), IAB 3/22/23, effective 4/26/23]

CHAPTER 2
OPERATION OF STATE SOIL CONSERVATION AND WATER QUALITY COMMITTEE

27—2.1(161A) Scope. This chapter governs the conduct of business by the state soil conservation and water quality committee. Rule-making proceedings held as part of committee meetings and contested case proceedings involving the committee are consistent with Iowa Code chapter 17A.

[ARC 3243C, IAB 8/2/17, effective 9/6/17]

27—2.2(161A) Time of meetings. The committee meets quarterly. The chairperson or a majority of the committee may establish meetings at more frequent intervals.

[ARC 6957C, IAB 3/22/23, effective 4/26/23]

27—2.3(161A) Place of meetings. Meetings are held in the Henry A. Wallace Building, 900 East Grand Avenue, Des Moines, Iowa, or at other locations as appropriate. The meeting place will be specified in the agenda.

27—2.4(161A) Notification of meetings. The director of the soil conservation and water quality division shall provide public notice of all meeting dates, locations, and tentative agenda.

2.4(1) Form of notice. Notice of meetings is given by posting the tentative agenda and by distribution upon request. The agenda lists the time, date, place, and topics to be discussed at the meeting. The agenda shall include an opportunity for the public to address the committee on any issue related to the duties and responsibilities of the committee, except as otherwise provided in these rules.

2.4(2) Posting of agenda. The tentative agenda for each meeting will be posted at the division's offices on the second floor, Henry A. Wallace Building, normally at least five days prior to the meeting. The agenda will be posted at least 24 hours prior to the meeting, unless, for good cause such notice is impossible or impractical, in which case as much notice as is reasonably possible will be given.

2.4(3) Distribution of agenda. Agenda will be mailed to anyone who files a request with the director. The request should state whether the agenda for a particular meeting is desired, or whether the requester desires to be on the division's mailing list to receive the agenda for all meetings of the state soil conservation and water quality committee.

2.4(4) Amendment to agenda. Any amendments to the agenda after posting and distribution under subrules 2.4(2) and 2.4(3) will be posted, but will not be mailed. The amended agenda will be posted at least 24 hours prior to the meeting, unless for good cause such notice is impossible or impractical, in which case as much notice as is reasonably possible will be given. The committee may adopt amendments to the agenda at the meeting only if good cause exists requiring expeditious discussion or action on such matters. The reasons and circumstances necessitating such agenda amendments, or those given less than 24 hours' notice by posting, shall be stated in the minutes of the meeting.

2.4(5) Supporting material. Written materials provided to the committee with the agenda may be examined and copied as provided in the public information rules of the department. The director may require a fee to cover the reasonable cost to the division to provide the copies, in accordance with rules of the department.

[ARC 3243C, IAB 8/2/17, effective 9/6/17]

27—2.5(161A) Attendance and participation by the public.

2.5(1) Attendance. All meetings are open to the public. The committee may exclude the public from portions of the meeting only in accordance with Iowa Code section 21.5.

2.5(2) Participation.

a. Items on agenda. Presentations to the committee may be made at the discretion of the chairperson.

b. Items not on agenda. Iowa Code section 21.4 requires the committee to give notice of its agenda. The committee will not take action on a matter not on the agenda, except in accordance with subrule 2.4(4). Presentations to the committee on subjects not on the agenda may be made at the discretion of

the chairperson. Persons who wish the committee to take action on a matter not on the agenda should file a request with the director to place that matter on the agenda of a subsequent meeting.

c. Meeting decorum. The chairperson may limit participation as necessary for the orderly conduct of agency business.

2.5(3) Use of cameras and recording devices. Cameras and recording devices may be used during meetings provided they do not interfere with the orderly conduct of the meeting. The chairperson may order the use of these devices be discontinued if they cause interference and may exclude those persons who fail to comply with that order.

27—2.6(161A) Quorum and voting requirements.

2.6(1) Quorum. A majority of the voting members of the committee constitutes a quorum.

2.6(2) Voting. The concurrence of a majority of the committee members is required to determine any matter before the committee for action, except for a vote to close a meeting which requires the concurrence of two-thirds of the members of the committee present.

[ARC 4956C, IAB 2/26/20, effective 4/1/20]

27—2.7(161A) Conduct of meeting.

2.7(1) General. Meetings will be conducted in accordance with Robert's Rules of Order unless otherwise provided in these rules. Voting shall be by voice or by roll call. Voting shall be by voice unless a voice vote is inconclusive, a member of the committee requests a roll call, or the vote is on a motion to close a portion of a meeting. The chairpersons shall announce the result of the vote.

2.7(2) Voice votes. All committee members present should respond when a voice vote is taken. The response shall be aye, nay, or abstain.

a. All members present shall be recorded as voting aye on any motions when there are no nay votes or abstentions heard.

b. Any member who abstains shall state at the time of the vote the reason for abstaining. The abstention and the reason for it shall be recorded in the minutes.

2.7(3) Provisions of information. The chairperson may recognize any agency staff member for the provision of information relative to an agenda item.

27—2.8(161A) Minutes, transcripts, and recordings of meetings.

2.8(1) Recordings. The director shall record by mechanized means each meeting and shall retain the recording for at least one month. Recordings of closed sessions shall be sealed and retained at least one year.

2.8(2) Transcripts. The division does not routinely prepare transcripts of meetings. The division will have transcripts of meetings, except for closed sessions, prepared upon receipt of a request for a transcript and payment of a fee to cover the cost to the division of preparing the transcript.

2.8(3) Minutes. The director shall keep minutes of each meeting. Minutes shall be reviewed and approved by the committee and retained permanently by the director. The approved minutes shall be signed by the director and the committee chairperson.

27—2.9(161A) Officers and duties.

2.9(1) Officers. The officers of the committee are the chairperson and the vice chairperson.

2.9(2) Duties. The chairperson shall preside at the meetings and shall exercise the powers conferred upon the chairperson. The vice chairperson shall perform the duties of the chairperson when the chairperson is absent or when directed by the chairperson.

27—2.10(161A) Election and succession of officers.

2.10(1) Elections. Officers shall be elected annually during June and shall assume office effective July 1.

2.10(2) Succession.

a. If the chairperson does not serve out the elected term, the vice chairperson shall succeed the chairperson for the remainder of the term. A special election shall be held to elect a new vice chairperson to serve the remainder of the term.

b. If the vice chairperson does not serve out the elected term, a special election shall be held to elect a new vice chairperson to serve the remainder of the term.

These rules are intended to implement Iowa Code chapter 161A.

[Filed 12/8/89, Notice 10/4/89—published 12/27/89, effective 1/31/90]

[Filed ARC 3243C (Notice ARC 3086C, IAB 6/7/17), IAB 8/2/17, effective 9/6/17]

[Filed ARC 4956C (Notice ARC 4839C, IAB 1/1/20), IAB 2/26/20, effective 4/1/20]

[Filed ARC 6957C (Notice ARC 6817C, IAB 1/11/23), IAB 3/22/23, effective 4/26/23]

IOWA FINANCE AUTHORITY[265]

[Prior to 7/26/85, Housing Finance Authority[495]]
 [Prior to 4/3/91, Iowa Finance Authority[524]]

CHAPTER 1

GENERAL

- 1.1(16) Purpose
- 1.2(16) Mission
- 1.3(16) Organization, programs and operations
- 1.4(16) Location where the public may submit requests or obtain information
- 1.5(16) Forms

CHAPTER 2

LOAN PROGRAMS

GENERAL PROVISIONS

- 2.1(16) Administrative agents

TERMS AND CONDITIONS

- 2.2(16) Interest and fees
- 2.3 Reserved
- 2.4(16) Loan conditions
- 2.5(16) Security for loans
- 2.6(16) Types of loans
- 2.7(16) Delinquency and foreclosure
- 2.8(16) Application processing
- 2.9(16) Mortgage purchase or loans to lenders for existing, newly built single-family or multifamily housing—general information
- 2.10(16) Assumption of mortgages

CHAPTER 3

MULTIFAMILY HOUSING

MULTIFAMILY LOAN PROGRAM

- 3.1(16) Purpose
- 3.2(16) Available funds
- 3.3(16) Intent of the authority
- 3.4(16) Application procedure
- 3.5(16) Program guidelines
- 3.6 and 3.7 Reserved
- 3.8(16) Multifamily loan program for workforce housing loan assistance
- 3.9 Reserved
- 3.10(16) Authority analysis of applications
- 3.11(16) Discretion of authority board

CHAPTER 4

GENERAL REVENUE BOND PROCEDURES

- 4.1(16) Revenue bonds authorized
- 4.2(16) Participating lenders
- 4.3(16) Procedures for project sponsors
- 4.4(16) Authority review
- 4.5 Reserved
- 4.6(16) Procedures following bond issuance
- 4.7(16) Right to audit

CHAPTERS 5 and 6
Reserved

CHAPTER 7
CONTESTED CASES

7.1(17A)	Scope and applicability
7.2(17A)	Definitions
7.3(17A)	Time requirements
7.4(17A)	Requests for contested case proceeding
7.5(17A)	Notice of hearing
7.6(17A)	Presiding officer
7.7(17A)	Waiver of procedures
7.8(17A)	Telephone or video proceedings
7.9(17A)	Disqualification
7.10(17A)	Consolidation—severance
7.11(17A)	Pleadings
7.12(17A)	Service and filing of pleadings and other papers
7.13(17A)	Discovery
7.14(17A)	Subpoenas
7.15(17A)	Motions
7.16(17A)	Prehearing conference
7.17(17A)	Continuances
7.18(17A)	Withdrawals
7.19(17A)	Intervention
7.20(17A)	Hearing procedures
7.21(17A)	Evidence
7.22(17A)	Default
7.23(17A)	Ex parte communication
7.24(17A)	Recording costs
7.25(17A)	Interlocutory appeals
7.26(17A)	Posthearing procedures and orders
7.27(17A)	Appeals and review
7.28(17A)	Applications for rehearing
7.29(17A)	Stays of authority actions
7.30(17A)	No factual dispute contested cases
7.31(17A)	Emergency adjudicative proceedings
7.32(17A,16)	Informal procedure prior to hearing

CHAPTER 8
PRIVATE ACTIVITY BOND ALLOCATION

8.1(7C)	General
8.2(7C)	Forms and applications
8.3(7C)	Applications for current allocation received prior to the calendar year for such allocation
8.4(7C)	Application for current allocation received during the calendar year
8.5(7C)	Certification of current allocation
8.6(7C)	State ceiling carryforwards
8.7(7C)	Expiration of applications and allocations
8.8(7C)	Resubmission of expired allocations
8.9(7C)	Use by political subdivisions
8.10(7C)	Application and allocation fees

CHAPTER 9
TITLE GUARANTY DIVISION

9.1(16)	Definitions
9.2(16)	Purpose
9.3(16)	Mission
9.4(16)	Organization
9.5(16)	Operation
9.6(16)	Participants
9.7(16)	Services offered
9.8(16)	Claims
9.9(16)	Mortgage release certificate
9.10(16)	Rules of construction
9.11(16)	Seal

CHAPTER 10
MORTGAGE CREDIT CERTIFICATES

10.1(16)	General
10.2(16)	Participating lenders
10.3(16)	Eligible borrowers
10.4(16)	MCC procedures

CHAPTER 11
IOWA MAIN STREET LOAN PROGRAM

11.1(16)	Program description
11.2(16)	Waiver
11.3(16)	Main street loan program
11.4(16)	Definitions
11.5(16)	Application
11.6(16)	Public benefit
11.7(16)	Loan criteria

CHAPTER 12
LOW-INCOME HOUSING TAX CREDITS

12.1(16)	Qualified allocation plans
12.2(16)	Location of copies of the plans

CHAPTER 13
PUBLIC RECORDS AND FAIR INFORMATION PRACTICES
(Uniform Rules)

13.1(17A,22)	Definitions
13.3(17A,22)	Requests for access to records
13.4(17A,22)	Access to confidential records
13.6(17A,22)	Procedure by which additions, dissents, or objections may be entered into certain records
13.9(17A,22)	Availability of records

CHAPTER 14
Reserved

CHAPTER 15
PURCHASING

15.1(16)	Applicability of competitive bidding
15.2(16)	Methods of obtaining bids or proposals used by the authority
15.3(16)	Items purchased through the department of administrative services

15.4(16)	Posting solicitations
15.5(16)	Contract purchases
15.6(16)	Blanket purchase agreements
15.7(16)	Bids and proposals to conform to specifications
15.8(16)	Time of delivery
15.9(16)	Cash discounts
15.10(16)	Ties
15.11(16)	Time of submission
15.12(16)	Modification or withdrawal of bids
15.13(16)	Financial security
15.14(16)	Rejection of bids and proposals
15.15(16)	Vendor appeals

CHAPTER 16 DECLARATORY ORDERS

16.1(17A)	Petition for declaratory order
16.2(17A)	Notice of petition
16.3(17A)	Intervention
16.4(17A)	Briefs
16.5(17A)	Inquiries
16.6(17A)	Service and filing of petitions and other papers
16.7(17A)	Consideration
16.8(17A)	Action on petition
16.9(17A)	Refusal to issue order
16.10(17A)	Contents of declaratory order—effective date
16.11(17A)	Copies of orders
16.12(17A)	Effect of a declaratory order

CHAPTER 17 PROCEDURE FOR RULE MAKING

17.1(17A)	Applicability
17.2(17A)	Advice on possible rules before notice of proposed rule adoption
17.3(17A)	Public rule-making docket
17.4(17A)	Notice of proposed rule making
17.5(17A)	Public participation
17.6(17A)	Regulatory analysis
17.7(17A,25B)	Fiscal impact statement
17.8(17A)	Time and manner of rule adoption
17.9(17A)	Variance between adopted rule and published notice of proposed rule adoption
17.10(17A)	Exemptions from public rule-making procedures
17.11(17A)	Concise statement of reasons
17.12(17A)	Contents, style, and form of rule
17.13(17A)	Authority rule-making record
17.14(17A)	Filing of rules
17.15(17A)	Effectiveness of rules prior to publication
17.16(17A)	General statements of policy
17.17(17A)	Review by authority of rules

CHAPTER 18 WAIVERS FROM ADMINISTRATIVE RULES

18.1(17A,16)	Definitions
18.2(17A,16)	Scope
18.3(17A,16)	Applicability of chapter

18.4(17A,16)	Criteria for waiver
18.5(17A,16)	Filing of petition
18.6(17A,16)	Content of petition
18.7(17A,16)	Additional information
18.8(17A,16)	Notice
18.9(17A,16)	Hearing procedures
18.10(17A,16)	Ruling
18.11(17A,16)	Public availability
18.12(17A,16)	Submission of waiver information
18.13(17A,16)	Voiding or cancellation
18.14(17A,16)	Violations
18.15(17A,16)	Defense
18.16(17A,16)	Judicial review

CHAPTER 19
STATE HOUSING TRUST FUND

19.1(16)	Trust fund allocation plans
19.2(16)	Location of copies of the plans

CHAPTER 20
SENIOR LIVING REVOLVING LOAN PROGRAM

20.1(16)	Purpose
20.2(16)	Priority of loan awards
20.3(16)	Application process
20.4(16)	Program guidelines
20.5(16)	Authority analysis of applications
20.6(16)	Discretion of authority board
20.7(16)	Closing/advance of funds

CHAPTER 21
HOME AND COMMUNITY-BASED SERVICES REVOLVING LOAN PROGRAM

21.1(16)	Purpose
21.2(16)	Available funds
21.3(16)	Intent of the authority
21.4(16)	Application procedure
21.5(16)	Program guidelines
21.6(16)	Authority analysis of applications
21.7(16)	Discretion of authority board
21.8(16)	Closing/advance of funds

CHAPTER 22
IOWA AFTERCARE SERVICES RENT SUBSIDY PROGRAM

22.1(16,PL106-169)	Purpose
22.2(16,PL106-169)	Definitions
22.3(16,PL106-169)	Eligibility requirements for direct rent subsidy
22.4(16,PL106-169)	Application for direct rent subsidy
22.5(16,PL106-169)	Amount of rent subsidy
22.6(16,PL106-169)	Redetermination of direct rent subsidy eligibility
22.7(16,PL106-169)	Termination of rent subsidy payments
22.8(16,PL106-169)	Eligibility requirements for transitional apartment subsidy
22.9(16,PL106-169)	Application for transitional apartment subsidy
22.10(16,PL106-169)	Amount of transitional apartment subsidy
22.11(16,PL106-169)	Redetermination of transitional apartment subsidy eligibility

- 22.12(16,PL106-169) Termination of transitional apartment subsidy payments
- 22.13(16,PL106-169) Fraudulent practices relating to the aftercare rent subsidy program
- 22.14(16,PL106-169) Appeals

CHAPTER 23

TRANSITIONAL HOUSING REVOLVING LOAN PROGRAM

- 23.1(16) Purpose
- 23.2(16) Priority of loan awards
- 23.3(16) Application process
- 23.4(16) Program guidelines
- 23.5(16) Authority analysis of applications
- 23.6(16) Discretion of authority board
- 23.7(16) Closing/advance of funds

CHAPTER 24

HOME AND COMMUNITY-BASED SERVICES RENT SUBSIDY PROGRAM

- 24.1(16) Purpose
- 24.2(16) Definitions
- 24.3(16) Eligibility requirements
- 24.4(16) Application
- 24.5(16) Amount of rent subsidy
- 24.6(16) Redetermination of eligibility
- 24.7(16) Termination of rent subsidy payments

CHAPTER 25

Reserved

CHAPTER 26

WATER POLLUTION CONTROL WORKS AND DRINKING WATER FACILITIES FINANCING

- 26.1(16) Statutory authority
- 26.2(16) Purpose
- 26.3(16) Definitions
- 26.4(16) Project funding
- 26.5(16) WPCSRF/DWSRF infrastructure construction loans
- 26.6(16) Planning and design loans
- 26.7(16) Disadvantaged community status
- 26.8(16) WPCSRF nonpoint source set-aside loan programs
- 26.9(16) Termination and rectification of disputes

CHAPTER 27

MILITARY SERVICE MEMBER HOME OWNERSHIP ASSISTANCE PROGRAM

- 27.1(16) Purpose
- 27.2(16) Definitions
- 27.3(16) Application procedure and determination of eligibility
- 27.4(16) MHOA award

CHAPTER 28

WASTEWATER AND DRINKING WATER TREATMENT FINANCIAL ASSISTANCE PROGRAM

- 28.1(16) Overview
- 28.2(16) Definitions
- 28.3(16) Project funding
- 28.4(16) Termination; rectification of deficiencies; disputes

CHAPTER 29

JUMP-START HOUSING ASSISTANCE PROGRAM

- 29.1(16) Purpose
- 29.2(16) Definitions
- 29.3(16) Grants to local government participants
- 29.4 Reserved
- 29.5(16) Eligible uses
- 29.6(16) Loan terms
- 29.7(16) Financial assistance subject to availability of funding
- 29.8(16) Funds allocated pursuant to 2009 Iowa Acts, House File 64, division I

CHAPTER 30

QUALIFIED MIDWESTERN DISASTER AREA BOND ALLOCATION

- 30.1(16) General
- 30.2(16) Forms
- 30.3(16) Eligibility for allocation
- 30.4(16) Allocation limit and Iowa department of economic development set-aside
- 30.5(16) Application for allocation
- 30.6(16) Certification of allocation
- 30.7(16) Expiration of allocations
- 30.8(16) Resubmission of expired allocations
- 30.9(16) Application and allocation fees

CHAPTER 31

COUNCIL ON HOMELESSNESS

- 31.1(16) General
- 31.2(16) Duties of the council

CHAPTER 32

IOWA JOBS PROGRAM

- 32.1(16) Purpose
- 32.2(16) Definitions
- 32.3(16) Allocation of funds
- 32.4(16) Local infrastructure competitive grant program
- 32.5(16) Noncompetitive grants
- 32.6(16) General grant conditions
- 32.7(16) Calculation of jobs created
- 32.8(16) Grant awards
- 32.9(16) Administration of awards

CHAPTER 33

WATER QUALITY FINANCIAL ASSISTANCE PROGRAM

- 33.1(16,83GA,SF376) Overview
- 33.2(16,83GA,SF376) Definitions
- 33.3(16,83GA,SF376) Small community assistance fund
- 33.4(16,83GA,SF376) Large community assistance fund
- 33.5(16,83GA,SF376) Project priority
- 33.6(16,83GA,SF376) Project funding
- 33.7(16,83GA,SF376) Termination and rectification of disputes

CHAPTERS 34 and 35

Reserved

CHAPTER 36

PUBLIC SERVICE SHELTER GRANT FUND

- 36.1(16,83GA,SF376) Public service shelter grant fund allocation plan
- 36.2(16,83GA,SF376) Location of copies of the plan

CHAPTER 37

RECOVERY ZONE BOND ALLOCATION

- 37.1(16) General
- 37.2(16) Forms
- 37.3(16) Notice from the authority to issuers
- 37.4(16) Notice from issuers to the authority
- 37.5(16) Waiver of RZ bonding authority
- 37.6(16) Application for allocation of recaptured or waived RZ bond authority
- 37.7(16) Allocations
- 37.8(16) Certification of allocation
- 37.9(16) Expiration of allocations
- 37.10(16) Resubmission of expired allocations
- 37.11(16) Application and allocation fees

CHAPTER 38

IOWA JOBS II PROGRAM

- 38.1(16) Purpose
- 38.2(16) Definitions
- 38.3(16) Allocation of funds
- 38.4(16) Iowa jobs II program
- 38.5(16) General grant conditions
- 38.6(16) Calculation of jobs created
- 38.7(16) Grant awards
- 38.8(16) Administration of awards

CHAPTER 39

HOME INVESTMENT PARTNERSHIPS PROGRAM

- 39.1(16) Purpose
- 39.2(16) Definitions
- 39.3(16) Eligible applicants
- 39.4(16) Eligible activities and forms of assistance
- 39.5(16) Application procedure
- 39.6(16) Application requirements
- 39.7(16) Application review criteria
- 39.8(16) Allocation of funds
- 39.9(16) Administration of awards

CHAPTER 40

IOWANS HELPING IOWANS HOUSING ASSISTANCE PROGRAM

- 40.1(16) Purpose
- 40.2(16) Definitions
- 40.3(16) Grants to local government participants
- 40.4 Reserved
- 40.5(16) Eligible uses
- 40.6(16) Loan terms
- 40.7(16) Financial assistance subject to availability of funding

CHAPTER 41
SHELTER ASSISTANCE FUND

- 41.1(16) Purpose
- 41.2(16) Definitions
- 41.3(16) Eligible applicants
- 41.4(16) Eligible activities
- 41.5(16) Ineligible activities
- 41.6(16) Application procedures
- 41.7(16) Application review process
- 41.8(16) Matching contributions
- 41.9(16) Funding awards
- 41.10(16) Requirements placed on recipients
- 41.11(16) Compliance with applicable federal and state laws and regulations
- 41.12(16) Administration

CHAPTER 42
EMERGENCY SOLUTIONS GRANT PROGRAM

- 42.1(16) Purpose
- 42.2(16) Definitions
- 42.3(16) Eligible applicants
- 42.4(16) Eligible activities
- 42.5(16) Ineligible activities
- 42.6(16) Application procedures
- 42.7(16) Application review process
- 42.8(16) Matching requirement
- 42.9(16) Funding awards
- 42.10(16) Compliance with applicable federal and state laws and regulations
- 42.11(16) Administration

CHAPTER 43
COMMUNITY HOUSING AND SERVICES FOR PERSONS WITH DISABILITIES
REVOLVING LOAN PROGRAM

- 43.1(16) Purpose
- 43.2(16) Definitions
- 43.3(16) Award of loan funds
- 43.4(16) Application process
- 43.5(16) Program guidelines
- 43.6(16) Authority analysis of applications
- 43.7(16) Discretion of authority board
- 43.8(16) Closing/advance of funds

CHAPTER 44
IOWA AGRICULTURAL DEVELOPMENT DIVISION

- 44.1(16) General
- 44.2(16) Definitions
- 44.3(16) Beginning farmer loan program eligibility
- 44.4(16) Beginning farmer loan program
- 44.5(16) Loan participation program
- 44.6(16) Beginning farmer tax credit program

CHAPTER 45
MANUFACTURED HOUSING
PROGRAM FUND

45.1(16)	Purpose
45.2(16)	Definitions
45.3(16)	Sources of funds
45.4(16)	Program overview
45.5(16)	Eligible financing
45.6(16)	Linked deposits
45.7(16)	Limits on linked deposits
45.8(16)	Availability of moneys for linked deposits

CHAPTER 46
WATER QUALITY FINANCING PROGRAM

46.1(16)	Overview
46.2(16)	Definitions
46.3(16)	Program administration
46.4(16)	Project funding
46.5(16)	Financial agreements
46.6(16)	Project scoring
46.7(16)	Scoring criteria
46.8(16)	Termination; rectification of deficiencies; disputes

CHAPTER 47
HOUSING RENEWAL PILOT PROGRAM

47.1(89GA, HF2564)	Purpose
47.2(89GA, HF2564)	Definitions
47.3(89GA, HF2564)	Agreement
47.4(89GA, HF2564)	Reporting

CHAPTER 31
COUNCIL ON HOMELESSNESS

265—31.1(16) General.**31.1(1) Location and staff.**

a. The main office of the council is located at the offices of the Iowa finance authority, located at the address set forth in rule 265—1.3(16). Office hours for the council shall be 8 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. Written requests may be submitted to the council at this address. Information about the council is available at this website address: www.iowafinance.com.

b. Staff assistance and administrative support shall be provided by the Iowa finance authority as approved by the executive director.

31.1(2) Composition. The powers of the council are vested in and exercised by 20 members, 11 of whom are voting members and 9 of whom are nonvoting members. Voting members are appointed by the governor in accordance with Iowa Code section 16.2D.

31.1(3) Terms. The 11 voting members selected from the general public shall each serve a two-year term. Terms shall be staggered so half of the voting members are appointed in one year and half are appointed in the year thereafter.

31.1(4) Board officers. The council shall annually elect a chairperson and a vice chairperson and may elect other officers as determined by the council.

31.1(5) Quorum and voting requirements. Seven voting members of the council shall constitute a quorum. Any action taken by the council must be adopted by an affirmative vote of a majority of the quorum. The majority shall not include any member who has a conflict of interest, and a statement by a member who asserts a conflict of interest shall be conclusive for this purpose.

31.1(6) Meetings. The council shall meet at least six times per year. Meetings may be held at the call of the chairperson or whenever a majority of the members so request. The council shall comply with the requirements of Iowa Code chapters 21 and 22. Interested parties are encouraged to attend and participate in council meetings where feasible.

31.1(7) Committees. The council may form an executive committee comprised of not more than six voting members of the council. The membership and size of the committee as well as the terms of the committee members will be established annually by the council. The members of the executive committee may elect a member to serve as a chairperson. A majority of voting committee members constitutes a quorum. Any action taken by the committee must be adopted by an affirmative vote of a majority of its voting members. The chairperson of the council may appoint nonvoting members of the council to serve on the executive committee as nonvoting members. The executive committee will be responsible for reviewing and making recommendations for amendments or changes to the internal rules of procedure. The executive committee will carry out the business of the council between regularly scheduled council meetings.

a. Nominating committee. The council shall annually at its March meeting elect five members, at least two of whom shall be nonvoting members and at least two of whom shall be voting members. The nominating committee shall nominate persons to the governor to fill the voting member positions when they become open. A majority of the members of the nominating committee shall constitute a quorum. Any action taken by the nominating committee must be adopted by an affirmative vote of a majority of its members.

b. Other committees. The council may establish other advisory committees and subcommittees comprised of members of the council to carry out various responsibilities of the council. A majority of the voting members of such a committee shall constitute a quorum. Any action taken by a committee must be adopted by an affirmative vote of a majority of its voting members.

c. Informal working groups. Informal working groups may be assembled from time to time by the chairperson for various tasks.

[ARC 7704B, IAB 4/8/09, effective 5/13/09; ARC 2005C, IAB 5/27/15, effective 7/1/15; ARC 4319C, IAB 2/27/19, effective 4/3/19; ARC 6955C, IAB 3/22/23, effective 4/26/23]

265—31.2(16) Duties of the council. The duties of the council shall be to:

1. Develop a process for evaluating state policies, programs, statutes, and rules to determine whether any state policies, programs, statutes, or rules should be revised to help prevent and alleviate homelessness.
2. Evaluate whether state agency resources could be more efficiently coordinated with other state agencies to prevent and alleviate homelessness.
3. Work to develop a coordinated and seamless service delivery system to prevent and alleviate homelessness.
4. Work to identify causes and effects of homelessness and increase awareness among policymakers and the general public.
5. Advise the governor's office, the Iowa finance authority, state agencies, and private organizations on strategies to prevent and eliminate homelessness.
6. Prepare a point-in-time report on homelessness in Iowa and file the report with the governor and the general assembly on or before the first day of December each year.
7. Assist in the completion of the state's continuum of care application to the U.S. Department of Housing and Urban Development.

[ARC 7704B, IAB 4/8/09, effective 5/13/09; ARC 6955C, IAB 3/22/23, effective 4/26/23]

These rules are intended to implement Iowa Code sections 16.5(1) "r" and 16.2D and 2022 Iowa Acts, House File 2258.

[Filed emergency 12/22/08—published 1/14/09, effective 12/22/08]

[Filed ARC 7704B (Notice ARC 7514B, IAB 1/14/09), IAB 4/8/09, effective 5/13/09]

[Filed ARC 2005C (Notice ARC 1864C, IAB 2/4/15), IAB 5/27/15, effective 7/1/15]

[Filed ARC 4319C (Notice ARC 4196C, IAB 1/2/19), IAB 2/27/19, effective 4/3/19]

[Filed ARC 6955C (Notice ARC 6818C, IAB 1/11/23), IAB 3/22/23, effective 4/26/23]

CHAPTER 47
HOUSING RENEWAL PILOT PROGRAM

265—47.1(89GA, HF2564) Purpose. Pursuant to 2022 Iowa Acts, House File 2564, division III, a housing renewal program fund is established under the control of the Iowa finance authority. The authority shall provide moneys from the fund to a nonprofit Iowa affiliate to award grants under the program. Grantees shall use the funds for the purpose of investing in housing rehabilitation and redevelopment for resale to an income-qualified buyer who occupies the home as the buyer's primary residence.

[ARC 6956C, IAB 3/22/23, effective 4/26/23]

265—47.2(89GA, HF2564) Definitions.

“*Authority*” means the Iowa finance authority created in Iowa Code section 16.1A.

“*Eligible participant*” means a city, a county, a consortium of local governments, or an organization exempt from taxation pursuant to Section 501(c)(3) of the Internal Revenue Code with whom a nonprofit affiliate elects to partner.

“*Grantee*” means an eligible participant awarded a grant under the program.

“*Nonprofit Iowa affiliate*” or “*nonprofit affiliate*” means a nonprofit Iowa affiliate of a nonprofit international organization whose primary activity is the promotion of the construction, remodeling, or rehabilitation of one-family or two-family dwellings for use by low-income families.

“*Redevelopment*” means activities including new construction of housing on vacant or demolished properties on infill lots or the conversion of property from a nonresidential use to housing.

“*Rehabilitation*” means renovation, remodeling and repair of existing housing units for continued residential use.

[ARC 6956C, IAB 3/22/23, effective 4/26/23]

265—47.3(89GA, HF2564) Agreement.

47.3(1) The authority will prepare an agreement between the authority and the nonprofit affiliate. The agreement may include terms and conditions reasonably necessary for implementation of the program pursuant to this chapter and 2022 Iowa Acts, House File 2564.

47.3(2) Any substantive change to the nonprofit affiliate's proposed uses of funds shall require an amendment to the agreement. Amendments shall be requested in writing. No amendment shall be valid until approved by the authority.

47.3(3) The nonprofit affiliate must prepare an agreement for each project approved for an award. The agreement will reflect the terms of the award and may include other terms and conditions reasonably necessary for implementation of the program pursuant to this chapter and 2022 Iowa Acts, House File 2564. The nonprofit affiliate and the grantee must execute an agreement before funds are disbursed by the nonprofit affiliate. The nonprofit affiliate must provide a copy of each agreement executed by the affiliate and a grantee to the authority within 30 days of execution.

[ARC 6956C, IAB 3/22/23, effective 4/26/23]

265—47.4(89GA, HF2564) Reporting.

47.4(1) The nonprofit affiliate must submit a report to the authority on or before November 30 of each year.

47.4(2) In addition to the requirements described in 2022 Iowa Acts, House File 2564, division III, the nonprofit affiliate must report on each of the following:

- a. A description of each grantee's project and grantee's progress toward completion of its projects.
- b. The sale price and sale closing date of each ownership unit.
- c. The income level of each homebuyer purchasing an ownership unit.
- d. The street address, city, zip code and county of each ownership unit.
- e. The amount of funds awarded to each grantee.
- f. The amount of funds expended by each grantee.
- g. The amount of funds obligated by each grantee.

- h.* The amount of funds leveraged by each grantee.
- i.* Any other information reasonably requested by the authority in sufficient detail to permit the authority to prepare any reports required by the authority, the general assembly or the governor's office.

[ARC 6956C, IAB 3/22/23, effective 4/26/23]

These rules are intended to implement 2022 Iowa Acts, House File 2564, division III.

[Filed ARC 6956C (Notice ARC 6819C, IAB 1/11/23), IAB 3/22/23, effective 4/26/23]

COSMETOLOGISTS

CHAPTER 60	LICENSURE OF COSMETOLOGISTS, ELECTROLOGISTS, ESTHETICIANS, MANICURISTS, NAIL TECHNOLOGISTS, AND INSTRUCTORS OF COSMETOLOGY ARTS AND SCIENCES
CHAPTER 61	LICENSURE OF SALONS AND SCHOOLS OF COSMETOLOGY ARTS AND SCIENCES
CHAPTER 62	RESERVED
CHAPTER 63	SANITATION FOR SALONS AND SCHOOLS OF COSMETOLOGY ARTS AND SCIENCES
CHAPTER 64	CONTINUING EDUCATION FOR COSMETOLOGY ARTS AND SCIENCES
CHAPTER 65	DISCIPLINE FOR COSMETOLOGY ARTS AND SCIENCES LICENSEES, INSTRUCTORS, SALONS, AND SCHOOLS

CHAPTER 60

LICENSURE OF COSMETOLOGISTS, ELECTROLOGISTS, ESTHETICIANS,
 MANICURISTS, NAIL TECHNOLOGISTS, AND INSTRUCTORS
 OF COSMETOLOGY ARTS AND SCIENCES

[Prior to 7/29/87, Health Department[470] Chs 149, 150]

645—60.1(157) Definitions. For purposes of these rules, the following definitions shall apply:

“Active license” means a license that is current and has not expired.

“Apprentice” means a person who is at least 16 years of age, who is employed in an apprenticeable occupation, who is a resident of the state of Iowa, and who is registered in Iowa by the Office of Apprenticeship of the United States Department of Labor.

“Apprenticeship program” means a program registered by the Office of Apprenticeship of the United States Department of Labor, which includes terms and conditions for the qualification, recruitment, selection, employment, and training of apprentices, including the requirement for a written apprenticeship agreement between an apprentice and an active licensee in an active licensed salon.

“Board” means the board of cosmetology arts and sciences.

“Certified laser product” means a product which is certified by a manufacturer pursuant to the requirements of 21 Code of Federal Regulations (CFR) Part 1040.

“Chemical exfoliation” means the removal of surface epidermal cells of the skin by using only non-medical-strength cosmetic preparations consistent with labeled instructions and as specified by rule. This procedure is not intended to elicit viable epidermal or dermal wounding, injury, or destruction.

“Core curriculum” means the basic core life sciences curriculum that is required for completion of any course of study of the cosmetology arts and sciences except for manicuring.

“Cosmetology arts and sciences” means any or all of the following disciplines performed with or without compensation by a licensee: cosmetology, electrology, esthetics, nail technology and manicuring.

“Depilatory” means an agent used for the temporary removal of superfluous hair by dissolving it at the epidermal surface.

“Examination” means any of the tests used to determine minimum competency prior to the issuance of a cosmetology arts and sciences license.

“Exfoliation” means the process whereby the superficial epidermal cells are removed from the skin.

“General supervision” means the supervising physician is not onsite for laser procedures or use of an intense pulsed light device for hair removal conducted on minors, but is available for direct communication, either in person or by telephone, radio, radiotelephone, television, or similar means.

“Grace period” means the 30-day period following expiration of a license when the license is still considered to be active. In order to renew a license during the grace period, a licensee is required to pay a late fee.

“Inactive license” means a license that has expired because it was not renewed by the end of the grace period. The category of “inactive license” may include licenses formerly known as lapsed, inactive, delinquent, closed, or retired.

“Intense pulsed light device” means a device that uses incoherent light to destroy the vein of the hair bulb.

“*Laser*” means light amplification by the stimulated emission of radiation.

“*Licensee*” means any person or entity licensed to practice pursuant to Iowa Code chapter 157 and 645—Chapters 60 to 65, Iowa Administrative Code.

“*Licensure by endorsement*” means the issuance of an Iowa license to practice cosmetology to an applicant who is or has been licensed in another state for 12 months during the last 24 months.

“*Mechanical exfoliation*” means the physical removal of surface epidermal cells by means that include but are not limited to brushing machines, granulated scrubs, peel-off masques, peeling creams or drying preparations that are rubbed off, and microdermabrasion.

“*Mentor*” means a licensee providing guidance in a mentoring program.

“*Mentoring*” means a program allowing students to experience cosmetology arts and sciences in a licensed salon under the guidance of a mentor.

“*Microdermabrasion*” means mechanical exfoliation using an abrasive material or apparatus to remove surface epidermal cells with a machine which is specified by rule.

“*Minor*” means an unmarried person who is under the age of 18 years.

“*NIC*” means the National-Interstate Council of State Boards of Cosmetology, Inc.

“*Pedicuring*” means the practice of cleaning, shaping or polishing the toenails.

“*Practice discipline*” means the practice of electrology, esthetics, nail technology, manicuring or cosmetology as recognized by the board of cosmetology arts and sciences.

“*Reactivate*” or “*reactivation*” means the process as outlined in rule 645—60.17(17A,147,272C) by which an inactive license is restored to active status.

“*Reciprocal license*” means the issuance of an Iowa license to practice cosmetology to an applicant who is currently licensed in another state and which state has a mutual agreement to license persons who have the same or similar qualifications to those required in Iowa.

“*Reinstatement*” means the process as outlined in 645—11.31(272C) by which a licensee who has had a license suspended or revoked or who has voluntarily surrendered a license may apply to have the license reinstated, with or without conditions. Once the license is reinstated, the licensee may apply for active status.

“*Testing service*” means a national testing service selected by the board.

[ARC 8515B, IAB 2/10/10, effective 3/17/10; ARC 6376C, IAB 6/29/22, effective 8/3/22]

645—60.2(157) Requirements for licensure.

60.2(1) Requirements for licensure. All persons providing services in one or more cosmetology arts and sciences disciplines shall hold a license issued by the board. The applicant shall:

- a. Submit a completed application for licensure.
- b. Direct the educational program to submit to the board a diploma or an official transcript indicating date of graduation and completion of required hours in each practice discipline for which the applicant is requesting licensure.
- c. If the applicant graduated from a school that is not licensed by the board, direct the school to provide an official transcript showing completion of a course of study that meets the requirements of rule 645—61.14(157).
- d. If the applicant has graduated from an apprenticeship program, the applicant must direct the United States Department of Labor to submit a certificate of completion.
- e. Foreign-trained applicants. If educated outside the United States, attach an original evaluation of the applicant’s education from World Education Services (WES) or any other accredited evaluation service. An applicant may obtain an application for evaluation by contacting WES online at www.wes.org or at (212)966-6311, or by writing to WES, P.O. Box 5087, Bowling Green Station, New York, New York 10274-5087.
- f. Examination requirements. Pass a national examination as prescribed by the board for the particular practice discipline with a score of 75 percent or greater.

(1) The applicant shall submit the test registration fee directly to the test service. NIC examinations are administered according to guidelines set forth by the National-Interstate Council of State Boards of Cosmetology.

(2) If applying for licensure by endorsement, the applicant shall complete the requirements set forth in rule 645—60.7(157).

60.2(2) Requirements for an instructor's license. An applicant for an instructor's license shall:

- a. Submit a completed application for licensure and the appropriate fee to the board;
- b. Be licensed in the state of Iowa in the specific practice discipline to be taught or be licensed as a cosmetologist who possesses the skill and knowledge required to instruct in that practice discipline;
- c. Provide documentation of completion of 1,000 hours of instructor's training or two years' active practice in the field of cosmetology within six years prior to application;
- d. Submit proof of completion of an instructor methods training course consisting of at least 16 hours;
- e. Submit proof of 60 hours of practical experience, excluding school hours, in the area of electrolysis prior to application for an instructor of electrolysis license.
- f. Pass an instructor's national examination, which, effective January 1, 2008, shall be the NIC instructor examination unless the applicant is applying for an instructor's license by endorsement as outlined in rule 645—60.7(157).

60.2(3) Conditions. The following conditions apply for all cosmetology arts and sciences licenses.

- a. Incomplete applications that have been on file in the board office for more than two years shall be considered invalid and shall be destroyed.
- b. The licensure fee is nonrefundable.
- c. Licensees who were issued their initial licenses within six months prior to the license renewal beginning date shall not be required to renew their licenses until the renewal month two years later.
- d. A new license granted by the board of cosmetology arts and sciences to an individual who holds multiple active licenses with the board shall have the same license expiration date as the licensee's existing license(s). If the licensee holds only one active license with the board, the license expiration date shall be in the current renewal period unless licensure is issued within six months of the end of the renewal cycle, in which case subrule 60.8(2) shall apply.

60.2(4) Licensure by work experience. An applicant who has relocated to Iowa from a state that did not require licensure to practice the profession may submit proof of work experience in lieu of educational and training requirements, if eligible, in accordance with rule 645—19.2(272C).

[ARC 3558C, IAB 1/3/18, effective 2/7/18; ARC 5755C, IAB 7/14/21, effective 8/18/21; ARC 6376C, IAB 6/29/22, effective 8/3/22; ARC 6952C, IAB 3/22/23, effective 4/26/23]

645—60.3(157) Criteria for licensure in specific practice disciplines.

60.3(1) A cosmetology license is not a requirement for an electrolysis, esthetics, nail technology or manicurist license.

60.3(2) Core life sciences curriculum hours shall be transferable in their entirety from one practice discipline to another practice discipline.

60.3(3) Theory hours earned in each practice discipline of cosmetology arts and sciences may be used in applying for a cosmetology license.

60.3(4) A cosmetologist licensed after July 1, 2005, is not eligible to be certified in chemical peels, microdermabrasion, laser or intense pulsed light (IPL) and shall not provide those services.

60.3(5) Pedicuring shall only be done by a cosmetologist or nail technologist.

60.3(6) Facial waxing shall only be done by a cosmetologist or esthetician.

60.3(7) An initial license to practice manicuring shall not be issued by the board after December 31, 2007. A manicurist license issued on or before December 31, 2007, may be renewed subject to licensure requirements identified by statute and administrative rule unless the license becomes inactive. A manicurist license that becomes inactive cannot be reactivated or renewed.

645—60.4(157) Practice-specific training requirements. The board shall approve a licensee to provide the appropriate services once a licensee has complied with training requirements and submitted a completed application, the required supporting evidence, and applicable fees as specified in these rules. The applicant shall receive a certification following board approval.

60.4(1) Microdermabrasion.

a. Microdermabrasion shall only be performed by a licensed, certified esthetician or a cosmetologist who was licensed prior to July 1, 2005, and is certified by the board.

b. To be eligible to perform microdermabrasion services, the licensee shall:

(1) Complete 14 contact hours of education specific to the material or apparatus used for microdermabrasion. Before an additional material or apparatus is utilized in the licensee's practice, the licensee shall provide official certification of training on the material or apparatus.

(2) Obtain from the program a certification of training that contains the following information:

1. Date, location, course title;
2. Number of contact hours; and
3. Specific identifying description of the microdermabrasion machine covered by the course.

(3) Complete a board-approved certification application form and submit to the board office the completed form, a copy of the certification of training, and the required fee pursuant to 645—subrule 5.5(14). The fee is nonrefundable.

60.4(2) Chemical exfoliation.

a. Chemical exfoliation shall only be performed by a cosmetologist who was licensed prior to July 1, 2005, and is certified by the board to perform those services. Additional certification is not required for licensed estheticians.

b. Chemical exfoliation procedures are limited to the removal of surface epidermal cells of the skin by using only non-medical-strength cosmetic preparations consistent with labeled instructions and as specified by these rules. This procedure is not intended to elicit viable epidermal or dermal wounding, injury, or destruction.

c. To be eligible to perform chemical peels, a cosmetologist who was licensed prior to July 1, 2005, shall:

(1) Complete 21 hours of training specific to the process and products to be used for chemical peels. Before an additional process or product is utilized in the licensee's practice, the licensee shall provide official certification of training on the new process or product.

(2) Obtain from the program a certification of training that contains the following information:

1. Date, location, course title;
2. Number of contact hours; and
3. Specific identifying description of the chemical peel process and products covered by the course.

(3) Complete a board-approved certification application form and submit to the board office the completed form, a copy of the certification of training, and the required fee pursuant to 645—subrule 5.5(15). The fee is nonrefundable.

60.4(3) Laser services.

a. A cosmetologist licensed after July 1, 2005, shall not use laser products.

b. An electrologist shall only provide hair removal services when using a laser.

c. Estheticians and cosmetologists shall use laser for cosmetic purposes only.

d. Cosmetologists licensed prior to July 1, 2005, electrologists and estheticians must be certified to perform laser services.

e. When a laser service is provided to a minor by a licensed cosmetologist, esthetician or electrologist who has been certified by the board, the licensee shall work under the general supervision of a physician. The parent or guardian shall sign a consent form prior to services being provided. Written permission shall remain in the client's permanent record for a period of five years.

f. To be eligible to perform laser services, a cosmetologist who was licensed on or before July 1, 2005, an electrologist, or an esthetician shall:

(1) Complete 40 hours of training specific to each laser machine, model or device to be used for laser services. Before an additional machine, model or device is utilized in the licensee's practice, the licensee shall submit official certification of training on the new machine, model or device.

(2) Obtain from the program a certification of training that contains the following information:

1. Date, location, course title;
2. Number of contact hours;

3. Specific identifying description of the laser equipment; and
4. Evidence that the training program includes a safety training component which provides a thorough understanding of the procedures to be performed. The training program shall address fundamentals of nonbeam hazards, management and employee responsibilities relating to control measures, and regulatory requirements.

(3) Complete a board-approved certification application form and submit to the board office the completed form, a copy of the certification of training, and the required fee pursuant to 645—subrule 5.5(14). The fee is nonrefundable.

60.4(4) IPL hair removal treatments.

- a. A cosmetologist licensed after July 1, 2005, shall not use IPL devices.
- b. An IPL device shall only be used for hair removal.
- c. Cosmetologists licensed prior to July 1, 2005, electrologists and estheticians must be certified to perform IPL services.
- d. When IPL hair removal services are provided to a minor by a licensed cosmetologist, esthetician or electrologist who has been certified by the board, the licensee shall work under the general supervision of a physician. The parent or guardian shall sign a consent form prior to services being provided. Written permission shall remain in the client's permanent record for a period of five years.

e. To be eligible to perform IPL hair removal services, a cosmetologist who was licensed on or before July 1, 2005, an electrologist, or an esthetician shall:

(1) Complete 40 hours of training specific to each IPL machine, model or device to be used for IPL hair removal services. Before an additional machine, model or device is utilized in the licensee's practice, the licensee shall submit official certification of training on the new machine, model or device.

(2) Obtain from the program a certification of training that contains the following information:

1. Date, location, course title;
2. Number of contact hours;
3. Specific identifying description of the IPL hair removal equipment; and
4. Evidence that the training program includes a safety training component which provides a thorough understanding of the procedures to be performed. The training program shall address fundamentals of nonbeam hazards, management and employee responsibilities relating to control measures, and regulatory requirements.

(3) Complete a board-approved certification application form and submit to the board office the completed form, a copy of the certification of training, and the required fee pursuant to 645—subrule 5.5(14). The fee is nonrefundable.

60.4(5) Health history and incident reporting.

a. Prior to providing laser or IPL hair removal, microdermabrasion or chemical peel services, the cosmetologist, esthetician, and electrologist shall complete a client health history of conditions related to the application for services and include it with the client's records. The history shall include but is not limited to items listed in paragraph 60.4(5) "b."

b. A licensed cosmetologist, esthetician, or electrologist who provides services related to the use of a certified laser product, IPL device, chemical peel, or microdermabrasion shall submit a report to the board within 30 days of any incident in which provision of such services resulted in physical injury requiring medical attention. Failure to comply with this requirement shall result in disciplinary action by the board. The report shall include the following:

- (1) A description of procedures;
- (2) A description of the physical condition of the client;
- (3) A description of any adverse occurrence, including:
 1. Symptoms of any complications including, but not limited to, onset and type of symptoms;
 2. A description of the services provided that caused the adverse occurrence;
 3. A description of the procedure that was followed by the licensee;
- (4) A description of the client's condition on termination of any procedures undertaken;
- (5) If a client is referred to a physician, a statement providing the physician's name and office location, if known;

(6) A copy of the consent form.

60.4(6) Failure to report. Failure to comply with paragraph 60.4(5) “b” when the adverse occurrence is related to the use of any procedure or device noted in the attestation may result in the licensee’s loss of authorization to administer the procedure or device noted in the attestation or may result in other sanctions provided by law.

60.4(7) A licensee shall not provide any services that constitute the practice of medicine.
[ARC 5755C, IAB 7/14/21, effective 8/18/21]

645—60.5(157) Licensure restrictions relating to practice.

60.5(1) A certified laser product or an intense pulsed light device shall only be used on surface epidermal layers of the skin except for hair removal.

60.5(2) A laser hair removal product or an intense pulsed light device shall not be used on a minor unless the minor is accompanied by a parent or guardian and then shall be used only under general supervision of a physician.

60.5(3) Persons licensed under Iowa Code chapter 157 shall not administer any practice of removing skin by means of a razor-edged instrument.

60.5(4) Persons licensed under this chapter who provide hair removal, manicuring and nail technology services shall not administer any procedure in which human tissue is cut, shaped, vaporized, or otherwise structurally altered, except for the use of a cuticle nipper.

60.5(5) Board-certified licensees providing microdermabrasion, chemical peels, laser or IPL hair removal treatments in a salon or barbershop setting shall not include any practice, activity, or treatment that constitutes the practice of medicine, osteopathic medicine, chiropractic or acupuncture.

60.5(6) Cosmetologists licensed prior to July 1, 2005, and licensed estheticians shall only perform medical aesthetic services in a medical spa under the delegation and supervision of a medical director as set forth by the Iowa board of medicine in rule 653—13.8(148,272C). The Iowa board of cosmetology arts and sciences does not license medical aestheticians.

60.5(7) Persons licensed under this chapter who provide apprenticeship programs must hold an active license sufficient to provide on-the-job training, must operate in an actively licensed establishment, and must comply with relevant United States Department of Labor laws and regulations for the operation of an apprenticeship program.
[ARC 2599C, IAB 6/22/16, effective 8/15/16; ARC 6376C, IAB 6/29/22, effective 8/3/22]

645—60.6(157) Consent form requirements. A licensed esthetician, cosmetologist, or electrologist, prior to providing services relating to a certified laser product, intense pulsed light device, chemical peel, or microdermabrasion, shall obtain from a client a consent form that:

1. Specifies in general terms the nature and purpose of the procedure(s);
2. Lists known risks associated with the procedure(s) if reasonably determinable;
3. States an acknowledgment that disclosure of information has been made and that questions asked about the procedure(s) have been satisfactorily answered;
4. Includes a signature of either the client for whom the procedure is performed or, if that client for any reason lacks legal capacity to consent, includes the signature of a person who has legal authority to consent on behalf of that client in those circumstances.

645—60.7(157) Licensure by endorsement. The board may issue a license by endorsement to any applicant from the District of Columbia or another state, territory, province or foreign country who has held an active license under the laws of another jurisdiction for at least 12 months during the past 24 months.

60.7(1) Applicants shall submit to the board a completed application and pay the licensure fee specified in 645—subrule 5.5(1).

60.7(2) Applicants shall provide verification of license in a cosmetology practice discipline from the jurisdiction in which the applicant has most recently been licensed, sent directly from the jurisdiction to the board office. The applicant must also disclose any public or pending complaints against the applicant

in any other jurisdiction. Web-based verification may be substituted for verification from a jurisdiction's board office if the verification includes:

- a. Licensee's name;
- b. Date of initial licensure;
- c. Current licensure status; and
- d. Any disciplinary action taken against the license.

60.7(3) Applicants who graduated from a cosmetology school prior to January 1, 2000, shall have passed the state written and practical examination required by the state in which the applicants were originally licensed.

60.7(4) Applicants who graduated from a cosmetology school after January 1, 2000, shall have passed a national theory examination.

60.7(5) Licensure by verification. A person who is licensed in another jurisdiction but who is unable to satisfy the requirements for licensure by endorsement may apply for licensure by verification, if eligible, in accordance with rule 645—19.1(272C).

[ARC 8515B, IAB 2/10/10, effective 3/17/10; ARC 5755C, IAB 7/14/21, effective 8/18/21; ARC 6952C, IAB 3/22/23, effective 4/26/23]

645—60.8(157) License renewal.

60.8(1) Biennial license renewal period for a license to practice cosmetology arts and sciences.

a. The renewal period shall begin on April 1 of one year and end on March 31 two years later. All licensees shall renew on a biennial basis.

b. The board may send a renewal notice by regular mail to each licensee at the address on record prior to the expiration of the license.

c. The licensee is responsible for renewing the license prior to its expiration. Failure of the licensee to receive the notice does not relieve the licensee of the responsibility for renewing the license.

d. A new or reactivated license granted by the board to a licensee who holds a current license in another practice discipline in cosmetology shall have the same license expiration date as the licensee's other license(s). If the licensee does not have another active license with the board, the license expiration date shall be in the current renewal period unless the license is issued within six months of the end of the renewal cycle and subrule 60.8(2) applies.

60.8(2) An individual who was issued a license within six months of the license renewal date will not be required to renew the license until the subsequent renewal two years later.

60.8(3) License renewal. A licensee seeking renewal shall:

a. Meet the continuing education requirements of rule 645—64.2(157). A licensee whose license was reactivated during the current renewal compliance period may use continuing education credit earned during the compliance period for the first renewal following reactivation; and

b. Submit the completed renewal application and renewal fee before the license expiration date.

c. Licensees currently licensed in Iowa but practicing exclusively in another state or serving honorably as active duty military or the spouse of active duty military service personnel may comply with Iowa continuing education requirements for license renewal by meeting the continuing education requirements of the state where the licensee practices. Those licensees living and practicing exclusively in a state which has no continuing education requirement for renewal of a license shall not be required to meet Iowa's continuing education requirement but shall pay all renewal fees when due.

60.8(4) Upon receiving the information required by this rule and the required fee, board staff shall administratively issue a two-year license. In the event the board receives adverse information on the renewal application, the board shall issue the renewal license but may refer the adverse information for further consideration or disciplinary investigation.

60.8(5) Late renewal. The license shall become late when the license has not been renewed by the expiration date on the renewal. The licensee shall be assessed a late fee as specified in 645—subrule 5.5(3). To renew a late license, the licensee shall complete the renewal requirements and submit the late fee within the grace period.

60.8(6) Inactive license. A licensee who fails to renew the license by the end of the grace period has an inactive license. A licensee whose license is inactive continues to hold the privilege of licensure in Iowa, but may not practice cosmetology arts and sciences in Iowa until the license is reactivated. A licensee who practices cosmetology arts and sciences in the state of Iowa with an inactive license may be subject to disciplinary action by the board, injunctive action pursuant to Iowa Code section 147.83, criminal sanctions pursuant to Iowa Code section 147.86, and other available legal remedies.

60.8(7) Those persons licensed for the first time shall not be required to complete continuing education as a prerequisite for the first renewal of their licenses. Continuing education hours acquired anytime from the initial licensing until the second license renewal may be used.

[ARC 3558C, IAB 1/3/18, effective 2/7/18; ARC 5755C, IAB 7/14/21, effective 8/18/21]

645—60.9 to 60.16 Reserved.

645—60.17(17A,147,272C) License reactivation. To apply for reactivation of an inactive license, a licensee shall:

60.17(1) Submit a reactivation application on a form provided by the board.

60.17(2) Pay the reactivation fee that is due as specified in rule 645—5.5(147,157).

60.17(3) Provide verification of current competence to practice cosmetology arts and sciences by satisfying one of the following criteria:

a. If the license has been on inactive status for five years or less, an applicant must provide the following:

(1) Verification of the license from the jurisdiction in which the applicant has most recently been practicing during the time period the Iowa license was inactive, sent directly from the jurisdiction to the board office. The applicant must also disclose any public or pending complaints against the applicant in any other jurisdiction. Web-based verification may be substituted for verification from a jurisdiction's board office if the verification includes:

1. Licensee's name;
2. Date of initial licensure;
3. Current licensure status; and
4. Any disciplinary action taken against the license; and

(2) Verification of completion of 6 hours of continuing education that meet the continuing education standards defined in rule 645—64.3(157,272C) within two years of application for reactivation; or verification of active practice, consisting of a minimum of 2,080 hours, in another state or jurisdiction during the two years preceding an application for reactivation.

b. If the license has been on inactive status for more than five years, an applicant must provide the following:

(1) Verification of the license from the jurisdiction in which the applicant has most recently been practicing during the time period the Iowa license was inactive, sent directly from the jurisdiction to the board office. The applicant must also disclose any public or pending complaints against the applicant in any other jurisdiction. Web-based verification may be substituted for verification from a jurisdiction's board office if the verification includes:

1. Licensee's name;
2. Date of initial licensure;
3. Current licensure status; and
4. Any disciplinary action taken against the license; and

(2) Verification of completion of 12 hours of continuing education that meet the continuing education standards defined in rule 645—64.3(157,272C) within two years of application for reactivation; or verification of active practice, consisting of a minimum of 2,080 hours, in another state or jurisdiction during the two years preceding an application for reactivation.

60.17(4) Licensees who are instructors of cosmetology arts and sciences shall obtain an additional 6 hours of continuing education in teaching methodology as prescribed in 645—Chapter 64.

60.17(5) Submit a sworn statement of previous cosmetology arts and sciences practice from an employer or professional associate, detailing places and dates of employment and verifying that the applicant has practiced cosmetology arts and sciences at least 2,080 hours or taught as the equivalent of a full-time faculty member for at least one of the immediately preceding years during the last two-year time period. Sole proprietors may submit the sworn statement on their own behalf.
 [ARC 3558C, IAB 1/3/18, effective 2/7/18; ARC 5755C, IAB 7/14/21, effective 8/18/21; ARC 6952C, IAB 3/22/23, effective 4/26/23]

645—60.18(17A,147,272C) License reinstatement. A licensee whose license has been revoked, suspended, or voluntarily surrendered must apply for and receive reinstatement of the license in accordance with 645—11.31(272C) and must apply for and be granted reactivation of the license in accordance with 645—60.17(17A,147,272C) prior to practicing cosmetology arts and sciences in this state.

These rules are intended to implement Iowa Code chapters 272C and 157.

[Filed prior to 7/1/52; amended 4/21/53, 5/15/53, 10/1/59, 4/19/71]

[Filed 8/5/77, Notice 6/1/77—published 8/24/77, effective 10/1/77]

[Filed 4/28/78, Notice 12/28/77—published 5/17/78, effective 6/21/78]

[Filed 10/19/79, Notice 8/22/79—published 11/14/79, effective 12/21/79]

[Filed 2/27/81, Notice 12/10/80—published 3/18/81, effective 4/22/81]

[Filed 11/15/82, Notice 9/1/82—published 12/8/82, effective 1/15/83]

[Filed 10/6/83, Notice 7/20/83—published 10/26/83, effective 11/30/83]

[Filed 4/15/85, Notice 2/27/85—published 5/8/85, effective 6/12/85]

[Filed 8/5/85, Notice 6/5/85—published 8/28/85, effective 10/2/85]

[Filed emergency 7/10/87—published 7/29/87, effective 7/10/87]

[Filed 4/29/88, Notice 3/23/88—published 5/18/88, effective 6/22/88]

[Filed 8/4/89, Notice 6/14/89—published 8/23/89, effective 9/27/89]

[Filed 9/29/89, Notice 8/23/89—published 10/18/89, effective 11/22/89]

[Filed 2/2/90, Notice 12/27/89—published 2/21/90, effective 3/28/90]

[Filed 9/27/91, Notice 6/12/91—published 10/16/91, effective 11/20/91]

[Filed 1/3/92, Notice 9/4/91—published 1/22/92, effective 2/26/92]¹

[Filed 12/4/92, Notice 8/5/92—published 12/23/92, effective 1/29/93]

[Filed 2/11/94, Notice 10/27/93—published 3/2/94, effective 4/6/94]

[Filed 4/19/95, Notice 2/1/95—published 5/10/95, effective 6/14/95]

[Filed 11/2/95, Notice 9/13/95—published 11/22/95, effective 12/27/95]

[Filed 11/15/96, Notice 9/11/96—published 12/4/96, effective 1/8/97]

[Filed 2/6/98, Notice 11/19/97—published 2/25/98, effective 4/1/98]

[Filed 2/19/99, Notice 12/2/98—published 3/10/99, effective 4/14/99]

[Filed 5/28/99, Notice 1/27/99—published 6/16/99, effective 7/21/99]

[Filed 2/1/01, Notice 11/29/00—published 2/21/01, effective 3/28/01]

[Filed 2/13/02, Notice 11/28/01—published 3/6/02, effective 4/10/02]

[Filed 8/14/02, Notice 5/29/02—published 9/4/02, effective 10/9/02]

[Filed 2/12/03, Notice 12/25/02—published 3/5/03, effective 4/9/03]

[Filed 8/14/03, Notice 5/28/03—published 9/3/03, effective 10/8/03]

[Filed 2/10/04, Notice 11/26/03—published 3/3/04, effective 4/7/04]

[Filed 2/3/05, Notice 11/24/04—published 3/2/05, effective 4/6/05]

[Filed 8/5/05, Notice 5/25/05—published 8/31/05, effective 10/5/05]

[Filed 11/4/05, Notice 9/14/05—published 11/23/05, effective 12/28/05][◇]

[Filed 11/4/05, Notice 9/28/05—published 11/23/05, effective 12/28/05][◇]

[Filed 2/1/06, Notice 12/7/05—published 3/1/06, effective 4/5/06]

[Filed without Notice 8/22/07—published 9/12/07, effective 1/1/08]

[Filed 10/24/07, Notice 9/12/07—published 11/21/07, effective 1/1/08]

[Filed 12/5/08, Notice 10/8/08—published 12/31/08, effective 2/4/09]

[Filed ARC 8515B (Notice ARC 8330B, IAB 12/2/09), IAB 2/10/10, effective 3/17/10]

[Filed ARC 2599C (Notice ARC 2467C, IAB 3/16/16), IAB 6/22/16, effective 8/15/16]
[Filed ARC 3558C (Notice ARC 3372C, IAB 10/11/17), IAB 1/3/18, effective 2/7/18]
[Filed ARC 5755C (Notice ARC 5455C, IAB 2/24/21), IAB 7/14/21, effective 8/18/21]
[Filed ARC 6376C (Notice ARC 6258C, IAB 3/23/22), IAB 6/29/22, effective 8/3/22]
[Filed ARC 6952C (Notice ARC 6662C, IAB 11/16/22), IAB 3/22/23, effective 4/26/23]

◊ Two or more ARCs

¹ Effective date of 2/26/92 delayed until adjournment of the 1992 General Assembly by the Administrative Rules Review Committee at its meeting held February 3, 1992.

CHAPTER 64
CONTINUING EDUCATION FOR COSMETOLOGY ARTS AND SCIENCES

[Prior to 7/29/87, Health Department[470] Ch 151]

[Prior to 12/23/92, see 645—Chapter 62]

645—64.1(157) Definitions. For the purpose of these rules, the following definitions shall apply:

“*Active license*” means a license that is current and has not expired.

“*Approved program/activity*” means a continuing education program/activity meeting the standards set forth in these rules.

“*Audit*” means the selection of licensees for verification of satisfactory completion of continuing education requirements during a specified time period.

“*Board*” means the board of cosmetology arts and sciences.

“*Continuing education*” means planned, organized learning acts acquired during licensure designed to maintain, improve, or expand a licensee’s knowledge and skills in order for the licensee to develop new knowledge and skills relevant to the enhancement of practice, education, or theory development to improve the safety and welfare of the public.

“*Hour of continuing education*” means at least 50 minutes spent by a licensee completing an approved continuing education activity through live, virtual, online or prerecorded means where the instructor provides proof of completion by the licensee as set forth in these rules.

“*Inactive license*” means a license that has expired because it was not renewed by the end of the grace period. The category of “inactive license” may include licenses formerly known as lapsed, inactive, delinquent, closed, or retired.

“*Independent study*” means a subject/program/activity that a person pursues autonomously that meets standards for approval criteria in the rules.

“*License*” means license to practice.

“*Licensee*” means any person or entity licensed to practice pursuant to Iowa Code chapter 157 and 645—Chapters 60 to 65, Iowa Administrative Code.

“*Prescribed practice*” means an area of specialty within the scope of cosmetology arts and sciences. [ARC 3558C, IAB 1/3/18, effective 2/7/18; ARC 6952C, IAB 3/22/23, effective 4/26/23]

645—64.2(157) Continuing education requirements.

64.2(1) The biennial continuing education compliance period shall begin on April 1 of one year and end on March 31 two years later.

64.2(2) Each biennium:

a. A licensee in this state shall be required to complete a minimum of 6 hours of continuing education that meets the requirements of rule 645—64.3(157,272C). A minimum of 4 hours of the 6 hours shall be in the prescribed practice discipline and a minimum of 2 hours of the 6 hours shall be in the content areas of Iowa cosmetology law and rules and sanitation. Individuals holding more than one active license shall obtain 4 hours of continuing education in each prescribed practice discipline and an additional 2 hours in the content areas of Iowa cosmetology law and rules and sanitation.

b. A licensee who is an instructor of cosmetology arts and sciences shall obtain 6 hours in teaching methodology in addition to meeting all continuing education requirements for renewal of the instructor’s practice license. A licensee must comply with all conditions of licensure including obtaining a minimum of 2 hours each biennium specific to Iowa cosmetology law and administrative rules as specified in subrule 64.3(2).

c. A licensee currently licensed in Iowa but practicing exclusively in another state may comply with Iowa continuing education requirements for license renewal by meeting the continuing education requirements of the state or states where the licensee practices. The licensee living and practicing in a state which has no continuing education requirement for renewal of a license shall not be required to meet Iowa’s continuing education requirement but shall pay all renewal fees when due.

d. A licensee shall be deemed to have complied with the continuing education requirements of this state during periods that the licensee:

(1) Serves honorably on active duty in the military services, or

- (2) Is the spouse of an active duty military service person, or
- (3) Is a government employee working in the person's licensed specialty and assigned to duty outside of the United States, or
- (4) Is engaged in active practice and absence from the state approved by the board.

64.2(3) Requirements of new licensees. Those persons licensed for the first time shall not be required to complete continuing education as a prerequisite for the first renewal of their licenses. Continuing education hours acquired anytime from the initial licensing until the second license renewal may be used.

64.2(4) Hours of continuing education credit may be obtained by attending and participating in a continuing education activity. These hours must be in accordance with these rules.

64.2(5) No hours of continuing education shall be carried over into the next biennium. A licensee whose license was reactivated during the current renewal compliance period may use continuing education earned during the compliance period for the first renewal following reactivation.

64.2(6) It is the responsibility of each licensee to finance the cost of continuing education.
[ARC 8515B, IAB 2/10/10, effective 3/17/10; ARC 3558C, IAB 1/3/18, effective 2/7/18]

645—64.3(157,272C) Standards.

64.3(1) *General criteria.* A continuing education activity which meets all of the following criteria is appropriate for continuing education credit if the continuing education activity:

- a. Constitutes an organized program of learning which contributes directly to the professional competency of the licensee;
- b. Pertains to subject matters which integrally relate to the practice of the profession;
- c. Is conducted by individuals who have specialized education, training and experience by reason of which said individuals should be considered qualified concerning the subject matter of the program. At the time of audit, the board may request the qualifications of presenters;
- d. Fulfills stated program goals, objectives, or both; and
- e. Provides proof of attendance to licensees in attendance including:
 - (1) Date, location, course title, presenter(s), sponsor(s);
 - (2) Number of program contact hours; and
 - (3) Certificate of completion or evidence of successful completion of the course provided by the course sponsor.

64.3(2) *Specific criteria.* A licensee shall obtain a minimum of 6 hours of continuing education credit every two years. A minimum of 4 hours of the 6 hours of continuing education shall be in each prescribed practice discipline. Two hours of continuing education per biennium must be specific to Iowa cosmetology law and administrative rules including infection control.

- a. The licensee may obtain continuing education hours of credit by:
 - (1) Attending workshops, conferences or symposiums.
 - (2) Accessing online training, such as viewing interactive conferences, attending webinars, or completing online training courses.
 - (3) Attending programs on product knowledge, methods and systems. Continuing education shall be directly related to the technique and theory specific to the practice of cosmetology arts and sciences. No direct selling of products is allowed as part of a continuing education offering.
 - (4) Attending business classes specific to owning or managing a salon are acceptable.
- b. In addition to fulfilling the requirements in rule 645—64.2(157), those persons holding an instructor's license must complete a minimum of 6 hours of continuing education approved by the board in the area of teaching methodology.
- c. Two hours of continuing education per biennium must be specific to Iowa cosmetology law and administrative rules.
- d. The licensee shall obtain at least 4 hours in each area of prescribed practice for each cosmetology arts and sciences license held.

64.3(3) *Specific criteria for providers and sponsors of continuing education.*

a. Continuing education shall be obtained by attending programs that meet the criteria in subrule 64.3(1). Individuals or groups may offer continuing education programs that meet the criteria in rule 645—64.3(157,272C) offered by or with express sponsorship in advance of delivery by the following organization(s).

- (1) National, state or local associations of cosmetology arts and sciences;
- (2) Schools and institutes of cosmetology arts and sciences;
- (3) Universities, colleges or community colleges;
- (4) National, state or local associations of barbers;
- (5) Barber schools or institutes;
- (6) Manufacturers of laser or microdermabrasion products;
- (7) Institutes of laser technology.

b. A licensee who is a presenter of a continuing education program that meets the criteria in rule 645—64.3(157,272C) may receive credit once per biennium for the initial presentation of the program. The presenter may receive the same number of hours granted the attendees.

[ARC 8515B, IAB 2/10/10, effective 3/17/10; ARC 3558C, IAB 1/3/18, effective 2/7/18]

645—64.4(157,272C) Audit of continuing education report. Rescinded IAB 12/31/08, effective 2/4/09.

645—64.5(157,272C) Automatic exemption. Rescinded IAB 12/31/08, effective 2/4/09.

645—64.6(157,272C) Grounds for disciplinary action. Rescinded IAB 12/31/08, effective 2/4/09.

645—64.7(157,272C) Continuing education waiver for active practitioners. Rescinded IAB 8/31/05, effective 10/5/05.

645—64.8(157,272C) Continuing education exemption for inactive practitioners. Rescinded IAB 8/31/05, effective 10/5/05.

645—64.9(157,272C) Continuing education exemption for disability or illness. Rescinded IAB 12/31/08, effective 2/4/09.

645—64.10(157,272C) Reinstatement of inactive practitioners. Rescinded IAB 8/31/05, effective 10/5/05.

645—64.11(272C) Hearings. Rescinded IAB 8/31/05, effective 10/5/05.

These rules are intended to implement Iowa Code section 272C.2 and chapter 157.

[Filed 6/20/78, Notice 5/3/78—published 7/12/78, effective 8/16/78]

[Filed 8/3/79, Notice 6/27/79—published 8/22/79, effective 9/26/79]

[Filed 2/12/82, Notice 12/23/81—published 3/3/82, effective 4/9/82]

[Filed 10/6/83, Notice 7/20/83—published 10/26/83, effective 11/30/83]

[Filed emergency 8/31/84—published 9/26/84, effective 8/31/84]

[Filed 10/4/85, Notice 8/28/85—published 10/23/85, effective 11/27/85]

[Filed emergency 7/10/87—published 7/29/87, effective 7/10/87]

[Filed 5/25/89, Notice 4/5/89—published 6/14/89, effective 7/19/89]

[Filed 8/4/89, Notice 6/14/89—published 8/23/89, effective 9/27/89]

[Filed 2/2/90, Notice 12/27/89—published 2/21/90, effective 3/28/90]

[Filed 12/4/92, Notice 8/5/92—published 12/23/92, effective 1/29/93]

[Filed 2/11/94, Notice 10/27/93—published 3/2/94, effective 4/6/94]

[Filed 4/19/95, Notice 2/1/95—published 5/10/95, effective 6/14/95]

[Filed 11/2/95, Notice 9/13/95—published 11/22/95, effective 12/27/95]

[Filed 11/15/96, Notice 9/11/96—published 12/4/96, effective 1/8/97]

[Filed 2/19/99, Notice 12/2/98—published 3/10/99, effective 4/14/99]

[Filed 2/1/01, Notice 11/29/00—published 2/21/01, effective 3/28/01]
[Filed 2/13/02, Notice 11/28/01—published 3/6/02, effective 4/10/02]
[Filed 8/5/05, Notice 5/25/05—published 8/31/05, effective 10/5/05][◇]
[Filed 11/4/05, Notice 9/28/05—published 11/23/05, effective 12/28/05]
[Filed 1/11/07, Notice 11/22/06—published 1/31/07, effective 3/7/07]
[Filed 10/24/07, Notice 9/12/07—published 11/21/07, effective 1/1/08]
[Filed 12/5/08, Notice 10/8/08—published 12/31/08, effective 2/4/09]
[Filed ARC 8515B (Notice ARC 8330B, IAB 12/2/09), IAB 2/10/10, effective 3/17/10]
[Filed ARC 3558C (Notice ARC 3372C, IAB 10/11/17), IAB 1/3/18, effective 2/7/18]
[Filed ARC 6952C (Notice ARC 6662C, IAB 11/16/22), IAB 3/22/23, effective 4/26/23]

[◇] Two or more ARCs

PHARMACY BOARD[657]

[Prior to 2/10/88, see Pharmacy Examiners, Board of [620], renamed Pharmacy Examiners Board[657]
under the “umbrella” of Public Health Department by 1986 Iowa Acts, ch 1245; renamed by 2007 Iowa Acts, Senate File 74]

CHAPTER 1

PURPOSE AND ORGANIZATION

- 1.1(17A) Board mission
- 1.2(17A,147,155A,272C) Description and organization of board
- 1.3(17A,272C) Responsibilities
- 1.4(17A,272C) Submission of complaints and requests
- 1.5(17A,21) Meetings
- 1.6(124,147,155A) Fee for returned check
- 1.7(124,124B,147,155A) Overpayment of fees
- 1.8(155A) Alternate board members

CHAPTER 2

PHARMACIST LICENSES

- 2.1(147,155A) Purpose and scope
- 2.2(147,155A) Definitions
- 2.3(147,155A) License and criminal history record check required
- 2.4(147,155A) Licensure by examination
- 2.5(155A) Application for examination—requirements
- 2.6(155A) Internship requirements
- 2.7(147) Reexamination applications and fees
- 2.8(155A) Licensure by score transfer
- 2.9(147,155A) Licensure by license transfer
- 2.10(155A) Foreign pharmacy graduates
- 2.11(147,155A) License expiration and renewal
- 2.12(272C) Continuing education requirements
- 2.13(272C) Active and inactive license status
- 2.14(147,155A) Fees for additional license certificates and verification
- 2.15(155A) Notifications to the board
- 2.16(235B,272C) Mandatory training for identifying and reporting abuse
- 2.17(272C) Continuing professional development portfolio
- 2.18(147,155A) Temporary license

CHAPTER 3

PHARMACY TECHNICIANS

- 3.1(155A) Definitions
- 3.2(155A) Purpose of registration
- 3.3(155A) Registration required
- 3.4 Reserved
- 3.5(155A) Certification of pharmacy technicians
- 3.6 and 3.7 Reserved
- 3.8(155A) Application
- 3.9(155A) Registration fee and term—technician trainee
- 3.10(155A) Registration fee, term, and renewal—certified pharmacy technician
- 3.11(155A) Verification fee
- 3.12 Reserved
- 3.13(155A) Notifications to the board
- 3.14 to 3.16 Reserved
- 3.17(155A) Training and utilization of pharmacy technicians

3.18(147,155A)	Identification of pharmacy technician
3.19	Reserved
3.20(155A)	Responsibility of supervising pharmacist
3.21(155A)	Delegation of functions
3.22	Reserved
3.23(155A)	Functions a pharmacy technician shall not perform
3.24 to 3.27	Reserved
3.28(147,155A)	Unethical conduct or practice
3.29(155A)	Denial of registration
3.30(155A)	Reporting discipline and criminal convictions
3.31(155A)	Discipline of pharmacy technicians

CHAPTER 4 PHARMACIST-INTERNS

4.1(155A)	Definitions
4.2(155A)	Goal and objectives of internship
4.3(155A)	1500-hour requirements
4.4(155A)	Iowa colleges of pharmacy clinical internship programs
4.5(155A)	Out-of-state internship programs
4.6(155A)	Registration, reporting, and authorized functions
4.7(155A)	Foreign pharmacy graduates
4.8(155A)	Fees
4.9(155A)	Preceptor requirements
4.10(155A)	Denial of pharmacist-intern registration
4.11(155A)	Notifications to the board
4.12(155A)	Discipline of pharmacist-interns

CHAPTER 5 PHARMACY SUPPORT PERSONS

5.1(155A)	Definitions
5.2(155A)	Purpose of registration
5.3	Reserved
5.4(155A)	Registration required
5.5(155A)	Exempt from registration
5.6	Reserved
5.7(155A)	Registration application form
5.8	Reserved
5.9(155A)	Registration fee, term, and renewal
5.10 to 5.12	Reserved
5.13(155A)	Registration verification
5.14(155A)	Notifications to the board
5.15(155A)	Identification of pharmacy support person
5.16	Reserved
5.17(155A)	Tasks a pharmacy support person shall not perform
5.18 and 5.19	Reserved
5.20(155A)	Training and utilization of pharmacy support persons
5.21(155A)	Delegation of functions and responsibility of supervising pharmacist
5.22(155A)	Delegation of nontechnical functions
5.23	Reserved
5.24(155A)	Denial of registration
5.25(147,155A)	Unethical conduct or practice

- 5.26(155A) Reporting discipline and criminal convictions
- 5.27(155A) Discipline of pharmacy support persons

CHAPTER 6

GENERAL PHARMACY PRACTICE

- 6.1(155A) Purpose and scope
- 6.2(155A) Pharmacist in charge
- 6.3(155A) Reference library
- 6.4(155A) Exemption from duplicate requirements
- 6.5 and 6.6 Reserved
- 6.7(124,155A) Security
- 6.8(124,155A) Prescription processing documentation
- 6.9(124,155A) Transfer of prescription
- 6.10(126,155A) Prescription label requirements
- 6.11 and 6.12 Reserved
- 6.13(155A) Patient record system
- 6.14(155A) Patient counseling and instruction
- 6.15(124,126) Return of drugs and devices
- 6.16(124,155A) Records

CHAPTER 7

HOSPITAL PHARMACY PRACTICE

- 7.1(155A) Purpose and scope
- 7.2(155A) Pharmacist in charge
- 7.3(155A) Reference library
- 7.4 Reserved
- 7.5(124,155A) Security
- 7.6(155A) Pharmacist absence
- 7.7(155A) Verification by remote pharmacist
- 7.8(124,126,155A) Drug distribution and control
- 7.9(124,155A) Drug information
- 7.10(124,155A) Ensuring rational drug therapy
- 7.11(124,126,155A) Outpatient services
- 7.12(124,126,155A) Drugs in the emergency department
- 7.13(124,155A) Records

CHAPTER 8

UNIVERSAL PRACTICE STANDARDS

- 8.1(155A) Purpose and scope
- 8.2(155A) Definitions
- 8.3(155A) Responsible parties
- 8.4(155A) Pharmacist identification and staff logs
- 8.5(155A) Environment and equipment requirements
- 8.6 Reserved
- 8.7(155A) Procurement, storage, and recall of drugs and devices
- 8.8 Reserved
- 8.9(124,155A) Records storage
- 8.10 Reserved
- 8.11(147,155A) Unethical conduct or practice
- 8.12(126,147) Advertising
- 8.13(135C,155A) Personnel histories
- 8.14(155A) Training and utilization of registered pharmacy staff
- 8.15(155A) Delivery of prescription drugs and devices

8.16(124,155A)	Confidential information
8.17	Reserved
8.18(124,155A)	Electronic prescription mandate
8.19(124,126,135,155A,280)	Manner of issuance of a prescription drug or medication order
8.20(155A)	Valid prescriber/patient relationship
8.21(155A)	Prospective drug use review
8.22(155A)	Notification of interchangeable biological product selection
8.23	Reserved
8.24(155A)	Documented verification
8.25	Reserved
8.26(155A)	Continuous quality improvement program
8.27 to 8.34	Reserved
8.35(155A)	Pharmacy license

CHAPTER 9

Reserved

CHAPTER 10

CONTROLLED SUBSTANCES

10.1(124)	Purpose and scope
10.2(124)	Definitions
10.3(124)	Who shall register
10.4	Reserved
10.5(124)	Application
10.6(124)	Registration renewal
10.7(124)	Separate registration for independent activities; coincident activities
10.8(124)	Separate registrations for separate locations; exemption from registration
10.9(124)	Modification or termination of registration
10.10(124)	Denial of application or discipline of registration
10.11(124,147,155A)	Registration verification
10.12(124)	Inspection
10.13(124)	Security requirements
10.14(124)	Accountability of controlled substances
10.15	Reserved
10.16(124)	Receipt and disbursement of controlled substances
10.17(124)	Ordering or distributing Schedule I or II controlled substances
10.18(124)	Schedule II perpetual inventory
10.19(124)	Physical count and record of inventory
10.20(124)	Schedule III through V accountability
10.21(124)	Report of theft or loss
10.22(124)	Disposal of registrant stock
10.23(124)	Disposal of previously dispensed controlled substances
10.24(124,126,155A)	Prescription requirements
10.25(124)	Dispensing records
10.26(124)	Schedule II emergency prescriptions
10.27(124)	Schedule II prescriptions—partial filling
10.28(124)	Schedule II medication order
10.29(124)	Schedule II—issuing multiple prescriptions
10.30(124)	Schedule II—changes to a prescription
10.31	Reserved
10.32(124)	Schedule III, IV, or V prescription
10.33(124,155A)	Dispensing Schedule V controlled substances without a prescription

10.34(124)	Dispensing products containing ephedrine, pseudoephedrine, or phenylpropanolamine without a prescription
10.35	Reserved
10.36(124,155A)	Records
10.37	Reserved
10.38(124)	Revision of controlled substances schedules
10.39(124)	Temporary designation of controlled substances
10.40(124)	Excluded and exempt substances
10.41(124)	Anabolic steroid defined
10.42	Reserved
10.43(124)	Reporting discipline and criminal convictions
10.44(124)	Discipline

CHAPTER 11

DRUGS IN EMERGENCY MEDICAL SERVICE PROGRAMS

11.1(124,147A,155A)	Definitions
11.2(124,147A,155A)	Responsibility
11.3(124,147A,155A)	Registration required
11.4(124,147A,155A)	Written agreement
11.5(124,147A,155A)	Termination of agreement
11.6 and 11.7	Reserved
11.8(124,147A,155A)	Identification
11.9	Reserved
11.10(124,147A,155A)	Ownership of prescription drugs
11.11(124,147A,155A)	Policies and procedures
11.12	Reserved
11.13(124,147A,155A)	Storage
11.14(124,147A,155A)	Protocols
11.15(124,147A,155A)	Administration of drugs beyond the limits of a written protocol
11.16(124,147A,155A)	Administration of Schedule II controlled substances—pharmacy-based service program
11.17 and 11.18	Reserved
11.19(124,147A,155A)	Patient care reports
11.20(124,147A,155A)	Prescription drugs in service programs
11.21	Reserved
11.22(124,147A,155A)	Return of drugs
11.23(124,147A,155A)	Out-of-date drugs or devices
11.24(124,147A,155A)	Product recall
11.25	Reserved
11.26(124,147A,155A)	Controlled substances records
11.27(124,147A,155A)	Ordering Schedule II controlled substances—medical director-based service programs
11.28	Reserved
11.29(124,147A,155A)	Schedule II controlled substances perpetual inventory
11.30(124,147A,155A)	Controlled substances annual inventory
11.31	Reserved
11.32(124,147A,155A)	Disposition of controlled substances
11.33(124,147A,155A)	Report of loss or theft of controlled substance
11.34(124,147A,155A)	Records

CHAPTER 12
PRECURSOR SUBSTANCES

12.1(124B)	Precursor substance identified
12.2(124B)	Reports required
12.3(124B)	Form of reports
12.4(124B)	Monthly reporting option
12.5(124B)	Exemptions
12.6(124B)	Identification of purchaser or other recipient
12.7(124B)	Permits
12.8(124B)	Denial, modification, suspension, or revocation of permit

CHAPTER 13
TELEPHARMACY PRACTICE

13.1(155A)	Purpose and scope
13.2(155A)	Definitions
13.3(124,155A)	Written agreement
13.4 to 13.7	Reserved
13.8(124,155A)	General requirements for telepharmacy site
13.9(155A)	General requirements for managing pharmacy
13.10(155A)	General requirements for verifying pharmacist
13.11(155A)	General requirements for telepharmacy technician
13.12(155A)	General requirements for telepharmacy support person
13.13 to 13.15	Reserved
13.16(124,155A)	Telepharmacy site—initial application
13.17	Reserved
13.18(155A)	Opening of traditional pharmacy
13.19 to 13.22	Reserved
13.23(124,155A)	Records

CHAPTER 14
PUBLIC INFORMATION AND INSPECTION OF RECORDS

14.1(22,124,155A)	Definitions
14.2(22,124,155A)	Purpose and scope
14.3(22,124,155A)	Requests for access to records
14.4(22,124,155A)	Access to confidential records
14.5(22,124,155A)	Requests for treatment of a record as a confidential record and its withholding from examination
14.6(22,124,155A)	Procedure by which additions, dissents, or objections may be entered into certain records
14.7(22,124,155A)	Consent to disclosure by the subject of a confidential record
14.8(22,124,155A)	Notice to suppliers of information
14.9(22,124,155A)	Disclosures without the consent of the subject
14.10(22,124,155A)	Routine use
14.11(22,124,155A)	Consensual disclosure of confidential records
14.12(22,124,155A)	Release to subject
14.13(22,124,155A)	Availability of records
14.14(22,124,155A)	Personally identifiable information
14.15(22,124,155A)	Other groups of records
14.16(22,124,155A)	Computer

CHAPTER 15
CORRECTIONAL PHARMACY PRACTICE

- 15.1(155A) Purpose and scope
- 15.2(126,155A) Definitions
- 15.3(155A) Responsibilities
- 15.4(155A) Reference library
- 15.5(124,155A) Security
- 15.6 Reserved
- 15.7(124,126,155A) Training and utilization of pharmacy technicians or pharmacy support persons
- 15.8(124,126,155A) Drug distribution and dispensing controls
- 15.9 Reserved
- 15.10(124,126,155A) Policies and procedures

CHAPTER 16
NUCLEAR PHARMACY PRACTICE

- 16.1(155A) Purpose and scope
- 16.2(155A) Definitions
- 16.3(155A) Training requirements for authorized nuclear pharmacist
- 16.4(155A) General requirements for a pharmacy providing radiopharmaceutical services
- 16.5(155A) Library
- 16.6(155A) Minimum equipment requirements
- 16.7(155A) Training and utilization of pharmacy support persons
- 16.8(155A) Sterile radiopharmaceutical preparations and compounding

CHAPTER 17
WHOLESALE DISTRIBUTOR LICENSES

- 17.1(155A) Purpose and scope
- 17.2(155A) Definitions
- 17.3(155A) Wholesale distributor license
- 17.4(155A) Grounds for denial
- 17.5 and 17.6 Reserved
- 17.7(124,155A) Compliance with federal and state laws
- 17.8(124,155A) Written policies and procedures
- 17.9(155A) Facilities
- 17.10(124,155A) Security
- 17.11(155A) Storage and handling
- 17.12 to 17.16 Reserved
- 17.17(155A) Reporting discipline and criminal convictions
- 17.18(155A) Discipline

CHAPTER 18
CENTRALIZED PRESCRIPTION FILLING AND PROCESSING

- 18.1(155A) Purpose and scope
- 18.2(155A) Definitions
- 18.3(155A) General requirements
- 18.4 Reserved
- 18.5(155A) Patient notification and authorization
- 18.6 to 18.9 Reserved
- 18.10(155A) Policy and procedures
- 18.11 to 18.14 Reserved
- 18.15(155A) Records

CHAPTER 19
NONRESIDENT PHARMACY PRACTICE

19.1(155A)	Definitions
19.2(155A)	Nonresident pharmacy license
19.3(155A)	Registered pharmacist in charge
19.4(124,155A)	Applicability of board rules
19.5 and 19.6	Reserved
19.7(155A)	Confidential data
19.8(124,155A)	Storage and shipment of drugs and devices
19.9(155A)	Patient record system, prospective drug use review, and patient counseling
19.10(155A)	Reporting discipline and criminal convictions
19.11(155A)	Discipline

CHAPTER 20
COMPOUNDING PRACTICES

20.1(124,126,155A)	Purpose and scope
20.2(124,126,155A)	Definitions
20.3(124,126,155A)	Nonsterile compounding
20.4(124,126,155A)	Sterile compounding
20.5(126,155A)	Delayed compliance
20.6(126,155A)	Compounding standards for outsourcing facilities
20.7 and 20.8	Reserved
20.9(124,155A)	Prescriber/patient/pharmacist relationship
20.10(126,155A)	Anticipatory compounding
20.11(126,155A)	Prohibition on resale of compounded preparations
20.12(126,155A)	Compounding copies of an approved drug
20.13(124,126,155A)	Use of flavoring agents
20.14	Reserved
20.15(124,126,155A)	Compounding for office use
20.16(124,126,155A)	Compounding for hospital use
20.17 and 20.18	Reserved
20.19(124,126,155A)	Labeling
20.20(126,155A)	Labeling for batch preparation compounding
20.21 and 20.22	Reserved
20.23(124,126,155A)	Records
20.24(155A)	Annual reporting of interstate distribution of compounded preparations

CHAPTER 21
ELECTRONIC DATA AND AUTOMATED SYSTEMS IN PHARMACY PRACTICE

21.1(124,155A)	Purpose and scope
21.2(124,155A)	Definitions
21.3(124,155A)	System security and safeguards
21.4	Reserved
21.5(124,155A)	Automated data processing systems
21.6(124,155A)	Electronic prescription applications
21.7(124,155A)	Facsimile transmission of a prescription
21.8(124,155A)	Electronic prescription mandate and exemptions
21.9(124,155A)	Exemption from electronic prescription mandate—petition
21.10(124,155A)	Automated medication distribution system (AMDS)
21.11(124,155A)	Pharmacist verification of controlled substance fills—daily printout or logbook

CHAPTER 22

UNIT DOSE, ALTERNATIVE PACKAGING, AND EMERGENCY BOXES

22.1(155A)	Unit dose dispensing systems
22.2	Reserved
22.3(126)	Prepackaging
22.4	Reserved
22.5(126,155A)	Patient med paks
22.6	Reserved
22.7(124,155A)	Emergency/first dose drug supply
22.8	Reserved
22.9(155A)	Home health agency/hospice emergency drugs

CHAPTER 23

CARE FACILITY PHARMACY PRACTICE

23.1(155A)	Purpose and scope
23.2(155A)	Definitions
23.3(124,155A)	Freedom of choice
23.4(124,155A)	Responsibilities
23.5(124,155A)	Emergency drugs
23.6	Reserved
23.7(124,155A)	Policies and procedures
23.8	Reserved
23.9(124,155A)	Medication orders
23.10	Reserved
23.11(124,155A)	Drugs dispensed—general requirements
23.12	Reserved
23.13(124,155A)	Labeling drugs under special circumstances
23.14(124,155A)	Provision of drugs to a facility for immunization or screening programs
23.15(124,155A)	Return and reuse of drugs and devices
23.16	Reserved
23.17(124,155A)	Accountability of controlled substances
23.18	Reserved
23.19(124,155A)	Dispensing Schedule II controlled substances
23.20(124,155A)	Partial filling of Schedule II controlled substances
23.21(124,155A)	Disposal of previously dispensed controlled substances

CHAPTER 24

Reserved

CHAPTER 25

CHILD SUPPORT NONCOMPLIANCE

25.1(252J)	Definitions
25.2(252J)	Issuance or renewal of license—denial
25.3(252J)	Suspension or revocation of a license
25.4(17A,22,252J)	Share information

CHAPTER 26

PETITIONS FOR RULE MAKING

(Uniform Rules)

26.1(17A)	Petition for rule making
26.2(17A)	Briefs
26.3(17A)	Inquiries
26.4(17A)	Board consideration

CHAPTER 27
DECLARATORY ORDERS
(Uniform Rules)

27.1(17A)	Petition for declaratory order
27.2(17A)	Notice of petition
27.3(17A)	Intervention
27.4(17A)	Briefs
27.5(17A)	Inquiries
27.6(17A)	Service and filing of petitions and other papers
27.7(17A)	Consideration
27.8(17A)	Action on petition
27.9(17A)	Refusal to issue order
27.10(17A)	Contents of declaratory order—effective date
27.11(17A)	Copies of orders
27.12(17A)	Effect of a declaratory order

CHAPTER 28
AGENCY PROCEDURE FOR RULE MAKING

28.1(17A)	Applicability
28.2(17A)	Definitions
28.3(17A)	Solicitation of comments before notice
28.4(17A)	Public rule-making docket
28.5(17A)	Public hearing proceedings
28.6(17A)	Regulatory analyses
28.7(17A,25B)	Fiscal impact statement
28.8(17A)	Time and manner of rule adoption
28.9(17A)	Variance between adopted rule and published notice of proposed rule adoption
28.10(17A)	Exemptions from public rule-making procedures
28.11(17A)	Concise statement of reasons
28.12(17A)	Style and form
28.13(17A)	Board rule-making record
28.14(17A)	Filing of rules
28.15(17A)	Effectiveness of rules prior to publication
28.16(17A)	Review by board of rules

CHAPTER 29
SALES OF GOODS AND SERVICES

29.1(68B)	Selling of goods or services by members of the board
29.2(68B)	Conditions of consent for board members
29.3(68B)	Authorized sales
29.4(68B)	Application for consent
29.5(68B)	Limitation of consent

CHAPTER 30
IOWA MONITORING PROGRAM FOR PHARMACY PROFESSIONALS

30.1(272C)	Iowa monitoring program for pharmacy professionals committee
30.2(272C)	Definitions
30.3(272C)	Organization of the committee
30.4(272C)	Eligibility
30.5(272C)	Terms of participation
30.6(272C)	Confidentiality
30.7(28E)	Authority for 28E agreements

CHAPTER 31
CRIMINAL CONVICTIONS

- 31.1(272C) Purpose and scope
- 31.2(272C) Definitions
- 31.3(272C) License application
- 31.4(272C) Eligibility determination
- 31.5(272C) Appeal
- 31.6(272C) Future petitions or applications

CHAPTER 32
NONPAYMENT OF STATE DEBT

- 32.1(272D) Definitions
- 32.2(272D) Issuance or renewal of a license—denial
- 32.3(272D) Suspension or revocation of a license
- 32.4(17A,22,272D) Share information

CHAPTER 33
MILITARY SERVICE AND VETERAN RECIPROCITY

- 33.1(272C) Definitions
- 33.2(272C) Military education, training, and service credit
- 33.3(272C) Veteran licensure or registration
- 33.4(272C) Request for contested case

CHAPTER 34
WAIVERS

- 34.1(17A) Definition
- 34.2(17A,124,126,147,155A,205,272C) Scope of chapter
- 34.3(17A,124,126,147,155A,205,272C) Applicability of chapter
- 34.4(17A) Criteria for waiver
- 34.5(17A,124,126,147,155A,205,272C) Filing of petition
- 34.6(17A) Content of petition
- 34.7(17A) Additional information and providing notice
- 34.8 and 34.9 Reserved
- 34.10(17A) Ruling
- 34.11(17A,22) Public availability
- 34.12(17A) Submission of waiver information
- 34.13(17A) Cancellation of a waiver
- 34.14(17A,124,126,147,155A,205,272C) Violations
- 34.15(17A,124,126,147,155A,205,272C) Defense
- 34.16(17A) Judicial review

CHAPTER 35
CONTESTED CASES

- 35.1(17A,124,124B,126,147,155A,205,272C) Scope and applicability
- 35.2(17A,272C) Definitions
- 35.3(17A) Time requirements
- 35.4(17A) Applicability of Iowa Rules of Civil Procedure
- 35.5(17A,272C) Combined statement of charges and settlement agreement
- 35.6(17A,124B,126,147,155A,205,272C) Notice of hearing
- 35.7(17A,272C) Statement of charges
- 35.8(13,272C) Legal representation
- 35.9(17A,272C) Presiding officer in a disciplinary contested case
- 35.10(17A,272C) Presiding officer for nondisciplinary hearings

- 35.11(17A,124B,147,155A,272C) Waiver of procedures
- 35.12(17A,272C) Telephone or electronic proceedings
- 35.13(17A) Disqualification
- 35.14(17A,272C) Consolidation—severance
- 35.15(17A,272C) Appearance
- 35.16(17A,272C) Answer
- 35.17(17A,272C) Service and filing of documents
- 35.18(272C) Investigative file
- 35.19(17A,272C) Discovery
- 35.20(17A,272C) Issuance of subpoenas in a contested case
- 35.21(17A,272C) Motions
- 35.22(17A,272C) Prehearing conference
- 35.23(17A,272C) Continuances
- 35.24(17A,272C) Settlement agreements
- 35.25(17A,124B,126,147,155A,205,272C) Hearing procedures in contested cases
- 35.26(17A,272C) Evidence
- 35.27(17A,272C) Default
- 35.28(17A,272C) Ex parte communication
- 35.29(17A,272C) Recording costs
- 35.30(17A,272C) Proposed decisions
- 35.31(17A) Final decision
- 35.32(17A,124B,126,147,155A,205,272C) Applications for rehearing
- 35.33(17A,272C) Stays of board actions
- 35.34(17A,272C) No factual dispute contested cases
- 35.35(17A,124B,126,147,155A,205,272C) Emergency adjudicative proceedings
- 35.36(17A,147,272C) Application for reinstatement
- 35.37(17A,22,272C) Dissemination of public records
- 35.38(17A) Judicial review

CHAPTER 36 DISCIPLINE

- 36.1(147,155A,272C) Authority
- 36.2(147,155A,272C) Definitions
- 36.3(147,155A,272C) Complaints, investigations, and board action
- 36.4(17A,147,152,272C) Issuance of investigatory subpoenas
- 36.5(147,272C) Peer review committee
- 36.6(147,155A,272C) Grounds for discipline
- 36.7(147,155A,272C) Disciplinary sanctions
- 36.8(147,272C) Voluntary surrender
- 36.9(155A,272C) Order for mental or physical examination
- 36.10(272C) Disciplinary hearings—fees and costs
- 36.11(88GA,SF304) Prohibited grounds for discipline

CHAPTER 37 IOWA PRESCRIPTION MONITORING PROGRAM

- 37.1(124) Purpose and scope
- 37.2(124) Definitions
- 37.3(124) Registration
- 37.4(124) Prescription monitoring program advisory council
- 37.5 Reserved
- 37.6(124) Security of PMP credentials
- 37.7(124) PMP reporting—exemptions

37.8(124)	PMP reporting—dispensing prescribers
37.9(124)	PMP reporting—pharmacies
37.10 and 37.11	Reserved
37.12(124)	Reporting requirements
37.13(124)	Opioid antagonist administration by first responders
37.14 and 37.15	Reserved
37.16(124)	Access to PMP information
37.17(124)	Integrated systems
37.18(124)	PMP administrator access
37.19(124)	Prescriber activity reports
37.20(124)	Proactive notifications
37.21(124)	Record retention
37.22(124)	Information errors
37.23(124)	Discipline

CHAPTER 38

Reserved

CHAPTER 39

EXPANDED PRACTICE STANDARDS

39.1(155A)	Purpose and scope
39.2 and 39.3	Reserved
39.4(155A)	Pharmaceutical care
39.5	Reserved
39.6(155A)	Statewide protocols
39.7(135,147A)	Opioid antagonist dispensing by pharmacist—standing order
39.8(155A)	Medications administered via prescription
39.9 to 39.12	Reserved
39.13(155A)	Collaborative pharmacy practice
39.14 and 39.15	Reserved
39.16(155A)	Pharmacy pilot or demonstration research projects

CHAPTER 40

TECHNOLOGY-ASSISTED TECHNICIAN PRODUCT VERIFICATION PROGRAMS

40.1(155A)	Purpose and scope
40.2(155A)	Definitions
40.3(155A)	TPV program requirements
40.4(155A)	Checking technician requirements
40.5 and 40.6	Reserved
40.7(155A)	Policies and procedures
40.8(155A)	TPV program quality assurance
40.9 and 40.10	Reserved
40.11(155A)	TPV program records

CHAPTER 41

OUTSOURCING FACILITIES

41.1(155A)	Purpose and scope
41.2(155A)	Definitions
41.3(155A)	Outsourcing facility license
41.4(155A)	Applicability of board rules
41.5(155A)	Reporting discipline and criminal convictions
41.6(155A)	Discipline

CHAPTER 42
LIMITED DISTRIBUTOR LICENSES

42.1(155A)	Purpose and scope
42.2(155A)	Definitions
42.3(155A)	Limited distributor license
42.4 and 42.5	Reserved
42.6(155A)	Grounds for denial
42.7(155A)	Policies and procedures
42.8 and 42.9	Reserved
42.10(155A)	Requirements
42.11	Reserved
42.12(155A)	Records
42.13	Reserved
42.14(155A)	Reporting discipline and criminal convictions
42.15(155A)	Discipline

CHAPTER 43
THIRD-PARTY LOGISTICS PROVIDER LICENSES

43.1(155A)	Purpose and scope
43.2(155A)	Definitions
43.3(155A)	3PL license
43.4	Reserved
43.5(155A)	Compliance with federal and state laws
43.6(155A)	Policies and procedures
43.7 and 43.8	Reserved
43.9(155A)	Reporting discipline and criminal conviction
43.10(155A)	Discipline

CHAPTERS 44 to 99
Reserved

CHAPTER 100
IOWA REAL-TIME ELECTRONIC PSEUDOEPHEDRINE
TRACKING SYSTEM

100.1(124)	Purpose and scope
100.2(124)	Definitions
100.3(124)	Electronic pseudoephedrine tracking system (PTS)
100.4(124)	Access to database information and confidentiality
100.5(124)	Violations

CHAPTER 3
PHARMACY TECHNICIANS

[Prior to 9/4/02, see 657—Ch 22]

657—3.1(155A) Definitions. For the purposes of this chapter, the following definitions shall apply:

“*Board*” means the Iowa board of pharmacy.

“*Cashier*” means a person whose duties within the pharmacy are limited to accessing finished, packaged prescription orders and processing payments for and delivering such orders to the patient or the patient’s representative.

“*Certified pharmacy technician*” or “*certified technician*” means an individual who holds a valid current national certification and who has registered with the board as a certified pharmacy technician.

“*Delivery*” means the transport and conveyance of a finished, securely packaged prescription order to the patient or the patient’s caregiver.

“*Nationally accredited program*” means a program and examination for the certification of pharmacy technicians that is accredited by the NCCA.

“*NCCA*” means the National Commission for Certifying Agencies.

“*Pharmacy support person*” means a person, other than a licensed pharmacist, a registered pharmacist-intern, or a registered pharmacy technician, who may perform nontechnical duties assigned by the pharmacist under the pharmacist’s responsibility and supervision pursuant to 657—Chapter 5.

“*Pharmacy technician*” or “*technician*” means a person who is employed in Iowa by a licensed pharmacy under the responsibility of an Iowa-licensed pharmacist to assist in the technical functions of the practice of pharmacy, as provided in rule 657—3.21(155A), and includes a certified pharmacy technician and a pharmacy technician trainee.

“*Pharmacy technician certification*” or “*national certification*” means a certificate issued by a national pharmacy technician certification authority accredited by the NCCA attesting that the technician has successfully completed the requirements of the certification program. The term includes evidence of renewal of the national certification.

“*Pharmacy technician trainee*” or “*technician trainee*” means an individual who is in training to become a pharmacy technician and who is in the process of acquiring national certification as a pharmacy technician as provided in rule 657—3.5(155A).

“*Pharmacy technician training*” or “*technician training*” means education or experience acquired for the purpose of qualifying for and preparing for national certification.

“*Supervising pharmacist*” means an Iowa-licensed pharmacist who is on duty in a licensed pharmacy in Iowa and who is responsible for the actions of a pharmacy technician or other supportive personnel. [ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.2(155A) Purpose of registration. A registration program for pharmacy technicians is established for the purposes of determining the competency of a pharmacy technician or of an applicant for registration as a certified pharmacy technician or pharmacy technician trainee and for the purposes of identification, tracking, and disciplinary action for violations of federal or state pharmacy or drug laws or regulations.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15]

657—3.3(155A) Registration required. Any person employed in Iowa as a pharmacy technician shall obtain and maintain during such employment a current registration as a certified pharmacy technician or pharmacy technician trainee pursuant to these rules. An individual commencing employment as a pharmacy technician in Iowa who fails to register as a certified pharmacy technician or pharmacy technician trainee as provided by these rules may be subject to disciplinary sanctions. A certified pharmacy technician commencing employment as a certified pharmacy technician in Iowa who fails to register as a certified pharmacy technician or who fails to maintain national certification may be subject to disciplinary sanctions. A pharmacist-intern with a current registration or a pharmacist with a current license is not required to obtain a pharmacy technician registration to work as a pharmacy technician.

3.3(1) Licensed health care provider. Except as provided in this rule, a licensed health care provider whose registration or license is in good standing with and not subject to current disciplinary sanctions or practice restrictions imposed by the licensee's professional licensing board and who assists in the technical functions of the practice of pharmacy shall be required to register as a certified pharmacy technician or pharmacy technician trainee pursuant to these rules. A registered nurse licensed pursuant to Iowa Code chapter 152 or 152E who is engaged in the administration of immunizations and vaccinations and the utilization of statewide protocols pursuant to a pharmacist's order is exempt from registration.

3.3(2) Registration required. Beginning July 1, 2021, any person not currently registered with the board as a pharmacy technician shall obtain registration prior to commencement of employment in an Iowa pharmacy as a pharmacy technician. Through June 30, 2021, any person not currently registered with the board as a pharmacy technician shall submit a completed application for registration within 30 days of accepting employment in an Iowa pharmacy as a pharmacy technician.

3.3(3) Technician training. A person who is enrolled in a college-based or American Society of Health-System Pharmacists (ASHP)-accredited technician training program shall obtain a pharmacy technician trainee registration prior to beginning on-site practical experience. A person who is employed in a pharmacy and who is receiving pharmacy technician training through work experience shall obtain a pharmacy technician trainee registration prior to the commencement of pharmacy technician training.

3.3(4) Registration number. Each pharmacy technician registered with the board will be assigned a unique registration number.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 9407B, IAB 3/9/11, effective 4/13/11; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5542C, IAB 4/7/21, effective 5/12/21; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—3.4 Reserved.

657—3.5(155A) Certification of pharmacy technicians. Except as provided in subrule 3.5(1), all pharmacy technicians shall be required to be nationally certified as provided by this rule. National certification acquired through successful completion of any NCCA-accredited pharmacy technician certification program and examination fulfills the requirement for national certification. National certification does not replace the need for licensed pharmacist control over the performance of delegated functions, nor does national certification exempt the pharmacy technician from registration pursuant to these rules. A certified pharmacy technician shall maintain the technician's national certification, in addition to the technician's Iowa registration, during any period of employment in or for an Iowa pharmacy as a certified pharmacy technician.

3.5(1) Pharmacy technician trainee. A person who is in the process of acquiring national certification as a pharmacy technician shall register with the board as a pharmacy technician trainee pursuant to rule 657—3.9(155A).

3.5(2) Certified pharmacy technician. All applicants for a new pharmacy technician registration except as provided by subrule 3.5(1), and all applicants for renewal of a pharmacy technician registration pursuant to rule 657—3.10(155A), shall provide proof of current national pharmacy technician certification and shall complete the application for certified pharmacy technician registration.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 9407B, IAB 3/9/11, effective 4/13/11; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 6076C, IAB 12/15/21, effective 1/19/22; ARC 6416C, IAB 7/27/22, effective 8/31/22]

657—3.6(155A) Extension of deadline for national certification. Rescinded ARC 1785C, IAB 12/10/14, effective 1/14/15.

657—3.7 Reserved.

657—3.8(155A) Application.

3.8(1) An applicant shall submit a completed application along with the appropriate nonrefundable application fee pursuant to rule 657—3.9(155A) or 657—3.10(155A).

3.8(2) The application shall include:

- a. Information sufficient to identify the applicant including, but not limited to, name, address, date of birth, gender, and social security number;
 - b. Current place or places of employment;
 - c. Criminal or disciplinary action history;
 - d. If the application is for certified pharmacy technician registration, documentation of current national pharmacy technician certification; and
 - e. Any other information deemed necessary by the board.
- [ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.9(155A) Registration fee and term—technician trainee.

3.9(1) Fee. The nonrefundable application fee for a pharmacy technician trainee registration shall be \$20. The nonrefundable application fee for a pharmacy technician trainee registration shall be submitted in the form of a personal check, certified check, cashier's check, or money order made payable to the Iowa Board of Pharmacy when submitted with a written application or by acceptable debit or credit card when submitted with an online application.

3.9(2) Term. A pharmacy technician trainee registration shall expire 12 months following the date of registration. A pharmacy technician trainee registration may be renewed only as provided in subrules 3.9(3) and 3.9(4).

a. *National certification completed.* When the registered pharmacy technician trainee completes national certification, and no later than the expiration of the pharmacy technician trainee registration, the technician shall submit a completed application and nonrefundable application fee for certified pharmacy technician registration.

b. *Expiration of registration.* Except as provided in subrules 3.9(3) and 3.9(4), the registration of a pharmacy technician trainee who fails to complete national certification prior to the expiration of the registration shall expire and the technician shall cease practice as a pharmacy technician.

3.9(3) Renewal. A technician trainee who is unable to complete national certification prior to the expiration of the registration may seek renewal of the registration in exceptional circumstances. To the extent practicable, the trainee should submit an application and nonrefundable fee of \$20 for technician trainee renewal, on forms provided by the board, at least 30 days prior to the expiration of the registration.

3.9(4) Reactivation. A technician trainee who was previously registered and left the practice of pharmacy prior to obtaining national certification may seek reactivation of the registration. The individual shall submit an application and nonrefundable fee of \$20 for technician trainee reactivation on forms provided by the board. Pursuant to rule 657—3.3(155A), a technician shall obtain registration prior to commencing employment as a technician trainee in an Iowa pharmacy.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5542C, IAB 4/7/21, effective 5/12/21; ARC 6076C, IAB 12/15/21, effective 1/19/22]

657—3.10(155A) Registration fee, term, and renewal—certified pharmacy technician.

3.10(1) Fee. The nonrefundable application fee for a certified pharmacy technician registration shall be \$40 per biennium. The nonrefundable application fee for a certified pharmacy technician registration shall be submitted in the form of a personal check, certified check, cashier's check, or money order made payable to the Iowa Board of Pharmacy when submitted with a written application or by acceptable debit or credit card when submitted with an online application.

3.10(2) Term. A certified pharmacy technician registration shall expire on the date that the technician's national certification expires.

3.10(3) Renewal. A certified pharmacy technician registration shall be renewed prior to the expiration of the registration.

a. *Delinquent registration grace period.* A certified pharmacy technician registration which is not renewed prior to the expiration of the registration shall be considered delinquent. Renewal during the month following the expiration date of the registration shall include the nonrefundable registration fee pursuant to subrule 3.10(1) and a nonrefundable late penalty fee of \$40. A registered certified pharmacy technician who renews during the month following the expiration date of the registration shall not be

subject to disciplinary action for continuing to practice as a pharmacy technician during the delinquency of the registration.

b. Registration reactivation beyond grace period. If the registration is not renewed prior to the expiration of the one-month grace period identified in paragraph 3.10(3)“a,” the technician shall cease the practice as a pharmacy technician until the registration is reactivated. A certified pharmacy technician without a current registration may apply for registration reactivation by submitting a completed application for reactivation and a nonrefundable reactivation fee of \$160. An individual who continues employment as a pharmacy technician without a current registration, in addition to the pharmacy and the pharmacist in charge that allow the individual to continue practice as a pharmacy technician, may be subject to disciplinary sanctions.

c. Voluntary cancellation. A registered certified pharmacy technician who ceases practice as a pharmacy technician and does not intend to renew the registration prior to its expiration may request that the board cancel the registration. If the certified pharmacy technician later seeks registration as a certified pharmacy technician, the technician shall not be assessed a late penalty fee or reactivation fee for renewal of the registration.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 0504C, IAB 12/12/12, effective 1/16/13; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.11(155A) Verification fee. The board may require the submission of a nonrefundable fee of \$15 for written verification of a registration.

[ARC 0504C, IAB 12/12/12, effective 1/16/13; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.12(155A) Registration certificates. Rescinded ARC 5542C, IAB 4/7/21, effective 5/12/21.

657—3.13(155A) Notifications to the board. A pharmacy technician shall report to the board within ten days a change of the technician’s name, address, or pharmacy employment status.

[ARC 9009B, IAB 8/11/10, effective 7/23/10]

657—3.14 to 3.16 Reserved.

657—3.17(155A) Training and utilization of pharmacy technicians.

3.17(1) Policies and procedures. All licensed pharmacies located in Iowa that utilize pharmacy technicians shall develop, implement, and periodically review written policies and procedures for the training and utilization of pharmacy technicians appropriate to the practice of pharmacy. Pharmacy policies shall specify the frequency of review.

3.17(2) Documented training. Pharmacy technician training shall be documented and maintained by the pharmacy for the duration of employment.

3.17(3) Vaccine administration training and continuing education. A pharmacy technician who administers a vaccine or immunization under the supervision of a pharmacist shall document successful completion of the requirements in paragraph 3.17(3)“a” or “b” and shall maintain competency by completing and maintaining documentation of the continuing education requirements in paragraph 3.17(3)“c.”

a. Initial qualification. Except as provided in paragraph 3.17(3)“b,” a technician shall have successfully completed an Accreditation Council for Pharmacy Education (ACPE)-accredited program on vaccine administration that is an evidence-based program that includes study material and hands-on training and techniques for administering vaccines, requires testing with a passing score, complies with current Centers for Disease Control and Prevention (CDC) guidelines, and provides instruction and experiential training in the following content areas:

- (1) Standards for immunization practices;
- (2) Basic immunology and vaccine protection;
- (3) Vaccine-preventable diseases;
- (4) Recommended immunization schedules;
- (5) Vaccine storage and management;
- (6) Informed consent;

- (7) Physiology and techniques for vaccine administration;
- (8) Immunization record management; and
- (9) Identification of adverse events.

b. Previous qualification. A technician who is currently licensed as a registered nurse shall be deemed to have met the training requirement.

c. Continuing education. During any technician registration renewal period, a technician who engages in the administration of vaccines shall complete and document at least one hour of ACPE-approved continuing education with the ACPE topic designator “06” followed by the letter “T” or “P.”

d. Certification maintained. During any period within which a technician may engage in the administration of vaccines, the technician shall maintain current certification in basic cardiac life support through a training program designated for health care providers that includes hands-on training. [ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 5820C, IAB 8/11/21, effective 7/15/21; ARC 6071C, IAB 12/15/21, effective 1/19/22]

657—3.18(147,155A) Identification of pharmacy technician.

3.18(1) Identification badge. A pharmacy technician shall wear a visible identification badge while on duty that clearly identifies the person as a pharmacy technician and that includes at least the technician’s first name.

3.18(2) Misrepresentation prohibited. A pharmacy technician shall not represent himself or herself in any manner as a pharmacist or pharmacist-intern. A pharmacy technician shall not represent himself or herself in any manner as a certified pharmacy technician unless the technician has attained national pharmacy technician certification.

[ARC 9009B, IAB 8/11/10, effective 7/23/10]

657—3.19 Reserved.

657—3.20(155A) Responsibility of supervising pharmacist. The ultimate responsibility for the actions of a pharmacy technician shall remain with the supervising pharmacist. A pharmacy license holder shall not infringe on the authority of a supervising pharmacist to delegate or decline to delegate specific functions to a pharmacy technician based on the supervising pharmacist’s professional judgment regarding the knowledge and training of the technician.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.21(155A) Delegation of functions.

3.21(1) Policies and procedures. Pursuant to established policies and procedures and the supervising pharmacist’s professional judgment, an Iowa-licensed supervising pharmacist may delegate any technical or nontechnical functions in the operation of the pharmacy, except those which are prohibited pursuant to rule 657—3.23(155A), to an appropriately trained and Iowa-registered pharmacy technician.

3.21(2) Remote supervision. A supervising pharmacist may delegate technical functions relating to prescription processing (e.g., data entry) to a certified pharmacy technician who is performing the delegated functions at a location that is not a licensed pharmacy only if the following conditions are met:

a. Adequate security and supervision are maintained at all times to prevent unauthorized access to, and unauthorized storage/transfer of, confidential patient information or patient records;

b. The supervising pharmacist has real-time access to the prescription processing system which the certified pharmacy technician is using or the patient record which the certified pharmacy technician is processing;

c. The supervising pharmacist is available to respond to certified pharmacy technician questions via a real-time communication mechanism at all times when delegated functions are being performed; and

d. The pharmacy’s prescription processing system is capable of documenting the functions performed by the certified pharmacy technician.

3.21(3) Pharmacist final verification required. Except as provided for an approved technician product verification program pursuant to 657—Chapter 40, the pharmacist shall provide and document the final verification for the accuracy, validity, completeness, and appropriateness of the patient's prescription or medication order prior to the delivery of the medication to the patient or the patient's representative.

3.21(4) Further delegation prohibited. A pharmacy technician shall not delegate technical functions to a pharmacy support person.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9783B, IAB 10/5/11, effective 11/9/11; ARC 4189C, IAB 12/19/18, effective 1/23/19; ARC 5007C, IAB 3/25/20, effective 4/29/20; ARC 5542C, IAB 4/7/21, effective 5/12/21; ARC 6416C, IAB 7/27/22, effective 8/31/22]

657—3.22(155A) Technical functions. Rescinded ARC 5542C, IAB 4/7/21, effective 5/12/21.

657—3.23(155A) Functions a pharmacy technician shall not perform.

3.23(1) Prohibited functions for all pharmacy technicians. A pharmacy technician shall not be authorized to perform any of the following functions:

a. Except for a certified pharmacy technician participating in an approved technician product verification program pursuant to 657—Chapter 40, provide the final verification for the accuracy, validity, completeness, or appropriateness of a filled prescription or medication order;

b. Conduct prospective drug use review or evaluate a patient's medication record for purposes identified in rule 657—8.21(155A);

c. Provide patient counseling, consultation, or patient-specific drug information, tender an offer of patient counseling on behalf of a pharmacist, or accept a refusal of patient counseling from a patient or patient's agent;

d. Make decisions that require a pharmacist's professional judgment, such as interpreting prescription drug orders or applying information;

e. Transfer a prescription drug order for a controlled substance to another pharmacy or receive the transfer of a prescription drug order for a controlled substance from another pharmacy;

f. Delegate technical functions to a pharmacy support person.

3.23(2) Prohibited functions for technician trainees. In addition to the prohibited functions in subrule 3.23(1), a technician trainee shall not be authorized to perform any of the following functions:

a. Accept new prescription drug orders or medication orders communicated to the pharmacy by a prescriber or the prescriber's agent.

b. Transfer or receive by transfer by any means the original prescription drug order information or prescription refill information of a prescription for any substance.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 9783B, IAB 10/5/11, effective 11/9/11; ARC 4189C, IAB 12/19/18, effective 1/23/19; ARC 5007C, IAB 3/25/20, effective 4/29/20; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.24(155A) New prescription drug orders or medication orders. Rescinded ARC 5542C, IAB 4/7/21, effective 5/12/21.

657—3.25(155A) Delegation of nontechnical functions. Rescinded IAB 4/7/10, effective 6/1/10.

657—3.26 and 3.27 Reserved.

657—3.28(147,155A) Unethical conduct or practice. Violation by a pharmacy technician of any of the provisions of this rule shall constitute unethical conduct or practice and may be grounds for disciplinary action as provided in rule 657—3.31(155A).

3.28(1) Misrepresentative deeds. A pharmacy technician shall not make any statement tending to deceive, misrepresent, or mislead anyone, or be a party to or an accessory to any fraudulent or deceitful practice or transaction in pharmacy or in the operation or conduct of a pharmacy.

3.28(2) Confidentiality. In the absence of express written authorization from the patient or written order or direction of a court, except where the best interests of the patient require, a pharmacy

technician shall not divulge or reveal to any person other than the patient or the patient's authorized representative, the prescriber or other licensed practitioner then caring for the patient, a licensed pharmacist, a person duly authorized by law to receive such information, or as otherwise provided in rule 657—8.16(124,155A), any of the following:

a. A patient's name, address, social security number, or any information that could be used to identify a patient;

b. The contents of any prescription drug order or medication order or the therapeutic effect thereof, or the nature of professional pharmaceutical services rendered to a patient;

c. The nature, extent, or degree of illness suffered by any patient; or

d. Any medical information furnished by the prescriber or the patient.

3.28(3) *Discrimination.* It is unethical to unlawfully discriminate between patients or groups of patients for reasons of religion, race, creed, color, gender, gender identity, sexual orientation, marital status, age, national origin, physical or mental disability, or disease state when providing pharmaceutical services.

3.28(4) *Unethical conduct or behavior.* A pharmacy technician shall not exhibit unethical behavior in connection with the technician's pharmacy employment. Unethical behavior shall include, but is not limited to, the following acts: verbal or physical abuse, coercion, intimidation, harassment, sexual advances, threats, degradation of character, indecent or obscene conduct, and theft.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.29(155A) Denial of registration. The executive director or designee may deny an application for registration as a certified pharmacy technician or pharmacy technician trainee for any violation of the laws of this state, another state, or the United States relating to prescription drugs, controlled substances, or nonprescription drugs or for any violation of Iowa Code chapter 124, 124B, 126, 147, 155A, or 205 or any rule of the board.

An individual whose application for registration as a certified pharmacy technician or pharmacy technician trainee is denied pursuant to this rule may, within 30 days after issuance of the notice of denial, appeal to the board for reconsideration of the application.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 3857C, IAB 6/20/18, effective 7/25/18]

657—3.30(155A) Reporting discipline and criminal convictions. A registered pharmacy technician shall provide to the board written notice of and unredacted documents related to any disciplinary or enforcement action imposed by any licensing agency or regulatory authority on any license or registration held by the registered pharmacy technician no later than 30 days after the final action. Discipline may include, but is not limited to, fine or civil penalty, citation or reprimand, probationary period, suspension, revocation, and voluntary surrender. A registered pharmacy technician shall provide written notice to the board of any criminal conviction that is related to the practice of pharmacy or controlled substances no later than 30 days after the conviction. The term "criminal conviction" includes instances when the judgment of conviction or sentence is deferred.

[ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—3.31(155A) Discipline of pharmacy technicians.

3.31(1) *Violations.* The board may impose discipline for any violation of the laws of this state, another state, or the United States relating to prescription drugs, controlled substances, or nonprescription drugs, or for any violation of Iowa Code chapter 124, 124B, 126, 147, 155A, or 205 or any rule of the board.

3.31(2) *Sanctions.* The board may impose the following disciplinary sanctions:

a. Revocation of a certified pharmacy technician or pharmacy technician trainee registration.

b. Suspension of a certified pharmacy technician or pharmacy technician trainee registration until further order of the board or for a specified period.

c. Nonrenewal of a certified pharmacy technician registration.

- d. Prohibition, permanently, until further order of the board, or for a specified period, from engaging in specified procedures, methods, or acts.
- e. Probation.
- f. The ordering of a physical or mental examination.
- g. The imposition of civil penalties not to exceed \$25,000.
- h. Issuance of a citation and warning.
- i. Such other sanctions allowed by law as may be appropriate.

[ARC 9009B, IAB 8/11/10, effective 7/23/10; ARC 1785C, IAB 12/10/14, effective 1/14/15; ARC 3857C, IAB 6/20/18, effective 7/25/18; ARC 5542C, IAB 4/7/21, effective 5/12/21]

These rules are intended to implement Iowa Code sections 147.72, 147.80, 147.107, 155A.6A, 155A.23, 155A.33, 155A.34, and 155A.39.

- [Filed 2/27/97, Notice 1/1/97—published 3/26/97, effective 4/30/97]
- [Filed 4/24/98, Notice 3/11/98—published 5/20/98, effective 6/24/98]
- [Filed 2/22/99, Notice 10/21/98—published 3/10/99, effective 4/14/99]
- [Filed 9/8/99, Notice 6/2/99—published 10/6/99, effective 11/10/99]
- [Filed 8/14/02, Notice 6/12/02—published 9/4/02, effective 10/9/02]
- [Filed 3/11/04, Notice 8/6/03—published 3/31/04, effective 5/5/04]
- [Filed emergency 7/16/04 after Notice 6/9/04—published 8/4/04, effective 7/16/04]
- [Filed 10/22/04, Notice 3/31/04—published 11/10/04, effective 12/15/04]
- [Filed emergency 6/30/05 after Notice 5/11/05—published 7/20/05, effective 7/1/05]
- [Filed 3/22/06, Notice 1/18/06—published 4/12/06, effective 5/17/06]
- [Filed 5/17/06, Notice 4/12/06—published 6/7/06, effective 7/12/06]
- [Filed 2/7/07, Notice 10/25/06—published 2/28/07, effective 4/4/07]
- [Filed emergency 11/13/07 after Notice 8/29/07—published 12/5/07, effective 11/13/07]
- [Filed 3/5/08, Notice 12/19/07—published 3/26/08, effective 4/30/08]¹
- [Filed emergency 6/9/08—published 7/2/08, effective 7/9/08]
- [Filed ARC 8673B (Notice ARC 8380B, IAB 12/16/09), IAB 4/7/10, effective 6/1/10]
- [Filed Emergency ARC 9009B, IAB 8/11/10, effective 7/23/10]
- [Editorial change: IAC Supplement 10/6/10]
- [Filed ARC 9407B (Notice ARC 9193B, IAB 11/3/10), IAB 3/9/11, effective 4/13/11]
- [Filed ARC 9502B (Notice ARC 9297B, IAB 12/29/10), IAB 5/18/11, effective 6/22/11]
- [Filed ARC 9783B (Notice ARC 9557B, IAB 6/15/11), IAB 10/5/11, effective 11/9/11]
- [Filed ARC 0504C (Notice ARC 0351C, IAB 10/3/12), IAB 12/12/12, effective 1/16/13]
- [Filed ARC 1785C (Notice ARC 1653C, IAB 10/1/14), IAB 12/10/14, effective 1/14/15]
- [Filed ARC 2194C (Notice ARC 1979C, IAB 4/29/15), IAB 10/14/15, effective 11/18/15]
- [Filed ARC 3857C (Notice ARC 3506C, IAB 12/20/17), IAB 6/20/18, effective 7/25/18]
- [Filed ARC 4189C (Notice ARC 4030C, IAB 9/26/18), IAB 12/19/18, effective 1/23/19]
- [Filed ARC 5007C (Notice ARC 4695C, IAB 10/9/19), IAB 3/25/20, effective 4/29/20]
- [Filed ARC 5542C (Notice ARC 5373C, IAB 1/13/21), IAB 4/7/21, effective 5/12/21]
- [Filed Emergency ARC 5820C, IAB 8/11/21, effective 7/15/21]
- [Filed ARC 6071C (Notice ARC 5831C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]
- [Filed ARC 6076C (Notice ARC 5833C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]
- [Filed ARC 6416C (Notice ARC 6279C, IAB 4/6/22), IAB 7/27/22, effective 8/31/22]
- [Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

¹ April 30, 2008, effective date delayed 70 days by the Administrative Rules Review Committee at its meeting held April 4, 2008.

CHAPTER 6
GENERAL PHARMACY PRACTICE
[Prior to 2/10/88, see Pharmacy Examiners[620] Ch 2]

657—6.1(155A) Purpose and scope. A general pharmacy is a location where a pharmacist provides pharmaceutical services or dispenses pharmaceutical products to patients in accordance with pharmacy laws. This chapter does not apply to a hospital pharmacy as defined in 657—Chapter 7. The requirements of these rules for general pharmacy practice are in addition to the requirements of 657—Chapter 8 and other rules of the board relating to services provided by the pharmacy.

657—6.2(155A) Pharmacist in charge. One professionally competent, legally qualified pharmacist in charge in each pharmacy shall be responsible for, at a minimum, the responsibilities identified in rule 657—8.3(155A).

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 0501C, IAB 12/12/12, effective 1/16/13; ARC 1961C, IAB 4/15/15, effective 5/20/15]

657—6.3(155A) Reference library. References may be printed or computer-accessed. A reference library shall be maintained which includes, at a minimum, one current reference from each of the following categories, including access to current periodic updates.

1. A reference including all pertinent Iowa laws, rules, and regulations that impact the pharmacy's practice.
2. A patient information reference that includes or provides patient information in compliance with rule 657—6.14(155A).
3. A reference on drug interactions.
4. A general information reference.
5. A drug equivalency reference.
6. A reference on natural or herbal medicines.
7. The readily accessible telephone number of a poison control center that serves the area.
8. Additional references as may be necessary for the pharmacist to adequately meet the needs of the patients served.

[ARC 2196C, IAB 10/14/15, effective 11/18/15]

657—6.4(155A) Exemption from duplicate requirements. A pharmacy established in the same location as another licensed pharmacy and with direct and immediate access to required references, patient counseling area, refrigerator, or sink with hot and cold running water may utilize the references, counseling area, refrigerator, or sink of the other pharmacy to satisfy the requirements of rule 657—6.3(155A), subrule 6.14(3), or rule 657—8.5(155A), paragraphs "1" and "2."

657—6.5 and 6.6 Reserved.

657—6.7(124,155A) Security. While on duty, each pharmacist shall be responsible for the security of the prescription department and of the provisions for effective control against theft of, diversion of, or unauthorized access to prescription drugs, including those collected through an authorized collection program, records for such drugs and authorized collection program activities, and patient records as provided in 657—Chapters 10 and 21 and federal regulations for authorized controlled substance collection programs, which can be found at www.deadiversion.usdoj.gov/drug_disposal/.

6.7(1) Department locked. The prescription department shall be locked by key or combination so as to prevent access when a pharmacist is not on site except as provided in subrules 6.7(2) and 6.7(4).

6.7(2) Temporary absence of pharmacist. In the temporary absence of the pharmacist, only the pharmacist in charge may designate pharmacy technicians or pharmacy support persons who may be present in the prescription department to perform technical or nontechnical functions, respectively, designated by the pharmacist in charge. Activities identified in subrule 6.7(3) may not be performed during such temporary absence of the pharmacist. A temporary absence is an absence of short duration not to exceed two hours.

a. In the absence of the pharmacist, the pharmacy shall be secured from public access and the pharmacy shall notify the public that the pharmacist is temporarily absent and that no prescriptions will be dispensed until the pharmacist returns. If the pharmacist in charge has authorized the presence in the pharmacy of a pharmacy technician or a pharmacy support person to perform designated functions when the pharmacy is closed, the pharmacy technician or the pharmacy support person may not dispense or deliver any drug, chemical, device, or prepared prescription to a patient or patient's agent.

b. A pharmacy technician or a pharmacy support person who is present in the pharmacy when the pharmacy is closed shall prepare and maintain in the pharmacy a log identifying each period of time that the pharmacy technician or pharmacy support person worked in the pharmacy while the pharmacy was closed and identifying each activity performed during that time period. Each entry shall be dated, and each daily record shall be signed by the pharmacy technician or pharmacy support person who prepared the record. The log shall be periodically reviewed by the pharmacist in charge, and documentation of such review shall be maintained for two years from the date of entry.

6.7(3) *Activities prohibited in absence of pharmacist.* Activities which shall not be designated and shall not be performed during the temporary absence of the pharmacist include:

- a. Dispensing or distributing any prescription drugs or devices to patients or others.
- b. Providing the final verification for the accuracy, validity, completeness, or appropriateness of a filled prescription or medication order.
- c. Conducting prospective drug use review or evaluating a patient's medication record for purposes identified in rule 657—8.21(155A).
- d. Providing patient counseling, consultation, or drug information.
- e. Making decisions that require a pharmacist's professional judgment such as interpreting or applying information.
- f. Transferring prescriptions to or from other pharmacies.

6.7(4) *Refill sales during pharmacist break.* At the discretion of the on-duty supervising pharmacist and pursuant to established policies and procedures, the pharmacist may delegate to a technician the dispensing of previously verified prescriptions which have been identified to not require pharmacist counseling pursuant to rule 657—6.14(155A) when the pharmacist is on a break of limited duration and is absent from the pharmacy department.

6.7(5) *Minimum physical security and monitoring system requirements.* Each pharmacy located in Iowa shall develop and implement policies and procedures to ensure appropriate physical security and monitoring of the pharmacy to prevent unauthorized access to prescription drugs, including controlled substances, and pharmacy records. The physical security and monitoring shall include the components identified herein, and the policies and procedures shall establish the utilization of such components commensurate with the pharmacy operation. The policies and procedures shall establish the retention of documentation of activities or recordings retained from the alarm and video surveillance systems, as well as contingencies when the systems are temporarily unavailable.

- a. No later than July 6, 2023, a basic alarm system.
- b. No later than July 6, 2023, a video surveillance system, except in areas where drugs are stored in an automated medication dispensing system or an alternative electronic storage unit which uses biometric restricted access or other electronic monitoring mechanism.
- c. Controlled access to computer records.
- d. A designated location that can be monitored, away from drug storage and handling areas, where personal items of pharmacy staff may be stored while on site.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 1308C, IAB 2/5/14, effective 3/12/14; ARC 2408C, IAB 2/17/16, effective 3/23/16; ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 4189C, IAB 12/19/18, effective 1/23/19; ARC 6330C, IAB 6/1/22, effective 7/6/22]

657—6.8(124,155A) Prescription processing documentation. All prescriptions shall be dated and assigned a unique identification number that shall be recorded on the original prescription, except as provided in 657—subrule 21.5(1). The original prescription shall be retained by the pharmacy filling the prescription and shall be maintained in the original format as received by the pharmacy. Dispensing documentation shall include the date of fill or refill; the name, strength, and National Drug Code (NDC)

of the actual drug product dispensed; and the initials or other unique identification of the pharmacist, pharmacist-intern, or technician in an approved technician product verification program. Dispensing documentation shall be maintained and be readily available.

[ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 5007C, IAB 3/25/20, effective 4/29/20]

657—6.9(124,155A) Transfer of prescription. The transmission of a prescription drug order from a pharmacy to a pharmacy engaged in centralized prescription filling or processing on behalf of the originating pharmacy pursuant to the requirements of 657—Chapter 18 shall not constitute the transfer of a prescription. Upon the request of a patient or the patient's caregiver, a pharmacy shall transfer original prescription drug order information and prescription refill information to a pharmacy designated by the patient or the patient's caregiver, central fill or processing pharmacies excepted, subject to the following requirements:

6.9(1) *Schedule III, IV, or V prescriptions.* The transfer of original prescription drug order information for controlled substances listed in Schedule III, IV, or V is permissible between pharmacies on a one-time basis except as provided in subrule 6.9(8).

6.9(2) *Noncontrolled substances prescriptions.* The transfer of original prescription drug order information for noncontrolled prescription drugs between pharmacies is permissible as long as the number of transfers does not exceed the number of originally authorized refills and the original prescription is still valid.

6.9(3) *Authorized individuals and means of transmission.* Individuals authorized to engage in the transfer of prescriptions include a pharmacist, a pharmacist-intern under the direct supervision of a pharmacist, and a certified pharmacy technician, except as prohibited in 657—subrule 3.23(1). The transferring individual may transmit the prescription and transfer information required under subrule 6.9(5) from the transferring pharmacy via electronic means pursuant to subrule 6.9(8) or, following direct communication between authorized individuals, via oral or facsimile transmission. The receiving individual shall ensure the prescription transfer record maintained in the receiving pharmacy contains all of the information required under subrule 6.9(7).

6.9(4) *Prescriptions maintained.* Both the original and the transferred prescription drug orders are maintained for a period of two years from the date of last activity.

6.9(5) *Record of transfer out.* The individual transferring the prescription drug order information shall:

- a. Invalidate the prescription drug order;
- b. Record on or with the invalidated prescription drug order the following information:
 - (1) The name, address, and, for a controlled substance, the DEA registration number of the pharmacy to which such prescription is transferred;
 - (2) The name of the individual receiving the prescription drug order information;
 - (3) The name of the individual transferring the prescription drug order information; and
 - (4) The date of the transfer.

6.9(6) *Original prescription status.* The original prescription drug order shall be invalidated in the data processing system for purposes of filling or refilling, but shall be maintained in the data processing system for refill history purposes.

6.9(7) *Record of transfer received.* The individual receiving the transferred prescription drug order information shall:

- a. Indicate that the prescription drug order has been transferred;
- b. Record on or with the transferred prescription drug order the following information:
 - (1) Original date of issuance and date of dispensing, if different from date of issuance;
 - (2) Original prescription number;
 - (3) Number of valid refills remaining, the date of last refill, and, for a controlled substance, the dates and locations of all previous refills;
 - (4) Name, address, and, for a controlled substance, the DEA registration number of the pharmacy from which such prescription drug order information is transferred;
 - (5) The date of the transfer;

- (6) Name of the individual receiving the prescription drug order information;
- (7) Name of the individual transferring the prescription drug order information; and
- (8) If transferring a controlled substance prescription from a pharmacy utilizing a shared electronic database system as described in subrule 6.9(8) to a pharmacy outside that shared system, the pharmacy name, location, DEA registration number, and prescription number from which the prescription was originally filled.

6.9(8) *Electronic transfer between pharmacies.* Pharmacies may electronically transfer prescription information, including controlled substance prescription information in compliance with federal regulations for controlled substances. For transfers of prescriptions for noncontrolled substances and controlled substances, pharmacies that share a real-time, online database may transfer up to the maximum refills permitted by law and the prescriber's authorization. A prescription for a controlled substance transferred between two pharmacies which do not share a real-time, online database may only be transferred one time.

[ARC 7634B, IAB 3/11/09, effective 4/15/09; ARC 8169B, IAB 9/23/09, effective 10/28/09; ARC 0343C, IAB 10/3/12, effective 11/7/12; ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 4189C, IAB 12/19/18, effective 1/23/19; ARC 5542C, IAB 4/7/21, effective 5/12/21]

657—6.10(126,155A) Prescription label requirements.

6.10(1) *Required information.* The label affixed to or on the dispensing container of any prescription drug or device dispensed by a pharmacy pursuant to a prescription drug order shall bear the following:

- a. Serial number (a unique identification number of the prescription);
- b. The name, telephone number, and address of the pharmacy;
- c. The name of the patient or, if such drug is prescribed for an animal, the species of the animal and the name of its owner, except as provided in 657—subrule 8.19(7) for epinephrine auto-injectors, 657—subrule 8.19(8) for opioid antagonists, 657—subrule 8.19(9) for expedited partner therapy, or 657—subrule 8.19(10) for bronchodilator canisters or bronchodilator canisters and spacers.
- d. The name of the prescribing practitioner;
- e. The date the prescription is dispensed;
- f. The directions or instructions for use, including precautions to be observed;
- g. Unless otherwise directed by the prescriber, the label shall bear the name, strength, and quantity of the drug dispensed.

(1) If a pharmacist selects an equivalent drug product for a brand name drug product prescribed by a practitioner, the prescription container label shall identify the generic drug and may identify the brand name drug for which the selection is made, such as “(generic name) Generic for (brand name product)”;

(2) If a pharmacist selects a brand name drug product for a generic drug product prescribed by a practitioner, the prescription container label shall identify the brand name drug product dispensed and may identify the generic drug product ordered by the prescriber, such as “(brand name product) for (generic name)”;

(3) If a pharmacist selects an interchangeable biological product for the biological product prescribed by a practitioner, the prescription container label shall identify the interchangeable biological product dispensed and may identify the biological product prescribed by the practitioner, such as “(interchangeable biological product) for (biological product)”;

h. The initials or other unique identification of the dispensing pharmacist, unless the identification of the pharmacist involved in each step of the prescription filling process is electronically documented and retrievable.

6.10(2) *Exceptions.* The requirements of subrule 6.10(1) do not apply to unit dose dispensing systems, 657—22.1(155A), and patient med paks, 657—22.5(126,155A).

[ARC 2194C, IAB 10/14/15, effective 11/18/15; ARC 2414C, IAB 2/17/16, effective 3/23/16; ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 4903C, IAB 2/12/20, effective 3/18/20; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—6.11 and 6.12 Reserved.

657—6.13(155A) Patient record system.

6.13(1) Information required. A patient record system shall be maintained by all pharmacies for patients for whom prescription drug orders are dispensed. The patient record system shall contain, at a minimum, the following information:

- a. Full name of the patient;
- b. Address and telephone number of the patient;
- c. Patient's date of birth;
- d. Patient's gender;
- e. Known allergies;
- f. A list of all prescription drug orders dispensed by the pharmacy during the two years immediately preceding the most recent entry showing the name of the drug or device, prescription number, name and strength of the drug, the quantity and date dispensed, and the name of the prescriber; and
- g. Pharmacist comments relevant to the patient's health care, including:
 - (1) Known drug reactions,
 - (2) Identified idiosyncrasies,
 - (3) Known chronic conditions or disease states of the patient,
 - (4) The identity of any other drugs, over-the-counter drugs, herbals, supplements, other alternative medications, or devices currently being used by the patient that may relate to prospective drug review.

6.13(2) Record retained. A patient record shall be maintained for a period of not less than two years from the date of the last entry in the patient record. This record may be a hard copy or a computerized form.

6.13(3) Confidential. Information in the patient record shall be deemed to be confidential and may be released only as provided in rule 657—8.16(124,155A).

6.13(4) Expedited partner therapy. When a pharmacy dispenses a prescription drug pursuant to Iowa Code section 139A.41 and 657—subrule 8.19(9) for expedited partner therapy, a pharmacy is only required to maintain the information about the patient who is known to the pharmacy.

[ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 4903C, IAB 2/12/20, effective 3/18/20]

657—6.14(155A) Patient counseling and instruction. Every pharmacy that is open to the public and located in Iowa shall post in every prescription pickup area, including in every drive-through prescription pickup lane, in a manner clearly visible to patients, a notice that Iowa law requires the pharmacist to discuss with the patient any prescriptions dispensed to the patient that are new or a change in drug therapy.

6.14(1) Counseling required. Upon receipt of a new prescription drug order, or upon receipt of a change in drug therapy including but not limited to a change of dose, directions, or drug formulation, and following a prospective drug use review pursuant to rule 657—8.21(155A), a pharmacist or pharmacist-intern shall counsel each patient or patient's caregiver. An offer to counsel shall not fulfill the requirements of this rule. Patient counseling shall be on matters which, in the pharmacist's professional judgment, will enhance or optimize drug therapy. Appropriate elements of patient counseling may include:

- a. The name and description of the drug;
- b. The dosage form, dose, route of administration, and duration of drug therapy;
- c. Intended use of the drug, if known, and expected action;
- d. Special directions and precautions for preparation, administration, and use by the patient;
- e. Common severe side effects or adverse effects or interactions and therapeutic contraindications that may be encountered, including their avoidance, and the action required if they occur;
- f. Techniques for self-monitoring drug therapy;
- g. Proper storage;
- h. Prescription refill information;
- i. Action to be taken in the event of a missed dose;
- j. Pharmacist comments relevant to the individual's drug therapy including any other information peculiar to the specific patient or drug.

6.14(2) Instruction. A pharmacist may instruct patients and demonstrate procedures for self-monitoring of medical conditions and for self-administration of drugs.

6.14(3) Counseling area. A pharmacy shall contain an area which is suitable for confidential patient counseling. Such area shall:

a. Be easily accessible to both patient and pharmacists and not allow patient access to prescription drugs;

b. Be designed to maintain the confidentiality and privacy of the pharmacist/patient communication.

6.14(4) Remote counseling. Patient counseling that is provided by the pharmacy via a pharmacist who is at a location other than the licensed pharmacy shall be provided via a real-time interactive communication mechanism.

6.14(5) Oral counseling not practicable. If in the pharmacist's professional judgment oral counseling is not practicable, the pharmacist may select and use alternative forms of patient information which shall include information for the patient or patient's caregiver to contact the pharmacist for further consultation. The manner in which the patient or caregiver contacts the pharmacist shall not cause the patient to incur any expense. "Not practicable" refers to patient variables including, but not limited to, the absence of the patient or patient's caregiver, the patient's or caregiver's hearing disorder, or a language barrier. "Not practicable" does not include pharmacy variables such as inadequate staffing, technology failure, or high prescription volume. A combination of oral counseling and alternative forms of counseling is encouraged.

6.14(6) Exception. Patient counseling, as described above, shall not be required for inpatients of an institution where other licensed health care professionals are authorized to administer the drugs.

6.14(7) Refusal of consultation. A pharmacist shall not be required to counsel a patient or caregiver when the patient or caregiver refuses such consultation. A patient's or caregiver's refusal of consultation shall be documented by the pharmacist. The absence of any record of a refusal of the pharmacist's attempt to counsel shall be presumed to signify that counseling was provided.

[ARC 8540B, IAB 2/24/10, effective 4/1/10; ARC 9910B, IAB 12/14/11, effective 1/18/12; ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 5913C, IAB 9/22/21, effective 10/27/21; ARC 6416C, IAB 7/27/22, effective 8/31/22]

657—6.15(124,126) Return of drugs and devices. For the protection of the public health and safety, prescription drugs and devices may be returned to the pharmacy for reuse or resale only as herein provided:

6.15(1) Integrity maintained. Prescription drugs and devices may be returned, exchanged, or resold only if, in the professional judgment of the pharmacist, the integrity of the prescription drug or device has not in any way been compromised.

6.15(2) Controlled substances. Under no circumstances shall pharmacy personnel accept from a patient or a patient's agent any controlled substances for return, exchange, or resale except to the same patient.

6.15(3) Unit dose returns. Prescription drugs dispensed in unit dose packaging, excluding controlled substances, may be returned and reused as authorized in 657—subrule 22.1(6).
[ARC 3638C, IAB 2/14/18, effective 3/21/18]

657—6.16(124,155A) Records. Every record required to be kept under Iowa Code chapters 124 and 155A or rules of the board shall be kept by the pharmacy and be available for inspection and copying by the board or its representative for at least two years from the date of the record or last activity except as specifically identified by law or rule. Controlled substances records shall be maintained in a readily retrievable manner in accordance with federal requirements and 657—Chapter 10.

6.16(1) Combined records. If controlled substances, prescription drugs, or nonprescription drug items are listed on the same record, the controlled substances shall be asterisked, red-lined, or in some other manner made readily identifiable from all other items appearing on the records.

6.16(2) Storage of records. Original hard-copy prescriptions and other pharmacy records shall be maintained by the pharmacy for a minimum of two years from the date of the record in accordance with this subrule.

a. Records shall be maintained within the licensed pharmacy department for a minimum of 12 months, except as provided herein. Pharmacy records less than 12 months old may be stored in a secure storage area outside the licensed pharmacy department, including at a remote location, if the pharmacy has retained an electronic copy of the records in the pharmacy that is immediately available and if the original records are available within 72 hours of a request by the board or its authorized agent, unless such remote storage is prohibited under federal law.

b. Records more than 12 months old may be maintained in a secure storage area outside the licensed pharmacy department, including at a remote location, if the records are retrievable within 72 hours of a request by the board or its authorized agent, unless such remote storage is prohibited under federal law.

6.16(3) *Number imprinted.* The original hard-copy prescription shall be imprinted with the prescription or control number assigned to the prescription drug order, except as provided in 657—subrule 21.5(1).

6.16(4) *Alternative data retention system.* Records, except when specifically required to be maintained in original or hard-copy form, may be maintained in an alternative data retention system, such as a data processing system or direct imaging system provided:

a. The records maintained in the alternative system contain all of the information required on the manual record;

b. The data processing system is capable of producing a hard copy of the record, within two business days, upon the request of the board, its representative, or other authorized local, state, or federal law enforcement or regulatory agencies; and

c. The information maintained in the alternative system is not obscured or rendered illegible due to security features of the original record.

[ARC 7636B, IAB 3/11/09, effective 4/15/09; ARC 8539B, IAB 2/24/10, effective 4/1/10; ARC 3638C, IAB 2/14/18, effective 3/21/18; ARC 5007C, IAB 3/25/20, effective 4/29/20]

These rules are intended to implement Iowa Code sections 124.301, 124.303, 124.306, 126.10, 126.11, 155A.6, 155A.13, 155A.27, 155A.28, 155A.31, and 155A.33 through 155A.36.

[Filed 5/16/67; amended 11/14/73]

[Filed 6/1/84, Notice 3/14/84—published 6/20/84, effective 7/25/84]

[Filed 5/14/86, Notice 4/9/86—published 6/4/86, effective 7/9/86]

[Filed 1/28/87, Notice 11/19/86—published 2/25/87, effective 4/1/87]

[Filed 11/25/87, Notice 10/7/87—published 12/16/87, effective 1/20/88]

[Filed emergency 1/21/88—published 2/10/88, effective 1/22/88]

[Filed 11/17/88, Notice 8/24/88—published 12/14/88, effective 1/18/89]

[Filed emergency 5/16/89—published 6/14/89, effective 5/17/89]

[Filed 9/12/89, Notice 6/14/89—published 10/4/89, effective 11/8/89]

[Filed emergency 5/10/91—published 5/29/91, effective 5/10/91]

[Filed 7/30/91, Notice 5/29/91—published 8/21/91, effective 9/25/91]

[Filed 9/23/93, Notice 5/26/93—published 10/13/93, effective 11/17/93]

[Filed 3/21/94, Notice 10/13/93—published 4/13/94, effective 5/18/94]

[Filed 2/27/97, Notice 1/1/97—published 3/26/97, effective 4/30/97]

[Filed 9/16/97, Notice 7/16/97—published 10/8/97, effective 11/12/97]

[Filed 4/24/98, Notice 3/11/98—published 5/20/98, effective 6/24/98]

[Filed 2/22/99, Notices 10/21/98—published 3/10/99, effective 4/14/99]^o

[Filed 4/22/99, Notice 3/10/99—published 5/19/99, effective 6/23/99]

[Filed 9/8/99, Notice 6/2/99—published 10/6/99, effective 11/10/99]

[Filed 2/7/01, Notice 10/18/00—published 3/7/01, effective 4/11/01]

[Filed 8/14/02, Notice 6/12/02—published 9/4/02, effective 10/9/02]

[Filed emergency 3/26/03 after Notice 11/13/02—published 4/16/03, effective 3/26/03]

[Filed 7/15/03, Notice 4/16/03—published 8/6/03, effective 9/10/03]

[Filed 10/22/04, Notice 3/31/04—published 11/10/04, effective 12/15/04]

[Filed 6/2/05, Notice 1/19/05—published 6/22/05, effective 7/27/05]

[Filed 6/2/05, Notice 3/16/05—published 6/22/05, effective 7/27/05]
[Filed 3/22/06, Notice 12/21/05—published 4/12/06, effective 5/17/06]
[Filed 3/22/06, Notice 1/18/06—published 4/12/06, effective 5/17/06]
[Filed 2/7/07, Notice 10/25/06—published 2/28/07, effective 4/4/07]
[Filed 8/2/07, Notice 6/20/07—published 8/29/07, effective 10/3/07]
[Filed 3/5/08, Notice 12/5/07—published 3/26/08, effective 4/30/08]
[Filed 11/24/08, Notice 10/8/08—published 12/17/08, effective 1/21/09]
[Filed ARC 7634B (Notice ARC 7447B, IAB 12/31/08), IAB 3/11/09, effective 4/15/09]
[Filed ARC 7636B (Notice ARC 7448B, IAB 12/31/08), IAB 3/11/09, effective 4/15/09]
[Filed ARC 8169B (Notice ARC 7926B, IAB 7/1/09), IAB 9/23/09, effective 10/28/09]
[Filed ARC 8540B (Notice ARC 8267B, IAB 11/4/09), IAB 2/24/10, effective 4/1/10]
[Filed ARC 8539B (Notice ARC 8269B, IAB 11/4/09), IAB 2/24/10, effective 4/1/10]
[Filed ARC 8673B (Notice ARC 8380B, IAB 12/16/09), IAB 4/7/10, effective 6/1/10]
[Filed ARC 9910B (Notice ARC 9787B, IAB 10/5/11), IAB 12/14/11, effective 1/18/12]
[Filed ARC 0343C (Notice ARC 0155C, IAB 6/13/12), IAB 10/3/12, effective 11/7/12]
[Filed ARC 0501C (Notice ARC 0375C, IAB 10/3/12), IAB 12/12/12, effective 1/16/13]
[Filed ARC 1308C (Notice ARC 1040C, IAB 10/2/13), IAB 2/5/14, effective 3/12/14]
[Filed ARC 1961C (Notice ARC 1793C, IAB 12/10/14), IAB 4/15/15, effective 5/20/15]
[Filed ARC 2194C (Notice ARC 1979C, IAB 4/29/15), IAB 10/14/15, effective 11/18/15]
[Filed ARC 2196C (Notice ARC 2065C, IAB 7/22/15), IAB 10/14/15, effective 11/18/15]
[Filed ARC 2408C (Notice ARC 2285C, IAB 12/9/15), IAB 2/17/16, effective 3/23/16]
[Filed ARC 2414C (Notice ARC 2288C, IAB 12/9/15), IAB 2/17/16, effective 3/23/16]
[Filed ARC 3638C (Notice ARC 3330C, IAB 9/27/17), IAB 2/14/18, effective 3/21/18]
[Filed ARC 4189C (Notice ARC 4030C, IAB 9/26/18), IAB 12/19/18, effective 1/23/19]
[Filed ARC 4903C (Notice ARC 4693C, IAB 10/9/19), IAB 2/12/20, effective 3/18/20]
[Filed ARC 5007C (Notice ARC 4695C, IAB 10/9/19), IAB 3/25/20, effective 4/29/20]
[Filed ARC 5542C (Notice ARC 5373C, IAB 1/13/21), IAB 4/7/21, effective 5/12/21]
[Filed ARC 5913C (Notice ARC 5704C, IAB 6/16/21), IAB 9/22/21, effective 10/27/21]
[Filed ARC 6330C (Amended Notice ARC 6179C, IAB 2/9/22; Notice ARC 5834C, IAB 8/11/21),
IAB 6/1/22, effective 7/6/22]
[Filed ARC 6416C (Notice ARC 6279C, IAB 4/6/22), IAB 7/27/22, effective 8/31/22]
[Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

◊ Two or more ARCs

CHAPTER 8
UNIVERSAL PRACTICE STANDARDS

[Prior to 2/10/88, see Pharmacy Examiners[620] Ch 6]

657—8.1(155A) Purpose and scope. The purpose of this chapter is to establish the minimum standards of pharmacy practice for the activities identified in this chapter. The requirements of these rules shall apply to all Iowa-licensed pharmacists, other registered pharmacy personnel, and all pharmacies, including owners, providing the services addressed in this chapter to patients in Iowa. These rules are in addition to rules of the board relating to specific types of pharmacy licenses issued by the board unless otherwise indicated by rule.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.2(155A) Definitions. For the purpose of this chapter, the following definitions shall apply:

“*Board*” means the Iowa board of pharmacy.

“*Confidential information*” means information accessed or maintained by the pharmacy in the patient’s or the pharmacy’s records which contains personally identifiable information that could be used to identify the patient. “Confidential information” includes but is not limited to patient name, address, telephone number, and social security number; prescriber name and address; and prescription and drug or device information such as therapeutic effect, diagnosis, allergies, disease state, pharmaceutical services rendered, medical information, and drug interactions.

“*DEA*” means the United States Department of Justice, Drug Enforcement Administration.

“*Pharmacy support person*” or “*PSP*” means a person, other than a member of the professional pharmacy staff, registered with the board who may perform nontechnical duties assigned by a supervising pharmacist under the pharmacist’s responsibility and supervision.

“*Professional pharmacy staff*” shall mean the professional employees of the pharmacy, including pharmacists, pharmacy technicians, and pharmacist-interns.

This rule is intended to implement Iowa Code chapter 155A.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.3(155A) Responsible parties.

8.3(1) Pharmacist in charge. One professionally competent, legally qualified pharmacist in charge in each pharmacy shall work cooperatively with the pharmacy, by and through its owner or license holder, and with all staff pharmacists to ensure the legal operation of the pharmacy, including meeting all inspection and other requirements of state and federal laws, rules, and regulations governing the practice of pharmacy. A part-time pharmacist in charge has the same obligations and responsibilities as a full-time pharmacist in charge.

8.3(2) Pharmacy. Each pharmacy, by and through its owner or license holder, shall work cooperatively with the pharmacist in charge and with all staff pharmacists to ensure the legal operation of the pharmacy, including meeting all inspection and other requirements of state and federal laws, rules, and regulations governing the practice of pharmacy. The pharmacy, by and through its owner or license holder, shall be responsible for employing a professionally competent, legally qualified pharmacist in charge. The pharmacy, by and through its owner or license holder, may be held responsible for unethical conduct or practices of any of the pharmacy staff.

8.3(3) Pharmacy and pharmacist in charge. The pharmacist in charge and the pharmacy, by and through its owner or license holder, shall share responsibility for, at a minimum, the following:

a. Ensuring that the pharmacy employs an adequate number of qualified personnel commensurate with the size and scope of services provided by the pharmacy.

b. Ensuring the availability of any equipment and references necessary for the particular practice of pharmacy.

c. Ensuring that there is adequate space within the prescription department or a locked room not accessible to the public for the storage of prescription drugs, including controlled substances, devices, and pharmacy records, and to support the operations of the pharmacy.

d. Ensuring that the license, registration, or certification of each professional pharmacy staff member and the registration of each pharmacy support person are maintained in current and active status.

e. Ensuring that the pharmacy provides adequate security to prevent unauthorized access and diversion.

8.3(4) *Pharmacist in charge and staff pharmacists.* The pharmacist in charge and staff pharmacists shall share responsibility for, at a minimum, the following:

a. Ensuring that a pharmacist performs prospective drug use review as specified in rule 657—8.21(155A).

b. Ensuring that a pharmacist or pharmacist-intern provides patient counseling as specified in rule 657—6.14(155A).

c. Dispensing drugs to patients, including the packaging, preparation, compounding, and labeling functions performed by pharmacy personnel.

d. Delivering drugs to the patient or the patient's agent.

e. Ensuring that patient medication records are maintained as specified in rule 657—6.13(155A).

f. Training and supervising pharmacist-interns, pharmacy technicians, pharmacy support persons, and other pharmacy employees.

g. Procuring and storing prescription drugs and devices and other products dispensed from the pharmacy.

h. Distributing and disposing of drugs from the pharmacy.

i. Maintaining records of all transactions of the pharmacy necessary to maintain accurate control over and accountability for all drugs as required by applicable state and federal laws, rules, and regulations.

j. Ensuring the legal operation of the pharmacy, including meeting all inspection and other requirements of state and federal laws, rules, and regulations governing the practice of pharmacy.

8.3(5) *Pharmacy, pharmacist in charge, and staff pharmacists.* The pharmacy, by and through its owner or license holder, the pharmacist in charge, and all staff pharmacists shall share responsibility for, at a minimum, the following:

a. Establishing and periodically reviewing (by the pharmacy and the pharmacist in charge), implementing (by the pharmacist in charge), and complying (by the pharmacist in charge and staff pharmacists) with policies and procedures for all operations of the pharmacy. The policies and procedures shall identify the frequency of review.

b. Establishing and maintaining effective controls against the theft or diversion of prescription drugs, including controlled substances, and records for such drugs.

c. Establishing (by the pharmacy and the pharmacist in charge), implementing (by the pharmacist in charge), and utilizing (by the pharmacist in charge and staff pharmacists) an ongoing, systematic program of continuous quality improvement for achieving performance enhancement and ensuring the quality of pharmaceutical services.

8.3(6) *Practice functions.* The pharmacist is responsible for all functions performed in the practice of pharmacy. The pharmacist maintains responsibility for any and all delegated functions including functions delegated to pharmacist-interns, pharmacy technicians, and pharmacy support persons.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 1576C, IAB 8/20/14, effective 9/24/14; ARC 1961C, IAB 4/15/15, effective 5/20/15; ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 6330C, IAB 6/1/22, effective 7/6/22]

657—8.4(155A) Pharmacist identification and staff logs.

8.4(1) *Display of pharmacist license.* During any period a pharmacist is working in a pharmacy, each pharmacist shall display, in a position visible to the public, an original license to practice pharmacy in Iowa. A current license renewal certificate, which may be a photocopy of an original renewal certificate, shall be displayed with the original license.

8.4(2) *Registration maintained of pharmacy personnel.* Each pharmacist-intern, pharmacy technician, and pharmacy support person shall maintain current registration with the board. The

registration certificate or a copy of the registration certificate shall be readily retrievable upon request of the board or its authorized agent.

8.4(3) *Identification codes.* A permanent log of the initials or identification code identifying by name each pharmacist, pharmacist-intern, pharmacy technician, and pharmacy support person shall be maintained for a minimum of two years and shall be available for inspection and copying by the board or its representative. The initials or identification code shall be unique to the individual to ensure that each pharmacist, pharmacist-intern, pharmacy technician, and pharmacy support person can be identified.

8.4(4) *Temporary or intermittent pharmacy staff.* The pharmacy shall maintain a log of all pharmacists, pharmacist-interns, pharmacy technicians, and pharmacy support persons who have worked at that pharmacy and who are not regularly staffed at that pharmacy. Such log shall include the dates and shifts worked by each pharmacist, pharmacist-intern, pharmacy technician, and pharmacy support person and shall be available for inspection and copying by the board or its representative for a minimum of two years following the date of the entry.

8.4(5) *Identification.* While on duty, pharmacy personnel shall wear visible identification that clearly identifies the person by licensed or registered title and includes at least the person's first name.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9409B, IAB 3/9/11, effective 4/13/11; ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.5(155A) *Environment and equipment requirements.* There shall be adequate space, equipment, and supplies for the professional and administrative functions of the pharmacy pursuant to rule 657—8.3(155A). Space and equipment shall be available in an amount and type to provide secure, environmentally controlled storage of drugs.

8.5(1) *Refrigeration.* The pharmacy shall maintain one or more refrigeration units, unless the pharmacy does not stock refrigerated items. The pharmacy shall document verification that the temperature of the refrigerator is maintained within a range compatible with the proper storage of drugs requiring refrigeration. If the temperature is manually or visually verified, a record of minimum daily verification shall be maintained.

8.5(2) *Sink.* The pharmacy shall have a sink with hot and cold running water located within the pharmacy department and available to all pharmacy personnel; the sink shall be maintained in a sanitary condition.

8.5(3) *Secure barrier.* A pharmacy department shall be closed and secured in the absence of the pharmacist except as provided in rule 657—6.7(124,155A) or 657—7.5(124,155A). To ensure that secure closure, the pharmacy department shall be surrounded by a physical barrier capable of being securely locked to prevent entry when the department is closed. A secure barrier may be constructed of other than a solid material with a continuous surface if the openings in the material are not large enough to permit removal of items from the pharmacy department by any means. Any material used in the construction of the barrier shall be of sufficient strength and thickness that it cannot be readily or easily removed, penetrated, or bent.

8.5(4) *Remodel or relocation—inspection.* A pharmacy planning to remodel or relocate a licensed pharmacy department on or within the premises currently occupied by the pharmacy department, or a pharmacy intending to remodel or install a sterile compounding facility or equipment, shall provide written notification to the board at least 30 days prior to commencement of the remodel, pharmacy relocation, or sterile compounding installation. The board may require on-site inspection of the facility, equipment, or pharmacy department prior to or during the pharmacy's remodel, relocation, or opening. The board may also require on-site inspection of a temporary pharmacy location intended to be utilized during the remodel, construction, or relocation of the pharmacy department.

8.5(5) *Orderly and clean.* The pharmacy shall be arranged in an orderly fashion and kept clean. All required equipment shall be in good operating condition and maintained in a sanitary manner. Animals shall not be allowed within a licensed pharmacy unless that pharmacy is exclusively providing services for the treatment of animals or unless the animal is a service animal or service-animal-in-training as defined in Iowa Code section 216C.1A.

8.5(6) *Light, ventilation, temperature, and humidity.* The pharmacy shall be properly lighted and ventilated. The temperature and humidity of the pharmacy shall be maintained within a range compatible with the proper storage of drugs.

8.5(7) *Other equipment.* The pharmacist in charge and the pharmacy, by and through its owner or license holder, shall share the responsibility for ensuring the availability of any other equipment necessary for the particular practice of pharmacy and to meet the needs of the patients served by the pharmacy.

8.5(8) *Bulk counting machines.* Unless bar-code scanning is required and utilized to verify the identity of each stock container of drugs utilized to restock a counting machine cell or bin, a pharmacist shall verify the accuracy of the drugs to be restocked prior to filling the counting machine cell or bin. A record identifying the individual who verified the drugs to be restocked, the individual who restocked the counting machine cell or bin, and the date shall be maintained. Established policies and procedures shall include a method to calibrate and verify the accuracy of the counting device. The pharmacy shall, at least quarterly, verify the accuracy of the device and maintain a dated record identifying the individual who performed the quarterly verification.

8.5(9) *Authorized collection program.* A pharmacy that is registered with the DEA to administer an authorized collection program shall provide adequate space, equipment, and supplies for such collection program pursuant to 657—Chapter 10 and federal regulations for authorized collection programs, which can be found at www.deadiversion.usdoj.gov/drug_disposal/.

8.5(10) *Health of personnel.* The pharmacist in charge or supervising pharmacist shall ensure that pharmacy personnel experiencing any health condition that may have an adverse effect on drug products or may pose a health or safety risk to others be prohibited from working in the pharmacy until such health condition is sufficiently resolved. All personnel who normally assist the pharmacist shall report to the pharmacist any health conditions that may have an adverse effect on drug products or may pose a health or safety risk to others.

8.5(11) *Hazardous drugs.* The pharmacy shall ensure pharmacy personnel and patients are adequately protected from unnecessary exposure to hazardous drugs. As of December 1, 2019, the pharmacy shall be in compliance with United States Pharmacopeia (USP) General Chapter 800 for handling hazardous drugs. A pharmacy engaged in compounding of hazardous drugs may request delayed compliance for specific requirements in USP General Chapter 800 pertaining to compounding, in accordance with rule 657—20.5(126,155A).

[ARC 8671B, IAB 4/7/10, effective 5/12/10; ARC 0503C, IAB 12/12/12, effective 1/16/13; ARC 1961C, IAB 4/15/15, effective 5/20/15; ARC 2408C, IAB 2/17/16, effective 3/23/16; ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 4267C, IAB 1/30/19, effective 3/6/19; ARC 4454C, IAB 5/22/19, effective 6/26/19; ARC 5348C, IAB 12/30/20, effective 2/3/21]

657—8.6(155A) Health of personnel. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.7(155A) Procurement, storage, and recall of drugs and devices.

8.7(1) *Source.* Procurement of prescription drugs and devices shall be from an Iowa-licensed distributor or, on a limited basis, from another licensed pharmacy or licensed practitioner located in the United States.

8.7(2) *Manner of storage.* Drugs and devices shall be stored in a manner to protect their identity and integrity.

8.7(3) *Storage temperatures.* All drugs and devices shall be stored at the proper temperature as provided in manufacturer labeling. In the absence of a specific temperature range, the pharmacy shall defer to storage conditions identified in United States Pharmacopeia chapter 659.

8.7(4) *Product recall.* There shall be a system for removing from use, including unit dose, any drugs and devices subjected to a product recall.

8.7(5) *Outdated drugs or devices.* Any drug or device bearing an expiration date shall not be dispensed for use beyond the expiration date of the drug or device. Outdated drugs or devices shall be removed from dispensing stock and shall be quarantined until such drugs or devices are properly disposed of.

8.7(6) Records. All pharmacies shall maintain supplier invoices of prescription drugs and controlled substances upon which the actual date of receipt of the drugs by the pharmacist or other responsible individual is clearly recorded. All pharmacies shall maintain supplier credit memos. Pharmacy records of invoices and credit memos shall be maintained for at least two years from the date of the record. If the original supplier invoice or credit memo is received electronically, hard-copy record is not required. [ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.8(124,155A) Out-of-date drugs or devices. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.9(124,155A) Records storage. Every record required to be maintained by a pharmacy pursuant to board rules or Iowa Code chapters 124 and 155A shall be maintained and be available for inspection and copying by the board or its representative for at least two years from the date of such record or the date of last activity on the record unless a longer retention period is specified for the particular record.

8.9(1) Records less than 12 months old. Records shall be maintained within the licensed pharmacy department for a minimum of 12 months, except as provided herein. Pharmacy records less than 12 months old may be stored in a secure storage area outside the licensed pharmacy department, including at a remote location, if the pharmacy has retained electronic copies of the records in the pharmacy that are immediately available and if the original records are available within 72 hours of a request by the board or its authorized agent, unless such remote storage is prohibited under federal law.

8.9(2) Records more than 12 months old. Records more than 12 months old may be maintained in a secure storage area outside the licensed pharmacy department, including at a remote location, if the records are retrievable within 72 hours of a request by the board or its authorized agent, unless such remote storage is prohibited under federal law.

[ARC 8539B, IAB 2/24/10, effective 4/1/10; ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 5007C, IAB 3/25/20, effective 4/29/20]

657—8.10 Reserved.

657—8.11(147,155A) Unethical conduct or practice. The provisions of this rule apply to licensed pharmacies, licensed pharmacists, registered pharmacy technicians, registered pharmacy support persons, and registered pharmacist-interns.

8.11(1) Misrepresentative deeds. A pharmacy, pharmacist, technician, support person, or pharmacist-intern shall not make any statement intended to deceive, misrepresent or mislead anyone, or be a party to or an accessory to any fraudulent or deceitful practice or transaction in pharmacy or in the operation or conduct of a pharmacy.

8.11(2) Unethical conduct.

a. A pharmacy, pharmacist, pharmacist-intern, technician, or support person shall not participate in any of the following types of unethical conduct:

(1) Any activity that negates a patient's freedom of choice of pharmacy services.

(2) Providing prescription blanks or forms bearing the pharmacy's name or other means of identification to any person authorized to prescribe, except that a hospital may make prescription blanks or forms bearing the hospital pharmacy's name or other means of identification available to hospital staff prescribers, emergency department prescribers, and prescribers granted hospital privileges for the prescribers' use during practice at or in the hospital.

(3) Any financial arrangement or transaction that would violate federal healthcare fraud, waste, and abuse laws, including but not limited to the Stark Law, the False Claims Act, and the Anti-Kickback Statute.

b. A purchasing pharmacist or pharmacy shall not engage in any activity or include in any agreement with a selling pharmacist or pharmacy any provision that would prevent or prohibit the prior notifications required in subrule 8.35(7).

8.11(3) Discrimination. A pharmacy, pharmacist, pharmacist-intern, technician, or pharmacy support person shall not discriminate between patients or groups of patients for reasons of religion, race,

creed, color, gender, gender identity, sexual orientation, marital status, age, national origin, physical or mental disability, or disease state when providing pharmaceutical services.

8.11(4) *Unprofessional conduct or behavior.* A pharmacy, pharmacist, pharmacist-intern, technician, or pharmacy support person shall not engage in unprofessional behavior in connection with the practice of pharmacy. Unprofessional behavior shall include, but not be limited to, the following acts: verbal abuse, coercion, intimidation, harassment, sexual advances, threats, degradation of character, indecent or obscene conduct, theft, and the refusal to provide reasonable information or answer reasonable questions for the benefit of the patient.

[ARC 9526B, IAB 6/1/11, effective 7/6/11; ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.12(126,147) Advertising. Prescription drug information, including price, may be provided to the public by a pharmacy so long as the information is not false or misleading and is not in violation of any federal or state laws applicable to the advertisement of such articles generally and if all of the following conditions are met:

1. All charges for services to the consumer shall be stated.
2. The effective dates for the prices listed shall be stated.
3. No reference shall be made to controlled substances listed in Schedules II through V of the

latest revision of the Iowa uniform controlled substances Act and the rules of the board.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.13(135C,155A) Personnel histories. Pursuant to the requirements of Iowa Code section 135C.33, the provisions of this rule shall apply to any pharmacy employing any person to provide patient care services in a patient's home. For the purposes of this rule, "employed by the pharmacy" shall include any individual who is paid to provide treatment or services to any patient in the patient's home, whether the individual is paid by the pharmacy or by any other entity such as a corporation, a temporary staffing agency, or an independent contractor. Specifically excluded from the requirements of this rule are individuals such as delivery persons or couriers who do not enter the patient's home for the purpose of instructing the patient or the patient's caregiver in the use or maintenance of the equipment, device, or drug being delivered, or who do not enter the patient's home for the purpose of setting up or servicing the equipment, device, or drug used to treat the patient in the patient's home.

8.13(1) *Applicant acknowledgment.* The pharmacy shall ask the following question of each person seeking employment in a position that will provide in-home services: "Do you have a record of founded child or dependent adult abuse or have you ever been convicted of a crime, in this state or any other state?" The applicant shall also be informed that a criminal history and child and dependent adult abuse record checks will be conducted. The applicant shall indicate, by signed acknowledgment, that the applicant has been informed that such record checks will be conducted.

8.13(2) *Criminal history check.* Prior to the employment of any person to provide in-home services as described by this rule, the pharmacy shall request that the department of public safety perform a criminal history check.

8.13(3) *Abuse history checks.* Prior to the employment of any person to provide in-home services as described by this rule, the pharmacy shall request that the department of human services perform a child and dependent adult abuse record check.

a. A person who has a criminal record, founded dependent adult abuse report, or founded child abuse report shall not be employed by a pharmacy to provide in-home services unless the department of human services has evaluated the crime or founded abuse report, has concluded that the crime or founded abuse does not merit prohibition from such employment, and has notified the pharmacy that the person may be employed to provide in-home services.

b. The pharmacy shall keep copies of all record checks and evaluations for a minimum of two years following receipt of the record or for a minimum of two years after the individual is no longer employed by the pharmacy, whichever is greater.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.14(155A) Training and utilization of registered pharmacy staff. Pursuant to rule 657—8.3(155A), all Iowa-licensed pharmacies utilizing pharmacist-interns, pharmacy technicians, or pharmacy support persons shall have written policies and procedures for the training and utilization of pharmacist-interns, pharmacy technicians, and pharmacy support persons appropriate to the practice of pharmacy at that licensed location. Training shall be documented and maintained by the pharmacy for at least two years from the last date of employment or internship and shall be available for inspection by the board or its authorized agent.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 1961C, IAB 4/15/15, effective 5/20/15; ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.15(155A) Delivery of prescription drugs and devices. A prescription order may be delivered to a patient at any location licensed as a pharmacy. Alternatively, a pharmacy may use the mail, a common carrier, or personal delivery to deliver a prescription order to any location requested by the patient. A pharmacy that delivers prescription orders by one or more alternate methods shall have policies and procedures to ensure patient confidentiality, prescription order accountability, and proper storage of prescription orders during delivery. When counseling is required pursuant to rule 657—6.14(155A), oral counseling shall be provided before the prescription order is delivered to the patient. Documentation of the delivery of prescription orders shall be maintained by the pharmacy for at least two years from the date of delivery. The term “patient” includes the patient and the patient’s authorized representatives.

[ARC 5007C, IAB 3/25/20, effective 4/29/20]

657—8.16(124,155A) Confidential information.

8.16(1) Release of confidential information. Confidential information may be released only as follows:

- a. Pursuant to the express written authorization of the patient or the order or direction of a court.
- b. To the patient or the patient’s authorized representative.
- c. To the prescriber or other licensed practitioner then caring for the patient.
- d. To another licensed pharmacist when the best interests of the patient require such release.
- e. To the board or its representative or to such other persons or governmental agencies duly authorized by law to receive such information.

A pharmacist shall utilize the resources available to determine, in the professional judgment of the pharmacist, that any persons requesting confidential patient information pursuant to this rule are entitled to receive that information.

8.16(2) Exceptions. Nothing in this rule shall prohibit a pharmacist from releasing confidential patient information as follows:

- a. Transferring a prescription to another pharmacy upon the request of the patient or the patient’s authorized representative or pursuant to subrule 8.35(7) when the pharmacy is discontinuing operations.
- b. Providing the patient with a copy of a nonrefillable prescription that is clearly marked as a copy and not to be filled.
- c. Providing drug therapy information to authorized practitioners for their patients.
- d. Disclosing information necessary for the processing of third-party payer claims on behalf of the patient.

8.16(3) Record disposal. Disposal of any materials containing or including patient-specific or confidential information shall be conducted in a manner to preserve patient confidentiality.

[ARC 9526B, IAB 6/1/11, effective 7/6/11; ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.17 Reserved.

657—8.18(124,155A) Electronic prescription mandate. Beginning January 1, 2020, all prescriptions shall be transmitted electronically to a pharmacy pursuant to rule 657—21.6(124,155A), except as provided in rule 657—21.8(124,155A). A pharmacist who receives a written, oral, or facsimile prescription shall not be required to verify that the prescription is subject to an exception provided in

rule 657—21.8(124,155A) and may dispense a prescription drug pursuant to an otherwise valid written, oral, or facsimile prescription pursuant to rule 657—8.19(124,126,155A).
[ARC 4580C, IAB 7/31/19, effective 9/4/19]

657—8.19(124,126,135,155A,280) Manner of issuance of a prescription drug or medication order. A prescription drug order or medication order that is issued prior to January 1, 2020, or that is exempt from the electronic prescription mandate pursuant to rule 657—21.8(124,155A) may be transmitted from a prescriber or a prescriber's agent to a pharmacy in written form, orally including telephone voice communication, by facsimile transmission as provided in rule 657—21.7(124,155A), or by electronic transmission in accordance with applicable federal and state laws, rules, and regulations. Any prescription drug order or medication order provided to a patient in written or printed form shall include the original, handwritten signature of the prescriber except as provided in rule 657—21.6(124,155A).

8.19(1) Requirements for a prescription. A valid prescription drug order shall be based on a valid patient-prescriber relationship except as provided in subrule 8.19(7) for epinephrine auto-injectors, subrule 8.19(8) for opioid antagonists, or subrule 8.19(10) for bronchodilator canisters or bronchodilator canisters and spacers.

a. Written, electronic, or facsimile prescription. In addition to the electronic prescription application and pharmacy prescription application requirements of this rule, a written, electronic, or facsimile prescription shall include:

- (1) The date issued.
- (2) The name and address of the patient except as provided in subrule 8.19(7) for epinephrine auto-injectors, subrule 8.19(8) for opioid antagonists, subrule 8.19(9) for expedited partner therapy, or subrule 8.19(10) for bronchodilator canisters or bronchodilator canisters and spacers.
- (3) The name, strength, and quantity of the drug or device prescribed.
- (4) The name and address of the prescriber and, if the prescription is for a controlled substance, the prescriber's DEA registration number.
- (5) The written or electronic signature of the prescriber.

b. Written prescription. In addition to the requirements of paragraph 8.19(1)“a,” a written prescription shall be manually signed, with ink or indelible pencil, by the prescriber. The requirement for manual signature shall not apply when an electronically prepared and signed prescription for a noncontrolled substance is printed on security paper as provided in 657—paragraph 21.6(2)“b.”

c. Facsimile prescription. In addition to the requirements of paragraph 8.19(1)“a,” a prescription transmitted via facsimile shall include:

- (1) The identification number of the facsimile machine used to transmit the prescription to the pharmacy.
- (2) The time and date of transmission of the prescription.
- (3) The name, address, telephone number, and facsimile number of the pharmacy to which the prescription is being transmitted.
- (4) If the prescription is for a controlled substance and in compliance with DEA regulations, the manual signature of the prescriber.

d. Electronic prescription. In addition to the requirements of paragraph 8.19(1)“a,” an electronically prepared prescription for a controlled or noncontrolled prescription drug or device that is electronically transmitted to a pharmacy shall include the prescriber's electronic signature, except as provided herein.

- (1) An electronically prepared prescription for a controlled substance that is printed out or faxed by the prescriber or the prescriber's agent shall be manually signed by the prescriber.
- (2) The prescriber shall ensure that the electronic prescription application used to prepare and transmit the electronic prescription complies with applicable state and federal laws, rules, and regulations regarding electronic prescriptions.
- (3) The prescriber or the prescriber's agent shall provide verbal verification of an electronic prescription upon the request of the pharmacy.

(4) An electronic prescription for a noncontrolled prescription drug or device that is transmitted by an authorized agent shall not be required to contain the prescriber's electronic signature.

8.19(2) Verification. The pharmacist shall exercise professional judgment regarding the accuracy, validity, and authenticity of any prescription drug order or medication order consistent with federal and state laws, rules, and regulations. In exercising professional judgment, the prescriber and the pharmacist shall take adequate measures to guard against the diversion of prescription drugs and controlled substances through prescription forgeries.

8.19(3) Transmitting agent. The prescriber may authorize an agent to transmit to the pharmacy a prescription drug order or medication order orally, by facsimile transmission, or by electronic transmission provided that the first and last names and title of the transmitting agent are included in the order.

a. New order. A new written or electronically prepared and transmitted prescription drug or medication order shall be manually or electronically signed by the prescriber, except as provided in paragraph 8.19(1) "d." If transmitted by the prescriber's agent, the first and last names and title of the transmitting agent shall be included in the order. If the prescription is for a controlled substance and is written or printed from an electronic prescription application, the prescription shall be manually signed by the prescriber. An electronically prepared prescription shall not be electronically transmitted to the pharmacy if the prescription has been printed prior to the electronic transmission. An electronically prepared and electronically transmitted prescription that is printed following the electronic transmission shall be clearly labeled as a copy, not valid for dispensing.

b. Refill order or renewal order. An authorization to refill a prescription drug or medication order, or to renew or continue an existing drug therapy, may be transmitted to professional pharmacy staff through oral communication, in writing, by facsimile transmission, or by electronic transmission initiated by or directed by the prescriber.

(1) If the transmission is completed by the prescriber's agent and the first and last names and title of the transmitting agent are included in the order, the prescriber's signature is not required on the fax or alternate electronic transmission.

(2) If the order differs in any manner from the original order, such as a change of the drug strength, dosage form, or directions for use, the prescriber shall sign the order as provided by paragraph 8.19(3) "a."

8.19(4) Receiving agent. Regardless of the means of transmission to a pharmacy, only professional pharmacy staff shall be authorized to receive a new prescription drug or medication order from a prescriber or the prescriber's agent. A technician trainee may receive a refill or renewal order from a prescriber or the prescriber's agent only if the technician's supervising pharmacist has authorized that function.

8.19(5) Legitimate purpose. The pharmacy and professional pharmacy staff shall ensure that the prescription drug or medication order, regardless of the means of transmission, has been issued for a legitimate medical purpose by a prescriber acting in the usual course of the prescriber's professional practice. A pharmacist shall not dispense a prescription drug if the pharmacist knows or should have known that the prescription was issued solely on the basis of an Internet-based questionnaire.

8.19(6) Refills. A refill is one or more dispensings of a prescription drug or device that result in the patient's receipt of the quantity authorized by the prescriber for a single fill as indicated on the prescription drug order.

a. Noncontrolled prescription drug or device. A prescription for a prescription drug or device that is not a controlled substance may authorize no more than 12 refills within 18 months following the date on which the prescription is issued.

b. Controlled substance. A prescription for a Schedule III, IV, or V controlled substance may authorize no more than 5 refills within 6 months following the date on which the prescription is issued.

8.19(7) Epinephrine auto-injector prescription issued to school or facility. A physician, an advanced registered nurse practitioner, or a physician assistant may issue a prescription for one or more epinephrine auto-injectors in the name of a facility as defined in Iowa Code subsection 135.185(1), a school district, or an accredited nonpublic school. The prescription shall comply with all requirements of subrule 8.19(1)

as applicable to the form of the prescription except that the prescription shall be issued in the name and address of the facility, the school district, or the accredited nonpublic school in lieu of the name and address of a patient. Provisions requiring a preexisting patient-prescriber relationship shall not apply to a prescription issued pursuant to this subrule.

a. The pharmacy's patient profile and record of dispensing of a prescription issued pursuant to this subrule shall be maintained in the name of the facility, school district, or accredited nonpublic school to which the prescription was issued and the drug was dispensed.

b. The label affixed to an epinephrine auto-injector dispensed pursuant to this subrule shall identify the name of the facility, school district, or accredited nonpublic school to which the prescription is dispensed.

8.19(8) *Opioid antagonist prescription issued to law enforcement, fire department, service program, or school district.* A physician, an advanced registered nurse practitioner, or a physician assistant may issue a prescription for one or more opioid antagonists in the name of a law enforcement agency, fire department, or service program pursuant to Iowa Code section 147A.18 and rule 657—39.7(135,147A), or in the name of a school district pursuant to Iowa Code section 135.190 and rule 657—39.7(135,147A). The prescription shall comply with all requirements of subrule 8.19(1) as applicable to the form of the prescription except that the prescription shall be issued in the name and address of the law enforcement agency, fire department, service program, or school district in lieu of the name and address of a patient. Provisions requiring a preexisting patient-prescriber relationship shall not apply to a prescription issued pursuant to this subrule.

a. The pharmacy's patient profile and record of dispensing of an opioid antagonist pursuant to this subrule shall be maintained in the name of the law enforcement agency, fire department, service program, or school district to which the prescription was issued and the drug was dispensed.

b. The label affixed to an opioid antagonist dispensed pursuant to this subrule shall identify the name of the law enforcement agency, fire department, service program, or school district to which the prescription is dispensed and shall be affixed such that the expiration date of the drug is not rendered illegible.

8.19(9) *Expedited partner therapy.* Pursuant to Iowa Code section 139A.41, a physician, physician assistant, or advanced registered nurse practitioner may issue a prescription to one or more sexual partners of an infected patient for an oral antibiotic intended to treat a sexually transmitted chlamydia or gonorrhea infection. The prescription shall comply with all requirements of subrule 8.19(1) as applicable to the form of the prescription except that the prescription shall not be required to contain the patient name and address. The prescription shall indicate the antibiotic is being issued for the purpose of expedited partner therapy. Provisions requiring a preexisting patient-prescriber relationship shall not apply to a prescription issued pursuant to this subrule.

8.19(10) *Bronchodilator canister or bronchodilator canister and spacer prescription issued to a school district or accredited nonpublic school.* A physician, an advanced registered nurse practitioner, or a physician assistant may issue a prescription for one or more bronchodilator canisters or bronchodilator canisters and spacers in the name of a school district or accredited nonpublic school pursuant to Iowa Code section 280.16A. The prescription shall comply with all requirements of subrule 8.19(1) as applicable to the form of the prescription except that the prescription shall be issued in the name and address of the school district or accredited nonpublic school in lieu of the name and address of a patient. Provisions requiring a preexisting patient-prescriber relationship shall not apply to a prescription issued pursuant to this subrule.

a. The pharmacy's patient profile and record of dispensing of a bronchodilator canister or bronchodilator canister and spacer pursuant to this subrule shall be maintained in the name of the school district or accredited nonpublic school to which the prescription was issued and the drug was dispensed.

b. The label affixed to a bronchodilator canister or bronchodilator canister and spacer dispensed pursuant to this subrule shall identify the name of the school district or accredited nonpublic school to

which the prescription is dispensed and shall be affixed such that the expiration date of the drug is not rendered illegible.

[ARC 8171B, IAB 9/23/09, effective 10/28/09; ARC 9912B, IAB 12/14/11, effective 1/18/12; ARC 2414C, IAB 2/17/16, effective 3/23/16; ARC 2827C, IAB 11/23/16, effective 11/3/16; ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 4580C, IAB 7/31/19, effective 9/4/19; ARC 4903C, IAB 2/12/20, effective 3/18/20; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—8.20(155A) Valid prescriber/patient relationship. Prescription drug orders and medication orders shall be valid as long as a prescriber/patient relationship exists. Once the prescriber/patient relationship is broken and the prescriber is no longer available to treat the patient or oversee the patient's use of a prescription drug, any remaining prescription refills may be dispensed at the discretion of the pharmacist for a suitable amount of time so that the patient can establish care with a new provider and a new order can be issued. In determining the duration of which prescriptions may be dispensed, the pharmacist shall consider the patient's health care status and access to health care services.

[ARC 3639C, IAB 2/14/18, effective 3/21/18]

657—8.21(155A) Prospective drug use review.

8.21(1) For purposes of promoting therapeutic appropriateness and ensuring rational drug therapy, a pharmacist shall review the patient record, information obtained from the patient, and each prescription drug or medication order to identify:

- a. Overutilization or underutilization;
- b. Therapeutic duplication;
- c. Drug-disease contraindications;
- d. Drug-drug interactions;
- e. Incorrect drug dosage or duration of drug treatment;
- f. Drug-allergy interactions;
- g. Clinical abuse/misuse;
- h. Drug-prescriber contraindications.

Upon recognizing any of the above, the pharmacist shall take appropriate steps to avoid or resolve the problem and shall, if necessary, include consultation with the prescriber. Information that shall be obtained for the purpose of drug utilization review includes, but is not limited to, a complete list of prescription and nonprescription medications being used by the patient, patient allergies, and patient disease states. The collection of patient information to be used for drug utilization review may be delegated to a pharmacy technician or pharmacist-intern. The review and assessment of patient records shall not be delegated to pharmacy technicians or pharmacy support persons but may be delegated to registered pharmacist-interns under the direct supervision of the pharmacist.

8.21(2) A pharmacist shall be exempt from the requirements of subrule 8.21(1) when dispensing a prescription issued to an unnamed patient for an oral antibiotic pursuant to Iowa Code section 139A.41. [ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 4903C, IAB 2/12/20, effective 3/18/20; ARC 5350C, IAB 12/30/20, effective 2/3/21]

657—8.22(155A) Notification of interchangeable biological product selection. Pursuant to Iowa Code section 155A.32, when a pharmacist substitutes a biological product that is an interchangeable biological product for the biological product prescribed, the pharmacist or pharmacist's designee shall, within five business days of dispensing the biological product, communicate to the prescriber the name and manufacturer of the biological product dispensed unless the prescription information has been entered into an electronic record system, such as an electronic medical record, electronic prescribing system, pharmacy benefit management system, or a pharmacy record to which the prescriber has access. The manner of communication to the prescriber may be via telephone, facsimile, electronic transmission, or other prevailing means.

This rule is intended to implement Iowa Code section 155A.32. [ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.23(124,155A) Individuals qualified to administer. Rescinded ARC 6076C, IAB 12/15/21, effective 1/19/22.

657—8.24(155A) Documented verification. The pharmacist shall provide, document, and retain a record of the final verification for the accuracy, validity, completeness, and appropriateness of the patient's prescription or medication order prior to the delivery of the medication to the patient or the patient's representative. In an approved technician product verification program, the checking technician shall provide, document, and retain a record of the final verification for the accuracy of the patient's prescription or medication order prior to the delivery of the medication to the patient or the patient's representative.

[ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 5007C, IAB 3/25/20, effective 4/29/20]

657—8.25 Reserved.

657—8.26(155A) Continuous quality improvement program. Pursuant to rule 657—8.3(155A), each pharmacy licensed to provide pharmaceutical services to patients in Iowa shall implement or participate in a continuous quality improvement program (CQI program). The CQI program is intended to be an ongoing, systematic program of standards and procedures to detect, identify, evaluate, and prevent medication errors, thereby improving medication therapy and the quality of patient care. A pharmacy that participates as an active member of a hospital or corporate CQI program that meets the objectives of this rule shall not be required to implement a new program pursuant to this rule.

8.26(1) Reportable program events. For purposes of this rule, a reportable program event or program event means a preventable medication error resulting in the incorrect dispensing of a prescribed drug received by or administered to the patient and includes but is not necessarily limited to:

- a. An incorrect drug;
- b. An incorrect drug strength;
- c. An incorrect dosage form;
- d. A drug received by the wrong patient;
- e. Inadequate or incorrect packaging, labeling, or directions; or
- f. Any incident related to a prescription dispensed to a patient that results in or has the potential to result in serious harm to the patient.

8.26(2) Responsibility. The pharmacist in charge may delegate program administration and monitoring, but the pharmacist in charge maintains ultimate responsibility for the validity and consistency of program activities.

8.26(3) Policies and procedures. Pursuant to rule 657—8.3(155A), each pharmacy shall have written policies and procedures for the operation and management of the pharmacy's CQI program. A copy of the pharmacy's CQI program description and policies and procedures shall be maintained and readily available to all pharmacy personnel. The policies and procedures shall address, at a minimum, a planned process to:

- a. Train all pharmacy personnel in relevant phases of the CQI program;
- b. Identify and document reportable program events;
- c. Minimize the impact of reportable program events on patients;
- d. Analyze data collected to assess the causes and any contributing factors relating to reportable program events;
- e. Use the findings to formulate an appropriate response and to develop pharmacy systems and workflow processes designed to prevent and reduce reportable program events; and
- f. Periodically, but at least quarterly, meet with appropriate pharmacy personnel to review findings and inform personnel of changes that have been made to pharmacy policies, procedures, systems, or processes as a result of CQI program findings.

8.26(4) Event discovery and notification. As provided by the procedures of the CQI program, the pharmacist in charge or appropriate designee shall be informed of and review all reported and documented program events. All pharmacy personnel shall be trained to immediately inform the pharmacist on duty of any discovered or suspected program event. When the pharmacist on duty determines that a reportable program event has occurred, the pharmacist shall ensure that all reasonably

necessary steps are taken to remedy any problems or potential problems for the patient and that those steps are documented. Necessary steps include, but are not limited to, the following:

- a. Notifying the patient or the patient's caregiver and the prescriber or other members of the patient's health care team as warranted;
- b. Identifying and communicating directions or processes for correcting the error; and
- c. Communicating instructions for minimizing any negative impact on the patient.

8.26(5) CQI program records. All CQI program records shall be maintained on site at the pharmacy or shall be accessible at the pharmacy and be available for inspection and copying by the board or its representative for at least two years from the date of the record. When a reportable program event occurs or is suspected to have occurred, the program event shall be documented in a written or electronic storage record created solely for that purpose. Records of program events shall be maintained in an orderly manner and shall be filed chronologically by date of discovery.

a. The program event shall initially be documented as soon as practicable but no more than three days following discovery of the event by the staff member who discovers the event or is informed of the event.

b. Program event documentation shall include a description of the event that provides sufficient information to permit categorization and analysis of the event and shall include:

- (1) The date and time the program event was discovered and the name of the staff person who discovered the event; and
- (2) The names of the individuals recording and reviewing or analyzing the program event information.

8.26(6) Program event analysis and response. The pharmacist in charge or designee shall review each reportable program event and determine if follow-up is necessary. When appropriate, information and data collected and documented shall be analyzed, individually and collectively, to assess the cause and any factors contributing to the program event. The analysis may include, but is not limited to, the following:

- a. A consideration of the effects on the quality of the pharmacy system related to workflow processes, technology utilization and support, personnel training, and both professional and technical staffing levels;
- b. Any recommendations for remedial changes to pharmacy policies, procedures, systems, or processes; and
- c. The development of a set of indicators that a pharmacy will utilize to measure its program standards over a designated period of time.

[ARC 1961C, IAB 4/15/15, effective 5/20/15; ARC 2413C, IAB 2/17/16, effective 3/23/16; ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—8.27 to 8.29 Reserved.

657—8.30(126,155A) Sterile products. Rescinded IAB 6/6/07, effective 7/11/07.

657—8.31(135,147A) Opioid antagonist dispensing by pharmacists by standing order. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.32(124,155A) Individuals qualified to administer. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.33(155A) Vaccine administration by pharmacists. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.34(155A) Collaborative drug therapy management. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

657—8.35(155A) Pharmacy license. A pharmacy license issued by the board is required for all sites where prescription drugs are offered for sale or dispensed under the supervision of a pharmacist. The current pharmacy license certificate shall be displayed in a position visible to the public. The board may issue any of the following types of pharmacy licenses: a general pharmacy license, a hospital pharmacy license, a limited use pharmacy license, or a nonresident pharmacy license. Nonresident pharmacy license applicants shall comply with board rules regarding nonresident pharmacy practice except when a waiver has been granted. Applicants for general or hospital pharmacy practice shall comply with board rules regarding general or hospital pharmacy practice except when a waiver has been granted. Any pharmacy that dispenses controlled substances to Iowa residents must also register pursuant to 657—Chapter 10.

8.35(1) Limited use pharmacy license. A limited use pharmacy license may be issued for nuclear pharmacy practice, correctional facility pharmacy practice, veterinary pharmacy practice, telepharmacy practice, and other limited use practice settings. Applications for a limited use pharmacy license shall be considered on a case-by-case basis.

8.35(2) Application. Applicants for initial licensure, license renewal, license reactivation, or license changes pursuant to subrule 8.35(6) shall complete the relevant pharmacy license application and shall include all required information and attachments. All pharmacy license applications require submission of a nonrefundable \$135 license fee plus applicable penalty fees. The application shall include the signature of the pharmacy owner's authorized representative and shall require at a minimum the following:

- a. Disclosure of pharmacy ownership information, including information about the pharmacy's registered agent;
- b. Identification and signature of the pharmacist in charge;
- c. The identification of and average number of hours worked by all pharmacists, pharmacist-interns, pharmacy technicians, and pharmacy support persons working in the pharmacy;
- d. Criminal and disciplinary history information;
- e. Description of the scope of services provided by the pharmacy; and
- f. If the pharmacy is located outside of Iowa, identification of a registered agent located in Iowa.

8.35(3) License renewal. A pharmacy license shall be renewed before January 1 of each year. An initial pharmacy license issued between November 1 and December 31 shall not require renewal until the following calendar year. The nonrefundable fee for a timely license renewal shall be \$135.

a. *Delinquent license grace period.* A pharmacy license renewal application that is postmarked or hand-delivered to the board after January 1 but prior to February 1 following expiration shall be considered delinquent and shall require the nonrefundable payment of the renewal fee plus a penalty fee of \$135. A pharmacy that submits a completed license renewal application, application fee, and penalty fee postmarked or delivered to the board office by January 31 shall not be subject to disciplinary action for continuing to operate in the month of January.

b. *Delinquent license reactivation beyond grace period.* If a pharmacy license is not renewed prior to the expiration of the one-month grace period identified in paragraph 8.35(3) "a," the pharmacy may not operate or provide pharmacy services to patients in the state of Iowa until the license is reactivated. A pharmacy without a current license may apply for license reactivation by submitting an application for reactivation and a nonrefundable \$540 reactivation fee. As part of the reactivation application, the pharmacy shall disclose the prescriptions dispensed and the services, if any, that were provided to Iowa patients while the license was delinquent. A pharmacy that continues to operate or provide pharmacy services in Iowa without a current license may be subject to disciplinary sanctions.

8.35(4) Inspection of pharmacy location.

a. A new pharmacy location in Iowa shall require an on-site inspection by an authorized agent of the board. Application for a pharmacy license and other required registrations shall be submitted to the board at least 14 days prior to the anticipated inspection. Any deficiencies identified during the inspection shall be corrected and verified by an authorized agent of the board prior to the issuance of the pharmacy license. Prescription drugs, including controlled substances, may not be delivered to a new pharmacy location prior to the delivery of the pharmacy license and registration certificates.

b. A pharmacy location in Iowa which is applying for a different license type than previously held may be subject to an inspection prior to the issuance of the new license.

8.35(5) Failure to complete licensure. An application for a pharmacy license, including any other required registration applications, will become null and void if the applicant fails to complete the licensure process within six months of acceptance by the board of the required applications. The licensure process shall be complete upon the pharmacy's opening for business at the licensed location following a satisfactory inspection by an agent of the board pursuant to this rule. When an applicant fails to timely complete the licensure process, fees submitted with applications will not be transferred or refunded. If the applicant intends to proceed with a pharmacy license, a new application and fee shall be required.

8.35(6) Pharmacy license changes. When a pharmacy changes its name, location, ownership, pharmacist in charge, or license type, a completed pharmacy license application with a nonrefundable \$135 fee shall be submitted to the board pursuant to subrule 8.35(2). Upon receipt of the completed application and fee, the board shall issue an updated pharmacy license certificate, pending any necessary inspection pursuant to paragraph 8.35(4) "b," unless the board identifies any ground for denial of the license. Any restrictions or disciplinary history associated with the previous pharmacy shall remain unchanged. A pharmacy wishing to disassociate itself from the previously licensed pharmacy restrictions or disciplinary history may petition the board for such disassociation. The burden is on the pharmacy to demonstrate that the current pharmacy is not associated with or responsible for the pharmacy as it previously existed. The old license certificate shall be returned to the board within ten days of receiving the updated license certificate.

a. Name. A change of the name under which the pharmacy is doing business shall require submission of a pharmacy license application and appropriate fee prior to the change of name.

b. Location. Except as provided in subrule 8.35(10) for a temporary relocation due to an exceptional circumstance, a change of pharmacy location shall require submission of a pharmacy license application and appropriate fee prior to the change of location. A pharmacy undergoing a change in location is required to notify patients of the change in accordance with paragraph 8.35(7) "d." A change of pharmacy location in Iowa may require an on-site inspection of the new location as provided in subrule 8.35(4).

c. Ownership. A change in ownership of a pharmacy shall require submission of a pharmacy license application and appropriate fee prior to the change in ownership. A change of ownership occurs when the owner listed on the pharmacy's most recent application changes. A pharmacy undergoing a change in ownership is required to notify the pharmacist in charge and patients of the change in accordance with subrule 8.35(7). A change of ownership effectively consists of closing a pharmacy and opening a new pharmacy.

d. Pharmacist in charge. In addition to the requirements of this paragraph, a change of pharmacist in charge for a nonresident pharmacy shall require registration of the new permanent pharmacist in charge if the pharmacist in charge is not currently registered by the board or licensed to practice pharmacy in Iowa.

(1) If a permanent pharmacist in charge has been identified by the time of the vacancy, a pharmacy license application identifying the new pharmacist in charge, along with the appropriate fee, shall be submitted to the board within ten days of the change.

(2) If a permanent pharmacist in charge has not been identified by the time of the vacancy, a temporary pharmacist in charge shall be identified. Written notification identifying the temporary pharmacist in charge shall be submitted to the board within ten days of the vacancy.

(3) If a permanent pharmacist in charge was not identified within ten days of the vacancy, the pharmacy shall, within 90 days of the vacancy, identify a permanent pharmacist in charge. A pharmacy license application identifying the permanent pharmacist in charge, along with appropriate fee, shall be submitted to the board within ten days of the appointment of a permanent pharmacist in charge. The pharmacy license application and the pharmacist in charge registration application, if needed, including appropriate fees, shall be received by the board within 90 days of the original vacancy of the permanent pharmacist in charge position.

(4) If a permanent pharmacist in charge is out of the pharmacy for an extended leave of absence of no more than 120 days, the pharmacy may identify an interim pharmacist in charge and provide notice of such to the board on forms provided by the board. Identification of an interim pharmacist in charge shall not require submission of a new pharmacy license application and shall not result in a permanent change in pharmacist in charge on the pharmacy license. If a permanent pharmacist in charge is out of the pharmacy for an extended leave of absence greater than 120 days, the pharmacy shall initiate a change of pharmacist in charge in accordance with this rule.

e. License type. A change in pharmacy license type shall require submission of a pharmacy license application and appropriate fee prior to the change in license type. A pharmacy changing license type shall notify the pharmacist in charge and patients of the change in accordance with subrule 8.35(7).

f. License change application submission. An application for license change shall be timely submitted pursuant to this subrule. A licensed pharmacy that has timely submitted an application for license change and fee may continue to service Iowa patients while the license change is pending final approval. An applicant who has submitted an application for license change after the required date of submission pursuant to this subrule but within 30 days of the required date of submission shall be assessed a nonrefundable late penalty fee of \$135 in addition to the license fee. An applicant who has submitted an application for license change 31 days or later following the required date of submission pursuant to this subrule shall be assessed a nonrefundable late penalty fee of \$540.

8.35(7) Closing or sale of a pharmacy. A closing pharmacy shall ensure that all pharmacy records are transferred to another licensed pharmacy that agrees to act as custodian of the records for at least two years. A pharmacy shall not execute a sale or closing of a pharmacy unless there exists an adequate period of time prior to the pharmacy's closing for delivery of the notifications to the pharmacist in charge, the board, the DEA, and pharmacy patients as required by this subrule. The executive director may exempt a pharmacy from one or more of the notification requirements in the event of an unforeseeable closure.

a. Pharmacist in charge notification. At least 40 days prior to the effective date of the sale or closing of a pharmacy, the pharmacist in charge of the closing pharmacy shall be notified of the proposed sale or closing. Information regarding the pending sale or closure of the pharmacy may be kept confidential until public notifications, which are required 30 days prior to the pharmacy's closing, are made. The pharmacist in charge of the closing pharmacy shall provide input and direction to the pharmacy owner regarding the responsibilities of the closing pharmacy, including the notifications, deadlines, and timelines established by this subrule. The pharmacist in charge of the purchasing or receiving pharmacy shall be notified of the pending transaction at least 30 days prior to the sale or closure of the pharmacy.

b. Board and DEA notifications. At least 30 days prior to the closing of a pharmacy, a written notice shall be sent to the board. Notification to the DEA shall be pursuant to federal regulation. Notification to the board shall include:

- (1) The anticipated date of closing or transfer of prescription drugs or records.
- (2) The name, address, DEA registration number, Iowa pharmacy license number, and Iowa controlled substances Act (CSA) registration number of the closing pharmacy and of the pharmacy to which prescription drugs will be transferred.
- (3) The name, address, DEA registration number, Iowa pharmacy license number, and CSA registration number of the location at which records will be maintained.

c. Terms of sale or purchase. If the closing is due to the sale of the pharmacy, a copy of the sale or purchase agreement, not including information regarding the monetary terms of the transaction, shall be submitted to the board upon the request of the board. The agreement shall include a written assurance from the closing pharmacy to the purchasing pharmacy that the closing pharmacy has given or will be giving notice to its patients as required by this subrule.

d. Patient notification. At least 30 days prior to closing, a closing pharmacy shall make a reasonable effort to notify all patients who had a prescription filled by the closing pharmacy within the last 18 months that the pharmacy intends to close, including the anticipated closing date.

(1) Written notification shall identify the pharmacy that will be receiving the patient's records. The notification shall advise patients that all patient records will be transferred to the identified pharmacy and that patients may contact the closing pharmacy to request the transfer of remaining refills to a pharmacy of the patient's choice. The notification shall also advise patients that after the date of closing, patients may contact the pharmacy to which the records have been transferred.

(2) Written notification shall be delivered to each patient at the patient's last address on file with the closing pharmacy by direct mail or personal delivery. A pharmacy shall not be required to provide written notice to more than one patient within the same household.

(3) Public notice shall be provided in a location and manner clearly visible to patients in the pharmacy pickup locations including drive-through prescription pickup lanes, on pharmacy or retail store entry and exit doors, and at pharmacy prescription counters.

e. Patient communication by receiving pharmacy. A pharmacy receiving the patient records of another pharmacy shall not contact the patients of the closing pharmacy until after the transfer of those patient records from the closing pharmacy to the receiving pharmacy and after the closure of the closing pharmacy.

f. Prescription drug inventory. A complete inventory of all prescription drugs being transferred shall be taken as of the close of business. The inventory shall serve as the ending inventory for the closing pharmacy as well as a record of additional or starting inventory for the pharmacy to which the drugs are transferred. A copy of the inventory shall be maintained in the records of the purchasing pharmacy for at least two years.

(1) DEA Form 222 is required for transfer of Schedule II controlled substances.

(2) The inventory of controlled substances shall be completed pursuant to the requirements in rule 657—10.19(124).

(3) The inventory of all noncontrolled prescription drugs shall include the name, strength, dosage form, and quantity, which may be estimated.

(4) Controlled substances and prescription drugs requiring destruction or other disposal shall be transferred in the same manner as all other drugs. The new owner is responsible for the disposal of these drugs.

g. Return of certificates and forms. The pharmacy license certificate and CSA registration certificate of the closing or selling pharmacy shall be returned to the board within ten days of closing or sale. The pharmacy shall be responsible for complying with federal DEA regulations for the cancellation and return of DEA forms and certificates.

h. Signs at closed pharmacy location. A location that no longer houses a licensed pharmacy shall not display any sign, placard, or other notification, visible to the public, which identifies the location as a pharmacy. A sign or other public notification that cannot feasibly be removed shall be covered so as to conceal the identification as a pharmacy. Nothing in this paragraph shall prohibit the display of a public notice to patients, as required in paragraph 8.35(7) "d," for a reasonable period not to exceed six months following the pharmacy's closing.

8.35(8) Reporting discipline and criminal convictions. A pharmacy shall, no later than 30 days after the final action, provide written notice to the board of any discipline imposed by any licensing authority on any license or registration held by the pharmacy. Discipline may include, but is not limited to, fine or civil penalty, citation or reprimand, probationary period, suspension, revocation, or voluntary surrender. A pharmacy shall, no later than 30 days after a conviction, provide written notice to the board of any criminal conviction of the pharmacy or of any pharmacy owner when that conviction is related to prescription drugs or to the operation of the pharmacy. The term criminal conviction includes instances when the judgment of conviction or sentence is deferred.

8.35(9) License verification fee. The board may require a nonrefundable fee of \$15 for completion of a request for written license verification of any pharmacy license.

8.35(10) Emergency temporary location changes. In response to a proclamation of disaster emergency or in the event of a natural or man-made disaster, fire, or other occurrence which results in sufficient damage to a pharmacy location as to render it unsafe to operate, a pharmacy may relocate to a temporary or mobile location only as provided herein.

a. Within one business day of the damage to the pharmacy rendering it unsafe, the pharmacy shall provide notice to the board of its intent to temporarily relocate pharmacy operations and provide the address of the temporary or mobile location.

b. A board compliance officer shall conduct an on-site inspection of the temporary or mobile location within five business days of the relocation.

c. A pharmacy may operate from the temporary or mobile location for no more than six months. If the pharmacy is not able to return to the original location within six months, the pharmacy shall submit an application and fee pursuant to paragraph 8.35(6) “b” prior to the expiration of the six-month temporary relocation period.

d. A pharmacy shall notify the board of its intent to return pharmacy operations to the original location at least five business days in advance of its return.

e. A board compliance officer shall conduct an on-site inspection of the original location prior to the return of pharmacy operations.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 9526B, IAB 6/1/11, effective 7/6/11 (See Delay note at end of chapter); ARC 9693B, IAB 9/7/11, effective 8/11/11; ARC 0504C, IAB 12/12/12, effective 1/16/13; ARC 1962C, IAB 4/15/15, effective 5/20/15; ARC 3236C, IAB 8/2/17, effective 9/6/17; ARC 3345C, IAB 9/27/17, effective 11/1/17; ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 4268C, IAB 1/30/19, effective 3/6/19; ARC 5007C, IAB 3/25/20, effective 4/29/20; ARC 6073C, IAB 12/15/21, effective 1/19/22; ARC 6417C, IAB 7/27/22, effective 8/31/22]

657—8.36 to 8.39 Reserved.

657—8.40(155A,84GA,ch63) Pharmacy pilot or demonstration research projects. Rescinded ARC 3858C, IAB 6/20/18, effective 7/25/18.

These rules are intended to implement Iowa Code sections 124.101, 124.301, 124.306, 124.308, 126.10, 126.11, 126.16, 135C.33, 147.7, 147.55, 147.72, 147.74, 147.76, 155A.2 through 155A.4, 155A.6, 155A.10, 155A.12 through 155A.15, 155A.19, 155A.20, 155A.27 through 155A.29, 155A.31 through 155A.35, and 155A.41.

[Filed 4/11/68; amended 11/14/73]

[Filed 11/24/76, Notice 10/20/76—published 12/15/76, effective 1/19/77]

[Filed 11/9/77, Notice 10/5/77—published 11/30/77, effective 1/4/78]

[Filed emergency 12/9/77—published 12/28/77, effective 12/9/77]

[Filed 10/20/78, Notice 8/9/78—published 11/15/78, effective 1/9/79]

[Filed 12/2/78, Notice 11/15/78—published 1/10/79, effective 2/14/79]

[Filed 12/21/78, Notice 11/15/78—published 1/10/79, effective 2/14/79]

[Filed 1/8/79, Notice 11/29/78—published 1/24/79, effective 2/28/79]

[Filed 8/28/79, Notice 5/30/79—published 9/19/79, effective 10/24/79]

[Filed 12/7/79, Notice 10/3/79—published 12/26/79, effective 1/30/80]

[Filed 2/22/80, Notice 10/3/79—published 3/19/80, effective 4/23/80]

[Filed emergency 4/22/80—published 5/14/80, effective 4/22/80]

[Filed 12/1/80, Notice 10/15/80—published 12/24/80, effective 1/28/81]

[Filed 2/12/81, Notice 12/24/80—published 3/4/81, effective 4/8/81]

[Filed 5/27/81, Notice 4/1/81—published 6/24/81, effective 7/29/81]

[Filed emergency 7/28/81—published 8/19/81, effective 8/1/81]

[Filed emergency 9/14/81—published 9/30/81, effective 9/30/81]

[Filed 7/28/82, Notice 3/17/82—published 8/18/82, effective 9/22/82]

[Filed emergency 8/26/82—published 9/15/82, effective 9/22/82]

[Filed 9/10/82, Notice 6/9/82—published 9/29/82, effective 11/8/82]

[Filed emergency 10/6/82—published 10/27/82, effective 10/27/82]^o

[Filed emergency 12/2/82—published 12/22/82, effective 12/22/82]

[Filed 11/18/83, Notice 8/3/83—published 12/7/83, effective 1/11/84]

[Filed 1/13/84, Notice 11/9/83—published 2/1/84, effective 3/7/84]

[Filed 6/22/84, Notice 4/11/84—published 7/18/84, effective 8/22/84]

[Filed emergency 7/13/84—published 8/1/84, effective 7/13/84]

- [Filed 9/21/84, Notice 7/18/84—published 10/10/84, effective 11/14/84]
[Filed 2/22/85, Notice 11/21/84—published 3/13/85, effective 4/18/85]
 [Filed emergency 6/18/85—published 7/3/85, effective 7/1/85]
[Filed 8/30/85, Notice 7/3/85—published 9/25/85, effective 10/30/85]◊
[Filed 11/27/85, Notice 8/28/85—published 12/18/85, effective 1/22/86]
[Filed 9/19/86, Notice 6/4/86—published 10/8/86, effective 11/12/86]
[Filed 1/28/87, Notice 11/19/86—published 2/25/87, effective 4/1/87]
 [Filed emergency 1/21/88—published 2/10/88, effective 1/22/88]
[Filed 1/21/88, Notice 11/4/87—published 2/10/88, effective 3/16/88]
[Filed 3/29/88, Notice 1/27/88—published 4/20/88, effective 5/25/88]
[Filed 3/29/88, Notice 2/10/88—published 4/20/88, effective 5/25/88]
[Filed 11/17/88, Notice 8/24/88—published 12/14/88, effective 1/18/89]◊
 [Filed emergency 5/16/89—published 6/14/89, effective 5/17/89]
[Filed 12/26/89, Notice 10/4/89—published 1/24/90, effective 2/28/90]
[Filed 3/19/90, Notice 1/10/90—published 4/18/90, effective 5/23/90]
[Filed 8/31/90, Notice 6/13/90—published 9/19/90, effective 10/24/90]
[Filed 1/29/91, Notice 6/13/90—published 2/20/91, effective 3/27/91]
[Filed 1/29/91, Notice 9/19/90—published 2/20/91, effective 3/27/91]
[Filed 4/26/91, Notice 2/20/91—published 5/15/91, effective 6/19/91]
 [Filed emergency 5/10/91—published 5/29/91, effective 5/10/91]
[Filed 7/30/91, Notice 5/29/91—published 8/21/91, effective 9/25/91]
[Filed 1/21/92, Notice 10/16/91—published 2/19/92, effective 3/25/92]
 [Filed 3/12/92, Notice 1/8/92—published 4/1/92, effective 5/6/92]
 [Filed 5/21/92, Notice 4/1/92—published 6/10/92, effective 7/15/92]
 [Filed 10/22/92, Notice 9/2/92—published 11/11/92, effective 1/1/93]
 [Filed 2/5/93, Notice 11/11/92—published 3/3/93, effective 4/8/93]
[Filed 9/23/93, Notice 5/26/93—published 10/13/93, effective 11/17/93]
[Filed 3/21/94, Notices 10/13/93, 12/8/93—published 4/13/94, effective 5/18/94]
 [Filed 6/24/94, Notice 4/13/94—published 7/20/94, effective 8/24/94]
[Filed 11/30/94, Notices 5/11/94, 7/20/94—published 12/21/94, effective 1/25/95]
 [Filed 3/22/95, Notice 11/9/94—published 4/12/95, effective 5/31/95]
[Filed 10/6/95, Notices 6/7/95, 8/16/95—published 10/25/95, effective 1/1/96]
 [Filed emergency 12/14/95—published 1/3/96, effective 1/1/96]
 [Filed 12/10/96, Notice 8/28/96—published 1/1/97, effective 2/5/97]
 [Filed 2/27/97, Notice 8/28/96—published 3/26/97, effective 4/30/97]
 [Filed 2/27/97, Notice 1/1/97—published 3/26/97, effective 4/30/97]
 [Filed 6/23/97, Notice 3/26/97—published 7/16/97, effective 8/20/97]
[Filed 11/19/97, Notice 10/8/97—published 12/17/97, effective 1/21/98]
 [Filed 4/24/98, Notice 3/11/98—published 5/20/98, effective 6/24/98]
 [Filed 7/31/98, Notice 5/20/98—published 8/26/98, effective 10/15/98]
 [Filed 4/22/99, Notice 3/10/99—published 5/19/99, effective 6/23/99]
 [Filed 11/23/99, Notice 6/2/99—published 12/15/99, effective 1/19/00]
 [Filed 2/18/00, Notice 12/15/99—published 3/22/00, effective 4/26/00]
 [Filed 11/9/00, Notice 4/19/00—published 11/29/00, effective 1/3/01]
 [Filed 8/14/02, Notice 6/12/02—published 9/4/02, effective 10/9/02]
 [Filed 3/11/04, Notice 8/6/03—published 3/31/04, effective 5/5/04]
[Filed emergency 7/16/04 after Notice 6/9/04—published 8/4/04, effective 7/16/04]
[Filed 10/22/04, Notice 3/31/04—published 11/10/04, effective 12/15/04]
[Filed 10/22/04, Notice 5/12/04—published 11/10/04, effective 12/15/04]
 [Filed 6/2/05, Notice 3/16/05—published 6/22/05, effective 7/27/05]
[Filed emergency 6/30/05 after Notice 5/11/05—published 7/20/05, effective 7/1/05]
 [Filed 3/22/06, Notice 1/18/06—published 4/12/06, effective 5/17/06]

- [Filed 5/17/06, Notice 4/12/06—published 6/7/06, effective 7/12/06]
[Filed 5/17/06, Notice 2/15/06—published 6/7/06, effective 10/1/06]
[Filed 11/30/06, Notice 9/27/06—published 12/20/06, effective 1/24/07]
[Filed 2/7/07, Notice 10/25/06—published 2/28/07, effective 4/4/07]
[Filed 5/14/07, Notice 2/28/07—published 6/6/07, effective 7/11/07]^o
[Filed 8/3/07, Notice 5/9/07—published 8/29/07, effective 10/3/07]
[Filed 8/3/07, Notice 6/20/07—published 8/29/07, effective 10/3/07]
[Filed emergency 11/13/07 after Notice 8/29/07—published 12/5/07, effective 11/13/07]
[Filed 11/13/07, Notice 8/29/07—published 12/5/07, effective 1/9/08]
[Filed 5/19/08, Notice 3/26/08—published 6/18/08, effective 7/23/08]
[Filed 9/5/08, Notice 7/2/08—published 9/24/08, effective 10/29/08]
[Filed ARC 7636B (Notice ARC 7448B, IAB 12/31/08), IAB 3/11/09, effective 4/15/09]
[Filed ARC 8171B (Notice ARC 7910B, IAB 7/1/09), IAB 9/23/09, effective 10/28/09]
[Filed ARC 8539B (Notice ARC 8269B, IAB 11/4/09), IAB 2/24/10, effective 4/1/10]
[Filed ARC 8673B (Notice ARC 8380B, IAB 12/16/09), IAB 4/7/10, effective 6/1/10]
[Filed ARC 8671B (Notice ARC 8414B, IAB 12/30/09), IAB 4/7/10, effective 5/12/10]
[Filed ARC 9409B (Notice ARC 9194B, IAB 11/3/10), IAB 3/9/11, effective 4/13/11]
[Filed ARC 9526B (Notice ARC 9295B, IAB 12/29/10), IAB 6/1/11, effective 7/6/11]¹
[Editorial change: IAC Supplement 6/29/11]
[Filed Emergency ARC 9693B, IAB 9/7/11, effective 8/11/11]
[Filed ARC 9912B (Notice ARC 9671B, IAB 8/10/11), IAB 12/14/11, effective 1/18/12]
[Filed ARC 0393C (Notice ARC 0256C, IAB 8/8/12), IAB 10/17/12, effective 11/21/12]
[Filed ARC 0503C (Notice ARC 0371C, IAB 10/3/12), IAB 12/12/12, effective 1/16/13]
[Filed ARC 0504C (Notice ARC 0351C, IAB 10/3/12), IAB 12/12/12, effective 1/16/13]
[Filed Emergency After Notice ARC 1030C (Notice ARC 0883C, IAB 7/24/13), IAB 9/18/13,
effective 9/1/13]
[Filed ARC 1032C (Notice ARC 0882C, IAB 7/24/13), IAB 9/18/13, effective 10/23/13]
[Filed ARC 1576C (Notice ARC 1411C, IAB 4/2/14), IAB 8/20/14, effective 9/24/14]
[Filed ARC 1786C (Notice ARC 1652C, IAB 10/1/14), IAB 12/10/14, effective 1/14/15]
[Filed ARC 1961C (Notice ARC 1793C, IAB 12/10/14), IAB 4/15/15, effective 5/20/15]
[Filed ARC 1962C (Notice ARC 1792C, IAB 12/10/14), IAB 4/15/15, effective 5/20/15]
[Filed ARC 2408C (Notice ARC 2285C, IAB 12/9/15), IAB 2/17/16, effective 3/23/16]
[Filed ARC 2413C (Notice ARC 2307C, IAB 12/9/15), IAB 2/17/16, effective 3/23/16]
[Filed ARC 2414C (Notice ARC 2288C, IAB 12/9/15), IAB 2/17/16, effective 3/23/16]
[Filed Emergency After Notice ARC 2827C (Notice ARC 2721C, IAB 9/28/16), IAB 11/23/16,
effective 11/3/16]
[Filed ARC 3236C (Notice ARC 3037C, IAB 4/26/17), IAB 8/2/17, effective 9/6/17]
[Filed ARC 3345C (Notice ARC 3136C, IAB 6/21/17), IAB 9/27/17, effective 11/1/17]
[Filed ARC 3639C (Notice ARC 3371C, IAB 10/11/17), IAB 2/14/18, effective 3/21/18]
[Filed ARC 3858C (Notice ARC 3509C, IAB 12/20/17), IAB 6/20/18, effective 7/25/18]
[Filed ARC 4267C (Notice ARC 4029C, IAB 9/26/18), IAB 1/30/19, effective 3/6/19]
[Filed ARC 4268C (Notice ARC 4092C, IAB 10/24/18), IAB 1/30/19, effective 3/6/19]
[Filed ARC 4454C (Amended Notice ARC 4172C, IAB 12/19/18; Notice ARC 3978C, IAB 8/29/18),
IAB 5/22/19, effective 6/26/19]
[Filed ARC 4580C (Notice ARC 4386C, IAB 4/10/19), IAB 7/31/19, effective 9/4/19]
[Filed ARC 4903C (Notice ARC 4693C, IAB 10/9/19), IAB 2/12/20, effective 3/18/20]
[Filed ARC 5007C (Notice ARC 4695C, IAB 10/9/19), IAB 3/25/20, effective 4/29/20]
[Filed ARC 5348C (Notice ARC 5113C, IAB 7/29/20), IAB 12/30/20, effective 2/3/21]
[Filed ARC 5350C (Notice ARC 5115C, IAB 7/29/20), IAB 12/30/20, effective 2/3/21]
[Filed ARC 6073C (Notice ARC 5835C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]
[Filed ARC 6076C (Notice ARC 5833C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]

[Filed ARC 6330C (Amended Notice ARC 6179C, IAB 2/9/22; Notice ARC 5834C, IAB 8/11/21),
IAB 6/1/22, effective 7/6/22]

[Filed ARC 6417C (Notice ARC 6281C, IAB 4/6/22), IAB 7/27/22, effective 8/31/22]

[Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

⁰ Two or more ARCs

¹ July 6, 2011, effective date of 8.35(7) delayed 70 days by the Administrative Rules Review Committee at its meeting held June 14, 2011.

CHAPTER 18
CENTRALIZED PRESCRIPTION FILLING AND PROCESSING

657—18.1(155A) Purpose and scope. The purpose of this chapter is to provide standards for centralized prescription drug order filling or centralized prescription processing by a pharmacy. Any facility established for the purpose of filling or processing prescription drug orders on behalf of other pharmacies shall be licensed as a pharmacy and shall hold all necessary registrations. A hospital pharmacy may participate in centralized prescription filling only of prescription drug orders for noncontrolled substances pursuant to these rules. A hospital pharmacy may engage in centralized prescription processing pursuant to the requirements of rule 657—7.7(155A). Except as specifically identified in the rules, the requirements of these rules for centralized prescription filling or centralized prescription processing are in addition to the requirements of 657—Chapters 6, 7, and 8, and other rules of the board relating to services provided by pharmacies.

657—18.2(155A) Definitions. For the purposes of this chapter, the following definitions shall apply:

“*Central fill pharmacy*” means a pharmacy contracting with an originating pharmacy, or having the same owner as an originating pharmacy, that provides centralized prescription drug order filling on behalf of the originating pharmacy pursuant to these rules.

“*Centralized prescription drug order filling*” or “*centralized filling*” means the filling of a prescription drug order by a pharmacy on behalf of another pharmacy. “Centralized filling” does not include the processing or dispensing of a prescription drug order but may include any of the following filling functions:

1. Receiving prescription drug orders from the originating pharmacy;
2. Interpreting or clarifying prescription drug orders;
3. Entering prescription drug order information into a pharmacy’s prescription record system;
4. Selecting, counting, and placing the prescribed drug into an appropriate prescription container;
5. Affixing the prescription label, including any auxiliary labels, to the prescription container;
6. Obtaining refill and substitution authorizations;
7. Verifying all filling processes performed by the central fill pharmacy.

“*Centralized prescription drug order processing*” or “*centralized processing*” means the processing of a prescription drug order by a pharmacy on behalf of another pharmacy. “Centralized processing” does not include the filling or dispensing of a prescription drug order but may include any of the following processing functions:

1. Interpreting or clarifying prescription drug orders;
2. Entering prescription drug order information into a pharmacy’s prescription record system;
3. Interpreting clinical data for prior authorization for dispensing;
4. Performing formulary-directed therapeutic interchange.

“*Central processing pharmacy*” means a pharmacy contracting with an originating pharmacy, or having the same owner as an originating pharmacy, that provides centralized prescription drug order processing on behalf of the originating pharmacy pursuant to these rules.

“*DEA*” means the U.S. Department of Justice, Drug Enforcement Administration.

“*Dispense*” means the delivery of a prescription drug or device to an ultimate user or the ultimate user’s agent by or pursuant to the lawful order of a practitioner. “Dispense” includes:

1. Receiving the prescription drug order from the patient, the patient’s agent, or the prescriber;
2. Delivering the filled prescription to the patient or the patient’s agent;
3. Providing drug information concerning a patient’s drug therapy;
4. Providing patient counseling;
5. Providing medication therapy management.

“*Hospital*” means a facility licensed pursuant to Iowa Code chapter 135B.

“*Hospital pharmacy*” means and includes a pharmacy licensed by the board and located within any hospital, health system, institution, or establishment which maintains and operates organized facilities

for the diagnosis, care, and treatment of human illnesses to which persons may or may not be admitted for overnight stay at the facility.

“*Mail order pharmacy*” means a pharmacy located within a United States jurisdiction whose primary business is to dispense a prescription drug or device pursuant to a valid prescription drug order and to deliver the drug or device to a patient, including a patient in this state, via the United States Postal Service, a common carrier, or a delivery service. “Mail order pharmacy” includes a pharmacy that does business via the Internet or other electronic media.

“*Medication therapy management*” means the review of drug therapy regimens of a patient by a pharmacist for the purpose of evaluating and rendering advice to a practitioner, or for the purpose of evaluating and modifying the drug regimen in accordance with a collaborative drug therapy management protocol pursuant to rule 657—39.13(155A).

“*Originating pharmacy*” means a pharmacy that receives a prescription drug order from a patient, the patient’s agent, or a prescriber, outsources prescription filling or processing functions to another pharmacy, and ultimately dispenses the prescription drug or device to the patient or the patient’s agent. [ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—18.3(155A) General requirements.

18.3(1) Essential qualifications. An originating pharmacy may outsource prescription drug filling to a central fill pharmacy or prescription drug order processing to a central processing pharmacy provided the pharmacies:

a. Have the same owner or have entered into a written contract or agreement, which is available for inspection and copying by the board or its authorized agent, that outlines the services to be provided and the responsibilities and accountabilities of each pharmacy in compliance with federal and state laws, rules, and regulations; and

b. Share a common electronic file or have appropriate technology to allow access to sufficient information necessary or required to perform the contracted functions.

18.3(2) Legal compliance. An originating pharmacy, a central fill pharmacy, and a central processing pharmacy shall comply with all provisions applicable to the pharmacy contained in federal and state laws, rules, and regulations to the extent applicable for the specific filling or processing activity and these rules, including but not limited to the following:

a. Each pharmacy located within Iowa shall maintain Iowa pharmacy licensure and, if the pharmacy dispenses controlled substances, the pharmacy shall maintain DEA and Iowa controlled substances registrations.

b. Each pharmacy located outside Iowa shall maintain Iowa nonresident pharmacy licensure in addition to the licensure requirements of the pharmacy’s home state.

c. Each pharmacist providing centralized prescription drug order processing or filling functions as an employee or agent of a central processing or central fill pharmacy located within Iowa shall maintain active licensure to practice pharmacy in Iowa.

d. Pharmacies shall comply with Iowa board rules relating to the duties that must be performed by a pharmacist.

e. Pharmacies shall comply with Iowa requirements for supervision of pharmacy technicians and pharmacy support persons.

18.3(3) Originating pharmacy responsibility. Except as specifically provided by this subrule, the originating pharmacy shall be responsible for all dispensing functions as the term “dispense” is defined in rule 657—18.2(155A). An originating pharmacy contracting only for centralized filling shall retain responsibility for all processing functions, and an originating pharmacy contracting only for centralized processing shall retain responsibility for all filling functions.

a. A mail order pharmacy engaged in the centralized filling of prescription drug orders may deliver a filled prescription directly to the patient and shall not be required to return the filled prescription to the originating pharmacy.

b. A central fill or a central processing pharmacy that shares a common central processing unit with the originating pharmacy may perform prospective drug use review (DUR) pursuant to

rule 657—8.21(155A). Only a pharmacist shall perform the DUR, and such review shall not be delegated. The pharmacist performing the DUR shall document in the shared patient record all concerns, recommendations, observations, and comments resulting from that review. The pharmacist at the originating pharmacy shall utilize the DUR notes in counseling the patient pursuant to rule 657—6.14(155A).

18.3(4) Central fill label requirements. The label affixed to the prescription container filled by a central fill pharmacy on behalf of an originating pharmacy shall include the following:

- a. A unique identifier indicating that the prescription was filled at the central fill pharmacy;
- b. Serial number (a unique identification number of the prescription) as assigned by the originating pharmacy;
- c. The name, address, and telephone number of the originating pharmacy;
- d. The name of the patient or, if such drug is prescribed for an animal, the species of the animal and the name of its owner, except as provided in 657—subrule 8.19(7) for epinephrine auto-injectors, 657—subrule 8.19(8) for opioid antagonists, 657—subrule 8.19(9) for expedited partner therapy, or 657—subrule 8.19(10) for bronchodilator canisters or bronchodilator canisters and spacers.
- e. The name of the prescribing practitioner;
- f. The date the prescription is filled by the central fill pharmacy;
- g. The directions or instructions for use, including precautions to be observed;
- h. Unless otherwise directed by the prescriber, the name, strength, and quantity of the drug dispensed.

(1) If a pharmacist selects an equivalent drug product for a brand name drug product prescribed by a practitioner, the prescription container label shall identify the generic drug and may identify the brand name drug for which the selection is made, such as “(generic name) Generic for (brand name product)”;

(2) If a pharmacist selects a brand name drug product for a generic drug product prescribed by a practitioner, the prescription container label shall identify the brand name drug product dispensed and may identify the generic drug product ordered by the prescriber, such as “(brand name product) for (generic name)”;

(3) If a pharmacist selects an interchangeable biological product for the biological product prescribed by a practitioner, the prescription container label shall identify the interchangeable biological product dispensed and may identify the biological product prescribed by the practitioner, such as “(interchangeable biological product) for (biological product)”;

i. The initials or other unique identification of the pharmacist who performed drug use review, unless the identification of the pharmacist involved in each step of the prescription filling process is electronically documented and retrievable.

[ARC 8673B, IAB 4/7/10, effective 6/1/10; ARC 3863C, IAB 6/20/18, effective 7/25/18; ARC 3985C, IAB 8/29/18, effective 10/3/18; ARC 4903C, IAB 2/12/20, effective 3/18/20; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—18.4 Reserved.

657—18.5(155A) Patient notification and authorization.

18.5(1) Prior notification and authorization. A pharmacy that outsources prescription drug order filling or prescription drug order processing to another pharmacy shall, prior to outsourcing a patient’s prescription:

a. Notify the patient or the patient’s agent that prescription filling or processing may be outsourced to another pharmacy.

b. Provide the name of the pharmacy that will be filling or processing the prescription or, if the pharmacy is part of a network of pharmacies under common ownership and any of the network pharmacies may fill or process the prescription, the patient shall be notified of this fact. Notification shall be provided through a notice to the patient or the patient’s agent by means of a sign prominently displayed in the originating pharmacy and through written notice provided to the patient or the patient’s agent prior to implementation of the program or upon commencement of services to a new patient, as applicable.

c. If a patient provides the originating pharmacy with notification that the patient no longer authorizes the originating pharmacy to outsource the patient's prescription drug orders, the originating pharmacy shall discontinue outsourcing the filling or processing of the patient's prescription drug orders.

18.5(2) Exception. The provisions of this rule do not apply to a patient in a facility, such as a hospital or care facility, where Iowa law requires that drugs be administered to the patient by a health care professional.

[ARC 3863C, IAB 6/20/18, effective 7/25/18]

657—18.6 to 18.9 Reserved.

657—18.10(155A) Policy and procedures. Pursuant to rule 657—8.3(155A), a policy and procedure manual relating to centralized filling or centralized processing activities shall be maintained at all pharmacies involved in centralized filling or centralized processing and shall be available for inspection and copying by the board or its authorized agent. The manual shall:

1. Outline the responsibilities of each of the pharmacies;
2. Include a list of the names, addresses, telephone numbers, and all license and registration numbers of the pharmacies involved in centralized filling or centralized processing; and
3. Include, but not necessarily be limited to, policies and procedures for:
 - Protecting the confidentiality and integrity of patient information;
 - Protecting each patient's freedom of choice of pharmacy services;
 - Maintaining appropriate records to identify the name, the initials or unique identification code, and the specific activities of each pharmacist or pharmacy technician who performed any centralized filling or centralized processing function; and
 - Operating a continuous quality improvement program for pharmacy services designed to objectively and systematically monitor and evaluate the quality and appropriateness of patient care, pursue opportunities to improve patient care, and resolve identified problems.

[ARC 1961C, IAB 4/15/15, effective 5/20/15; ARC 3863C, IAB 6/20/18, effective 7/25/18]

657—18.11 to 18.14 Reserved.

657—18.15(155A) Records. Central fill or central processing pharmacies shall maintain appropriate records that identify, by prescription drug order, the initials or unique identification code of each pharmacist or pharmacy technician who performs a centralized filling or centralized processing function for a prescription drug order. Originating pharmacies shall maintain appropriate records that identify, by prescription drug order, the initials or unique identification code of the pharmacist who performed drug use review. These records may be maintained separately by each pharmacy or in a common electronic file as long as the data processing system is capable of producing a printout that lists the functions performed by each pharmacy and pharmacist or technician and identifies the pharmacist or technician who performed each function.

[ARC 3863C, IAB 6/20/18, effective 7/25/18]

These rules are intended to implement Iowa Code sections 124.301, 124.306, 124.308, 155A.13, and 155A.28.

[Filed 6/2/05, Notice 1/19/05—published 6/22/05, effective 7/27/05]

[Filed 3/6/08, Notice 12/19/07—published 3/26/08, effective 4/30/08¹]

[Filed emergency 6/9/08—published 7/2/08, effective 7/9/08]

[Filed 11/24/08, Notice 10/8/08—published 12/17/08, effective 1/21/09]

[Filed ARC 8673B (Notice ARC 8380B, IAB 12/16/09), IAB 4/7/10, effective 6/1/10]

[Filed ARC 1961C (Notice ARC 1793C, IAB 12/10/14), IAB 4/15/15, effective 5/20/15]

[Filed ARC 3858C (Notice ARC 3509C, IAB 12/20/17), IAB 6/20/18, effective 7/25/18]

[Filed ARC 3863C (Notice ARC 3512C, IAB 12/20/17), IAB 6/20/18, effective 7/25/18]

[Filed ARC 3985C (Notice ARC 3764C, IAB 4/25/18), IAB 8/29/18, effective 10/3/18]

[Filed ARC 4903C (Notice ARC 4693C, IAB 10/9/19), IAB 2/12/20, effective 3/18/20]

[Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

¹ April 30, 2008, effective date of ARC 6671B delayed 70 days by the Administrative Rules Review Committee at its meeting held April 4, 2008.

CHAPTER 21
ELECTRONIC DATA AND AUTOMATED SYSTEMS IN PHARMACY PRACTICE

657—21.1(124,155A) Purpose and scope. The purpose of this chapter is to provide the minimum standards for the utilization of electronic data and automated systems in the practice of pharmacy and shall apply to all pharmacies located in Iowa.

[ARC 3640C, IAB 2/14/18, effective 3/21/18]

657—21.2(124,155A) Definitions. For the purpose of this chapter, the following definitions shall apply:

“Automated data processing system” means an application that is used for prescription, patient, drug, and prescriber information; installed on a pharmacy’s computer or server; and controlled by the pharmacy.

“Automated medication distribution system” or *“AMDS”* includes, but is not limited to, an automated device or series of devices operated by an electronic interface with one or more computers that is used to prepare, package, or dispense specified dosage units of drugs for administration or dispensing. *“AMDS”* does not include electronic storage devices that do not have an electronic interface with one or more computers of the pharmacy.

“CSA” means the Iowa uniform controlled substances Act.

“CSA registration” means the registration issued by the board pursuant to the CSA that signifies the registrant’s authorization to engage in registered activities with controlled substances.

“DEA” means the U.S. Department of Justice, Drug Enforcement Administration.

“Electronically prepared prescription” means a prescription that is generated utilizing an electronic prescription application.

“Electronic device” means an electronic, mechanical, or other device which is used to intercept communications and includes but is not limited to network, file and print servers; desktop workstations; laptop computers; tablets; mini-computers; smart phones; and similar devices.

“Electronic prescription” means an electronically prepared prescription that is authorized and transmitted from the prescriber to the pharmacy by means of electronic transmission.

“Electronic prescription application” means software that is used to create electronic prescriptions and that is intended to be installed on a prescriber’s computers and servers where access and records are controlled by the prescriber.

“Electronic signature” means a confidential personalized digital key, code, number, or other method used for secure electronic data transmissions which identifies a particular person as the source of the message, authenticates the signatory of the message, and indicates the person’s approval of the information contained in the transmission.

“Electronic transmission” means the transmission of an electronic prescription, formatted as an electronic data file, from a prescriber’s electronic prescription application to a pharmacy’s computer, where the data file is imported into the pharmacy prescription application.

“Facsimile transmission” or *“fax transmission”* means the transmission of a digital image of a prescription from the prescriber or the prescriber’s agent to the pharmacy. *“Facsimile transmission”* includes but is not limited to transmission of a written prescription between the prescriber’s fax machine and the pharmacy’s fax machine; transmission of an electronically prepared prescription from the prescriber’s electronic prescription application to the pharmacy’s fax machine or printer; or transmission of an electronically prepared prescription from the prescriber’s fax machine to the pharmacy’s fax machine, computer, or printer.

“Intermediary” means any technology system that receives and transmits an electronic prescription between the prescriber and the pharmacy.

“Pharmacist verification” or *“verified by a pharmacist”* means the accuracy of a prescription drug is verified by a pharmacist, pharmacist-intern, or technician in an approved technician product verification program.

“*Prescription drug order*” or “*prescription*” means a lawful order of a practitioner for a drug or device for a specific patient that is communicated to a pharmacy, regardless of whether the communication is oral, electronic, via facsimile, or in printed form.

“*Readily retrievable*” means that hard-copy or electronic records can be separated out from all other records within 72 hours of a request from the board or other authorized agent.

“*Written prescription*” means a prescription that is created on paper, a prescription that is electronically prepared and printed, or a prescription that is electronically prepared and transmitted from the prescriber’s electronic device to a pharmacy via facsimile. A written prescription for a controlled substance shall be manually signed by the prescriber in compliance with federal and state laws, rules, and regulations.

[ARC 3640C, IAB 2/14/18, effective 3/21/18; ARC 4580C, IAB 7/31/19, effective 9/4/19; ARC 5007C, IAB 3/25/20, effective 4/29/20]

657—21.3(124,155A) System security and safeguards. To maintain the integrity and confidentiality of patient records and prescription drug orders, any system, computer, or electronic device utilized shall have adequate security including system safeguards designed to prevent and detect unauthorized access, modification, or manipulation of patient records and prescription drug orders. Authentication credentials shall be securely maintained by the individual to whom the credentials are issued and shall not be shared with or disclosed to any other individual. Once a drug or device has been dispensed, any alterations in either the prescription drug order data or the patient record shall be documented and shall include the identification of all pharmacy personnel who were involved in making the alteration as well as the responsible pharmacist. An automated data processing system used for the receipt and processing of electronic transmissions from a prescriber’s electronic prescription application shall comply with DEA requirements relating to electronic prescriptions and shall be certified compliant with DEA regulations.
[ARC 3640C, IAB 2/14/18, effective 3/21/18]

657—21.4 Reserved.

657—21.5(124,155A) Automated data processing systems. An automated data processing system may be used, subject to the requirements contained in this rule, for the storage and retrieval of prescription, patient, prescriber and drug data as well as data relating to the pharmacy staff utilization of the system.

21.5(1) *Electronic storage of hard-copy prescriptions.* A pharmacy that maintains an electronic copy of an original hard-copy prescription for a noncontrolled substance shall retain, in a readily retrievable format, the original hard-copy prescription as required in rule 657—6.8(155A) but shall be exempt from the requirement to record on the original hard-copy prescription the date and unique identification number of the prescription.

21.5(2) *Data retrievable and printable.* Any automated data processing system shall be capable of immediate retrieval (via computer monitor or hard-copy printout) of, at a minimum, any prescription, patient, prescriber, and drug data as well as data relating to pharmacy staff utilization of the system.

21.5(3) *Auxiliary procedure for system downtime.* A pharmacy utilizing an automated data processing system shall have a procedure that will maintain security and confidentiality of all data as well as ensure the legal dispensing of any prescription drug order in the event the system experiences downtime.

[ARC 3640C, IAB 2/14/18, effective 3/21/18]

657—21.6(124,155A) Electronic prescription applications. Each prescription shall be transmitted electronically to a pharmacy except as provided in rule 657—21.8(124,155A). The prescription drug order shall contain all information required by Iowa Code section 155A.27. The receiving pharmacist shall be responsible for verifying the authenticity of an electronically prescribed prescription pursuant to rule 657—8.19(124,126,155A). A prescription that is subject to exemption as provided in rule 657—21.8(124,155A) may be transmitted to a pharmacy via electronic or facsimile transmission or printed in hard-copy format for delivery to the pharmacy. A prescription that is transmitted by a

prescriber's agent via electronic or facsimile transmission shall include the first and last names and title of the agent responsible for the transmission.

21.6(1) *Electronic transmission.* A prescription prepared pursuant to this rule shall be transmitted electronically to a pharmacy, unless exempt pursuant to rule 657—21.8(124,155A). A pharmacy shall be certified compliant with DEA regulations relating to electronic prescriptions prior to electronically receiving prescriptions for controlled substances. The electronic record shall serve as the original record and shall be maintained for two years from the date of last activity on the prescription. Any annotations shall be made and retained on the electronic record.

a. An electronically prepared and transmitted prescription that is printed following transmission shall be clearly labeled as a copy, not valid for dispensing.

b. The authenticity of a prescription transmitted via electronic transmission between a DEA-certified electronic prescription application and a DEA-certified electronic automated data processing system shall be deemed verified by virtue of the security processes included in those applications.

c. A pharmacy shall ensure that no intermediary has the ability to change the content of the prescription drug order or compromise its confidentiality during the transmission process. The electronic format of the prescription drug order may be changed by the intermediary to facilitate the transmission between electronic applications as long as the content of the prescription drug order remains unchanged.

d. In addition to the information requirements for a prescription, an electronically transmitted prescription shall identify the transmitter's telephone number for verbal confirmation, the telephone number where the prescriber can be contacted for timely consultation about patient care matters, the time and date of transmission, and the pharmacy intended to receive the transmission as well as any other information required by federal or state laws, rules, or regulations.

e. If the transmission of an electronic prescription fails, the prescriber may print the prescription, manually sign the printed prescription, and deliver the prescription to the pharmacy via facsimile transmission in accordance with subrule 21.6(2).

21.6(2) *Printed (hard-copy) prescriptions.* A prescription that is exempt from the electronic prescription mandate as provided in rule 657—21.8(124,155A) may be printed in hard-copy format for facsimile transmission or delivery to the pharmacy.

a. A prescription for a controlled substance shall include the prescriber's manual signature. Printed or hard-copy prescriptions for Schedule II controlled substances shall not be transmitted to a pharmacy via facsimile transmission, except as authorized in rule 657—21.7(124,155A).

b. If the prescriber authenticates a prescription for a noncontrolled prescription drug utilizing an electronic signature, the printed prescription shall be printed on security paper. Security features of the paper shall ensure that prescription information is not obscured or rendered illegible when transmitted via facsimile or when scanned into an electronic record system.

c. If the facsimile transmission of a printed prescription is a result of a failed electronic transmission, the facsimile shall indicate that it was originally transmitted to the named pharmacy, the date and time of the original electronic transmission, and the fact that the original transmission failed.

[ARC 3640C, IAB 2/14/18, effective 3/21/18; ARC 4580C, IAB 7/31/19, effective 9/4/19; ARC 5350C, IAB 12/30/20, effective 2/3/21; ARC 6077C, IAB 12/15/21, effective 1/19/22]

657—21.7(124,155A) *Facsimile transmission of a prescription.* A pharmacist may dispense noncontrolled and controlled drugs, including Schedule II controlled substances only as provided in this rule, pursuant to a prescription faxed to the pharmacy by the prescribing practitioner or the practitioner's agent. The means of transmission via facsimile shall ensure that prescription information is not obscured or rendered illegible due to security features of the paper utilized by the prescriber to prepare a written prescription. The faxed prescription shall serve as the original record, except as provided in subrule 21.7(1), shall be maintained for a minimum of two years from the date of the last activity on the prescription, and shall contain all information required by Iowa Code section 155A.27, including the prescriber's signature. If the prescription is transmitted by an agent of the prescriber, the facsimile transmission shall include the first and last names and title of the agent responsible for the

transmission. The pharmacist shall be responsible for verifying the authenticity of the prescription as to the source of the facsimile transmission.

21.7(1) *Schedule II controlled substances—emergency situations.* A pharmacist may, in an emergency situation as defined in 657—subrule 10.26(1), dispense a Schedule II controlled substance pursuant to a facsimile transmission to the pharmacy of a written, signed prescription from the prescriber or the prescriber’s agent pursuant to the requirements of rule 657—10.26(124). The facsimile shall serve as the temporary written record required by 657—subrule 10.26(2).

21.7(2) *Schedule II controlled substances—compounded injectable.* A prescription for a Schedule II narcotic substance to be compounded for the direct administration to a patient by parenteral, intravenous, intramuscular, subcutaneous, or intraspinal infusion may be transmitted by a prescriber or the prescriber’s agent to a pharmacy via facsimile.

21.7(3) *Schedule II controlled substances—care facility patients.* A prescription for any Schedule II controlled substance for a resident of a care facility, as defined in rule 657—23.2(155A), may be transmitted by the prescriber or the prescriber’s agent to a pharmacy via facsimile. The prescription shall identify that the patient is a resident of a care facility.

21.7(4) *Schedule II controlled substances—hospice patients.* A prescription for any Schedule II controlled substance for a patient in a hospice program licensed pursuant to Iowa Code chapter 135J or a program certified or paid for by Medicare under Title XVIII may be transmitted via facsimile by the prescriber or the prescriber’s agent to the pharmacy. The prescription shall identify that the patient is a hospice patient.

[ARC 3640C, IAB 2/14/18, effective 3/21/18; ARC 5350C, IAB 12/30/20, effective 2/3/21; ARC 6077C, IAB 12/15/21, effective 1/19/22]

657—21.8(124,155A) Electronic prescription mandate and exemptions. All prescriptions shall be transmitted electronically to a pharmacy except as provided in this rule.

21.8(1) *Prescriptions exempt.* Prescriptions which shall be exempt from electronic transmission include:

- a. A prescription for a patient residing in a nursing home, long-term care facility, correctional facility, or jail.
- b. A prescription authorized by a licensed veterinarian.
- c. A prescription for a device.
- d. A prescription dispensed by a department of veterans affairs pharmacy.
- e. A prescription requiring information that makes electronic transmission impractical, such as complicated or lengthy directions for use or attachments.
- f. A prescription for a compounded preparation containing two or more components.
- g. A prescription issued in response to a public health emergency in a situation where a non-patient-specific prescription would be permitted.
- h. A prescription issued for an opioid antagonist pursuant to Iowa Code section 135.190, a prescription issued for epinephrine pursuant to Iowa Code section 135.185, or a prescription issued for a bronchodilator canister or bronchodilator canister and spacer pursuant to Iowa Code section 280.16A.
- i. A prescription issued during a temporary technical or electronic failure at the location of the prescriber or pharmacy, provided that a prescription issued pursuant to this paragraph shall indicate on the prescription that the prescriber or pharmacy is experiencing a temporary technical or electronic failure.
- j. A prescription issued pursuant to an established and valid collaborative practice agreement, standing order, or drug research protocol.
- k. A prescription issued in an emergency situation pursuant to federal law and regulation and rules of the board. An emergency situation may include, but is not limited to, the issuance of a prescription to meet the immediate care need of a patient after hours when a prescriber is unable to access electronic prescribing capabilities. Such prescription shall be limited to a quantity sufficient to meet the acute need of the patient with no authorized refills.

21.8(2) *Prescriber, medical group, institution, or pharmacy exemption.* A prescriber, medical group, institution, or pharmacy that has been granted an exemption to the electronic prescription

mandate pursuant to rule 657—21.9(124,155A) shall be exempt from the electronic prescription mandate only for the duration of the approved exemption, and the exemption shall not apply retroactively to prescriptions issued prior to approval. Upon expiration of an approved exemption, the prescriber, medical group, institution, or pharmacy shall either comply with the electronic prescription mandate or timely petition the board for renewal of the exemption pursuant to rule 657—21.9(124,155A). A prescriber, medical group, institution or pharmacy that has been granted an exemption to the electronic prescription mandate pursuant to rule 657—21.9(124,155A) shall identify the exemption on each prescription issued and transmitted by any nonelectronic means. A pharmacist shall not be required to verify that the prescription or prescriber is subject to an exemption.

[ARC 4580C, IAB 7/31/19, effective 9/4/19; ARC 6077C, IAB 12/15/21, effective 1/19/22; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—21.9(124,155A) Exemption from electronic prescription mandate—petition. A prescriber, medical group, institution, or pharmacy that is unable to comply with the electronic prescription mandate in rule 657—21.8(124,155A) may petition the board, on forms provided by the board, for an exemption from the requirements based upon economic hardship; technical limitations that the prescriber, medical group, institution, or pharmacy cannot control; or other exceptional circumstances. A timely petition for renewal of a previously approved exemption shall be submitted at least 60 days in advance of the expiration of the previously approved exemption.

21.9(1) Petition information. A petition for exemption from the electronic prescription mandate shall include, but not be limited to, all of the following:

a. The name and address of the prescriber, medical group, institution, or pharmacy seeking the exemption. For medical groups and institutions, a list of the names, professional license numbers, and CSA registration numbers of all prescribers who would be covered by the exemption shall be maintained by the petitioner for the duration of any approved exemption and shall not be required to be submitted with the petition.

b. Whether the petitioner is seeking an exemption for controlled substance prescriptions, non-controlled substance prescriptions, or both.

c. The petitioner's current electronic prescribing capabilities.

d. The reason, such as economic hardship, technological limitations, or other exceptional circumstances, the petitioner is seeking exemption, including any supporting documentation to justify the reason.

e. Anticipated date of compliance with the electronic prescription mandate.

f. If the petition seeks renewal of a previously approved exemption, information relating to the petitioner's actions during the previous exemption period to work toward compliance with the electronic prescription mandate or an explanation as to why no progress has been made.

21.9(2) Criteria for board consideration of a petition. The board shall consider all information provided in a petition seeking exemption to the electronic prescription mandate and shall approve or deny a petition for exemption based on whether there is a compelling reason to justify the exemption and the nature and volume of prescriptions impacted. Except for petitions citing the exceptional circumstances listed below, which will be administratively reviewed for approval, each petition will be reviewed on a case-by-case basis.

a. A free or low-income clinic where health care is provided at no cost or at a reduced cost to the patient without reimbursement from a third-party payer that requests an exemption for noncontrolled substances only.

b. A licensed prescriber who issues no more than 50 noncontrolled substance prescriptions per year who requests an exemption for noncontrolled substances only.

c. The department of veterans affairs for prescriptions that are not filled at a veterans affairs pharmacy.

d. A prescriber at a student health center based at a college or university for noncontrolled substances only.

e. A dentist seeking an exemption for prescriptions limited to toothpastes and mouthwashes.

f. A compounding pharmacy that dispenses no more than 50 prescriptions for commercially available prescription medications per year that requests an exemption for noncontrolled substances only.

21.9(3) *Duration of approved exemption.* The board may approve an exemption, or the renewal of an exemption, to the electronic prescription mandate for a specified period of time not to exceed one year from the date of approval.

[ARC 4580C, IAB 7/31/19, effective 9/4/19; ARC 6077C, IAB 12/15/21, effective 1/19/22]

657—21.10(124,155A) Automated medication distribution system (AMDS). Any pharmacy that utilizes an AMDS shall comply with these rules in addition to all applicable federal and state laws, rules, and regulations.

21.10(1) *Policies and procedures.* Pursuant to the requirements regarding policies and procedures in 657—subrule 8.3(5), each pharmacy utilizing an AMDS shall have policies and procedures that address all aspects of the operation of the AMDS to include, at a minimum:

- a.* Access to drugs and patient information,
- b.* Pharmacy personnel training in the proper operation of the AMDS,
- c.* Methods to ensure accurate stocking of the AMDS pursuant to subrule 21.10(2),
- d.* Confidentiality of patient information,
- e.* Routine and preventative maintenance of the AMDS according to manufacturer recommendations,
- f.* Packaging and labeling of prescription drugs loaded into or dispensed from the AMDS that is in compliance with federal and state laws, rules, and regulations, and
- g.* Security and control of the prescription drugs maintained and utilized in the AMDS to include:
 - (1) Drug loading, storage, and records.
 - (2) Drugs removed from system components but not used.
 - (3) Inventory.
 - (4) Cross contamination.
 - (5) Lot number control.
 - (6) Wasted or discarded drugs.
 - (7) Controlled substances.

21.10(2) *Stocking the AMDS.* The pharmacy shall have adequate procedures in place to ensure the accurate stocking of drugs into an AMDS using barcode scanning technology. Only a pharmacy technician, pharmacist-intern, or pharmacist shall be allowed to participate in the stocking of the AMDS.

21.10(3) *Pharmacist verification of drugs dispensed from AMDS.*

a. When an AMDS only dispenses drugs that were prepackaged and verified by a pharmacist prior to being stocked in the AMDS and there was no further manipulation of the drug or package other than affixing a patient-specific label, such drugs shall not require additional pharmacist verification prior to administration or dispensing to the patient or authorized representative.

b. When a drug is stocked in an AMDS and undergoes further manipulation, such as counting and packaging, such drugs shall require pharmacist verification prior to dispensing to the patient. Such verification shall be documented.

21.10(4) *Placement of AMDS.*

a. An AMDS placed outside a pharmacist's direct supervision shall only dispense pharmacist-verified packages in compliance with paragraph 21.10(3) "a."

b. An AMDS that manipulates, including but not limited to counting, packaging, or labeling, prescription drugs for subsequent patient dispensing shall only be utilized in a pharmacy under the direct supervision of a pharmacist, except in an approved telepharmacy pursuant to 657—Chapter 13.

[ARC 3640C, IAB 2/14/18, effective 3/21/18]

657—21.11(124,155A) Pharmacist verification of controlled substance fills—daily printout or logbook. The individual pharmacist who makes use of the pharmacy prescription application shall provide documentation of the fact that the fill information entered into the pharmacy prescription application each time the pharmacist fills a prescription order for a controlled substance is correct. If

the pharmacy prescription application provides a hard-copy printout of each day's controlled substance prescription order fill data, that printout shall be verified, dated, and signed by each individual pharmacist who filled a controlled substance prescription order. Each individual pharmacist must verify that the data indicated is correct and sign this document in the same manner as the pharmacist would sign a check or legal document (e.g., J. H. Smith or John H. Smith). This document shall be maintained in a separate file at that pharmacy for a period of two years from the dispensing date. This printout of the day's controlled substance prescription order fill data shall be generated by and available at each pharmacy using a computerized pharmacy prescription application within 48 hours of the date on which the prescription was dispensed. The printout shall be verified and signed by each pharmacist involved with such dispensing. In lieu of preparing and maintaining printouts as provided above, the pharmacy may maintain a bound logbook or separate file. The logbook or file shall include a statement signed each day by each individual pharmacist involved in each day's dispensing that attests to the fact that the prescription information entered into the pharmacy prescription application that day has been reviewed by the pharmacist and is correct as shown. Pharmacist statements shall be signed in the manner previously described. The logbook or file shall be maintained at the pharmacy for a period of two years after the date of dispensing.

[ARC 3640C, IAB 2/14/18, effective 3/21/18]

These rules are intended to implement Iowa Code sections 124.301, 124.306, 124.308, 147.107, 155A.27, 155A.33, and 155A.35.

[Filed 9/16/97, Notice 7/16/97—published 10/8/97, effective 11/12/97]

[Filed 7/31/98, Notice 5/20/98—published 8/26/98, effective 9/30/98]

[Filed 8/14/02, Notice 6/12/02—published 9/4/02, effective 10/9/02]

[Filed 10/22/04, Notice 3/31/04—published 11/10/04, effective 12/15/04]

[Filed 6/2/05, Notice 1/19/05—published 6/22/05, effective 7/27/05]

[Filed 6/2/05, Notice 3/16/05—published 6/22/05, effective 7/27/05]

[Filed 2/7/07, Notice 10/25/06—published 2/28/07, effective 4/4/07]

[Filed 8/3/07, Notice 6/20/07—published 8/29/07, effective 10/3/07]

[Filed ARC 7636B (Notice ARC 7448B, IAB 12/31/08), IAB 3/11/09, effective 4/15/09]

[Filed ARC 8171B (Notice ARC 7910B, IAB 7/1/09), IAB 9/23/09, effective 10/28/09]

[Filed ARC 9912B (Notice ARC 9671B, IAB 8/10/11), IAB 12/14/11, effective 1/18/12]

[Filed ARC 2639C (Notice ARC 2498C, IAB 4/13/16), IAB 8/3/16, effective 9/7/16]

[Filed ARC 3345C (Notice ARC 3136C, IAB 6/21/17), IAB 9/27/17, effective 11/1/17]

[Filed ARC 3640C (Notice ARC 3329C, IAB 9/27/17), IAB 2/14/18, effective 3/21/18]

[Filed ARC 4580C (Notice ARC 4386C, IAB 4/10/19), IAB 7/31/19, effective 9/4/19]

[Filed ARC 5007C (Notice ARC 4695C, IAB 10/9/19), IAB 3/25/20, effective 4/29/20]

[Filed ARC 5350C (Notice ARC 5115C, IAB 7/29/20), IAB 12/30/20, effective 2/3/21]

[Filed ARC 6077C (Notice ARC 5836C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]

[Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

CHAPTER 39
EXPANDED PRACTICE STANDARDS

657—39.1(155A) Purpose and scope. The purpose of this chapter is to establish the minimum standards for the programs and activities identified in this chapter. These rules shall apply to all licensed pharmacists, other registered pharmacy personnel, and all pharmacies, including owners, engaged in the state of Iowa in the programs and activities identified in this chapter. These rules are in addition to rules of the board relating to the practice of pharmacy unless otherwise indicated by rule.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—39.2 and 39.3 Reserved.

657—39.4(155A) Pharmaceutical care. Pharmaceutical care is a comprehensive, patient-centered, outcomes-oriented pharmacy practice in which the pharmacist accepts responsibility for assisting the prescriber and the patient in optimizing the patient's drug therapy plan and works to promote health, to prevent disease, and to optimize drug therapy. Pharmaceutical care does not include the prescribing of drugs without the consent of the prescriber.

39.4(1) Drug therapy problems. In providing pharmaceutical care, the pharmacist shall strive to identify, resolve, and prevent drug therapy problems.

39.4(2) Drug therapy plan. In providing pharmaceutical care, the pharmacist shall access and evaluate patient-specific information, identify drug therapy problems, and utilize that information in a documented plan of therapy that assists the patient or the patient's caregiver in achieving optimal drug therapy. In concert with the patient, the patient's prescribing practitioner, and the patient's other health care providers, the pharmacist shall assess, monitor, and suggest modifications of the drug therapy plan as appropriate.

[ARC 3858C, IAB 6/20/18, effective 7/25/18]

657—39.5 Reserved.

657—39.6(155A) Statewide protocols. To the extent authorized in Iowa Code section 155A.46, a pharmacist may, pursuant to statewide protocols developed by the board in consultation with the department of public health and available on the board's website at pharmacy.iowa.gov, order and dispense medications pursuant to the requirements identified in the statewide protocols. For the purpose of this rule, the order shall constitute a prescription.

[ARC 4270C, IAB 1/30/19, effective 3/6/19; see Delay note at end of chapter; ARC 4387C, IAB 4/10/19, effective 4/5/19; ARC 4583C, IAB 7/31/19, effective 9/4/19; ARC 6076C, IAB 12/15/21, effective 1/19/22]

657—39.7(135,147A) Opioid antagonist dispensing by pharmacist—standing order. An authorized pharmacist may dispense an opioid antagonist pursuant to a standing order established by the department, which standing order can be found via the board's website, or pursuant to a standing order authorized by an individual licensed health care professional in compliance with the requirements of this rule. An authorized pharmacist may only delegate the dispensing of an opioid antagonist to an authorized pharmacist-intern under the direct supervision of an authorized pharmacist. Nothing in this rule prohibits a prescriber or facility from establishing and implementing standing orders or protocols under the authority granted to the prescriber or facility.

39.7(1) Definitions. For the purposes of this rule, the following definitions shall apply:

“Authorized pharmacist” means an Iowa-licensed pharmacist who has completed the training requirements of this rule. *“Authorized pharmacist”* also includes an Iowa-registered pharmacist-intern who has completed the training requirements of this rule and is working under the direct supervision of an authorized Iowa-licensed pharmacist.

“Department” means the Iowa department of public health.

“First responder” means an emergency medical care provider, a registered nurse staffing an authorized service program under Iowa Code section 147A.12, a physician assistant staffing an

authorized service program under Iowa Code section 147A.13, a firefighter, or a peace officer as defined in Iowa Code section 801.4 who is trained and authorized to administer an opioid antagonist.

“Licensed health care professional” means a person licensed under Iowa Code chapter 148 to practice medicine and surgery or osteopathic medicine and surgery, an advanced registered nurse practitioner licensed under Iowa Code chapter 152 or 152E and registered with the board of nursing, or a physician assistant licensed to practice under the supervision of a physician as authorized in Iowa Code chapters 147 and 148C.

“Opioid antagonist” means the same as defined in Iowa Code section 147A.1.

“Opioid-related overdose” means the same as defined in Iowa Code section 147A.1.

“Person in a position to assist” means a family member, friend, caregiver, health care provider, employee of a substance abuse treatment facility, school employee, or other person who may be in a position to render aid to a person at risk of experiencing an opioid-related overdose.

“Recipient” means an individual at risk of an opioid-related overdose or a person in a position to assist an individual at risk of an opioid-related overdose.

“Standing order” means a preauthorized medication order with specific instructions from the licensed health care professional to dispense a medication under clearly defined circumstances.

39.7(2) Authorized pharmacist training. An authorized pharmacist shall document successful completion of an ACPE-approved continuing education program of at least one-hour duration related to opioid antagonist utilization prior to dispensing opioid antagonists pursuant to a standing order.

39.7(3) Additional supply. Notwithstanding a standing order to the contrary, an authorized pharmacist shall only dispense an opioid antagonist after completing an eligibility assessment and providing training and education to the recipient.

39.7(4) Assessment. An authorized pharmacist shall assess an individual for eligibility to receive an opioid antagonist pursuant to a standing order. In addition to the criteria identified in a standing order, an authorized pharmacist shall also take into consideration the following criteria to determine the eligibility of the recipient to receive and possess an opioid antagonist:

a. The person at risk of an opioid-related overdose for which the opioid antagonist is intended to be administered has no known sensitivity or allergy to naloxone, unless the person at risk is not known to the recipient, including but not limited to a first responder or member of law enforcement.

b. The recipient is oriented to person, place, and time and able to understand and learn the essential components of opioid-related overdose, appropriate response, and opioid antagonist administration.

39.7(5) Recipient training and education. Upon assessment and determination that an individual is eligible to receive and possess an opioid antagonist pursuant to a standing order, an authorized pharmacist shall, prior to dispensing an opioid antagonist pursuant to a standing order, provide training and education to the recipient including, but not limited to, the information identified in this subrule. An authorized pharmacist shall require the recipient to attest that, if the product will be accessible to any other individual for administration, the recipient will make available to such individual all received training and education materials. An authorized pharmacist may provide to the recipient written materials that include, but may not be limited to, the information identified in this subrule, but the written materials shall not be in lieu of direct pharmacist consultation with the recipient.

a. The signs and symptoms of opioid-related overdose as described in the standing order.

b. The importance of calling 911 as soon as possible and the potential need for rescue breathing.

c. The appropriate use and directions for administration of the opioid antagonist to be dispensed pursuant to the standing order.

d. Adverse reactions of the opioid antagonist as well as reactions resulting from opioid withdrawal following administration.

e. The proper storage conditions, including temperature excursions, of the opioid antagonist being dispensed.

f. The expiration date of the opioid antagonist being dispensed and the appropriate disposal of the opioid antagonist upon expiration.

g. The prohibition of the recipient from further distributing the opioid antagonist to another individual, unless that individual has received appropriate training and education.

h. Information about substance abuse or behavioral health treatment programs.

39.7(6) Labeling. Upon the determination that a recipient is eligible to receive and possess an opioid antagonist, an authorized pharmacist shall label the product pursuant to rule 657—6.10(126,155A) and 657—subrule 8.19(8). An authorized pharmacist shall ensure that the labeling does not render the expiration date of the product illegible. The medication shall be dispensed in the name of the eligible recipient.

39.7(7) Reporting. A copy of the assessment form shall be submitted to the department as provided on the assessment form within seven days of the dispensing of the opioid antagonist or within seven days of a denial of eligibility.

39.7(8) Records. An authorized pharmacist shall create and maintain an original record of each individual assessment on forms provided by the board, regardless of the eligibility determination following assessment, and dispensing of opioid antagonists pursuant to a standing order. These records shall be available for inspection and copying by the board or its authorized agent for at least two years. [ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 6953C, IAB 3/22/23, effective 4/26/23]

657—39.8(155A) Medications administered via prescription.

39.8(1) Vaccine administration. A pharmacist who is authorized to administer vaccines pursuant to the statewide protocol may administer, including via delegation to authorized pharmacy personnel, any vaccine pursuant to a prescription or medication order for an individual patient. In case of a serious complication, the pharmacist shall notify the prescriber who authorized the prescription within 24 hours and shall submit a report to the Vaccine Adverse Event Reporting System (VAERS).

39.8(2) Medication administration. A pharmacist may administer, including via delegation to authorized pharmacy personnel if so delegated or authorized by the prescriber, any medication pursuant to a prescription or medication order for an individual patient. In case of a serious complication, the pharmacist shall notify the prescriber who issued the prescription within 24 hours and shall submit a report to the United States Food and Drug Administration Adverse Event Reporting System (FAERS). [ARC 6076C, IAB 12/15/21, effective 1/19/22]

657—39.9(155A) Statewide protocol—nicotine replacement tobacco cessation products. Rescinded ARC 6076C, IAB 12/15/21, effective 1/19/22.

657—39.10(155A) Vaccine administration by pharmacists—physician-approved protocol. Rescinded ARC 6076C, IAB 12/15/21, effective 1/19/22.

657—39.11(155A) Vaccine administration by pharmacists—statewide protocol. Rescinded ARC 6076C, IAB 12/15/21, effective 1/19/22.

657—39.12 Reserved.

657—39.13(155A) Collaborative pharmacy practice.

39.13(1) Definitions. For the purpose of this rule, the following definitions shall apply:

“*Collaborative pharmacy practice*” means a practice of pharmacy whereby one or more pharmacists provides patient care and drug therapy management services not otherwise permitted to be performed by a pharmacist to patients under a collaborative pharmacy practice agreement with one or more practitioners which defines the nature, scope, conditions, and limitations of the patient care and drug therapy management services to be provided by the pharmacist(s) in order to ensure that a patient achieves the desired outcomes.

“*Practitioner*” means a physician, dentist, podiatric physician, veterinarian, optometrist, or advanced registered nurse practitioner who holds an active license to practice in Iowa.

39.13(2) Collaborative practice agreement.

a. Pursuant to these rules, a pharmacist or pharmacy may engage in collaborative pharmacy practice under a collaborative pharmacy practice agreement with one or more practitioners, or as

established by a health system pharmacy and therapeutics committee, to provide patient care and drug therapy management services to one or more patients.

b. A collaborative pharmacy practice agreement shall include:

(1) The identification of the parties to the agreement, including the name(s) or category of the pharmacist(s), including registered pharmacist-intern(s) under the supervision of a pharmacist, who are authorized to perform delegated activities under the agreement and the name(s) or category of the practitioner(s) who are delegating activities under the agreement;

(2) The establishment of the delegating practitioner's scope of practice authorized in the agreement and a description of the permitted activities and decisions to be performed by the pharmacist(s);

(3) The protocol, formulary, or clinical guidelines that describe or limit the pharmacist's authority to perform the patient care or drug therapy management services and, as applicable, the drug name, class or category provided under drug therapy management;

(4) A description of the process to monitor compliance with the agreement and clinical outcomes of patients;

(5) The effective date;

(6) A provision addressing termination of the agreement; and

(7) The signatures of the parties to the agreement and dates of signing, unless established by a health system pharmacy and therapeutics committee.

c. Parties to the collaborative pharmacy practice agreement shall review and revise such agreement as appropriate, but no less than every two years.

d. Any collaborative pharmacy practice agreement shall be maintained by the pharmacist(s) or pharmacy and be available upon request or inspection.

e. Prior to engaging in patient care or drug therapy management services under a collaborative pharmacy practice agreement, including when the agreement is updated, each pharmacist practicing under the agreement shall attest that the pharmacist has read and understands the agreement. Documentation of pharmacist attestation shall be maintained for at least two years from the attestation date and be available upon request or inspection.

[ARC 6174C, IAB 2/9/22, effective 3/16/22]

657—39.14 and 39.15 Reserved.

657—39.16(155A) Pharmacy pilot or demonstration research projects. The purpose of this rule is to specify the procedures to be followed in applying for approval of a pilot or demonstration research project for innovative applications in the practice of pharmacy. In reviewing projects, the board will consider only projects that expand pharmaceutical care services that contribute to positive patient outcomes. The board will not consider any project intended only to provide a competitive advantage to a single applicant or group of applicants.

39.16(1) Definitions. For the purposes of this rule, the following definitions shall apply:

“Act” means Iowa Code chapter 155A, the Iowa pharmacy practice Act.

“Board” means the Iowa board of pharmacy.

“Practice of pharmacy” means the practice of pharmacy as defined in Iowa Code section 155A.3(37).

“Project” means a pilot or demonstration research project as described in this rule.

39.16(2) Scope of project. A project may not expand the definition of the practice of pharmacy. A project may include therapeutic substitution or substitution of medical devices used in patient care if such substitution is included under a collaborative pharmacy practice agreement pursuant to rule 657—39.13(155A).

39.16(3) Board approval of a project. Board approval of a project may include the grant of an exception to or a waiver of rules adopted under the Act or under any law relating to the authority of prescription verification and the ability of a pharmacist to provide enhanced patient care in the practice of pharmacy. Project approval, including exception to or waiver of board rules, shall initially be for a specified period of time not exceeding 18 months from commencement of the project. The board may

approve the extension or renewal of a project following consideration of a petition that clearly identifies the project, that includes a report similar to the final project report described in paragraph 39.16(6) "a," that describes and explains any proposed changes to the originally approved and implemented project, and that justifies the need for extending or renewing the term of the project.

39.16(4) *Applying for approval of a project.* A person who wishes the board to consider approval of a project shall submit to the board a petition for approval that contains at least the following information:

a. Responsible pharmacist. Name, address, telephone number, and pharmacist license number of each pharmacist responsible for overseeing the project.

b. Location of project. Name, address, and telephone number of each specific location and, if a location is a pharmacy, the pharmacy license number where the proposed project will be conducted.

c. Project summary. A detailed summary of the proposed project that includes at least the following information:

(1) The goals, hypothesis, and objectives of the proposed project.

(2) A full explanation of the project and how it will be conducted.

(3) The time frame for the project including the proposed start date and length of study. The time frame may not exceed 18 months from the proposed start date of the project.

(4) Background information or literature review to support the proposed project.

(5) The rule or rules to be waived in order to complete the project and a request to waive the rule or rules.

(6) Procedures to be used during the project to ensure that the public health and safety are not compromised as a result of the waiver.

39.16(5) *Review and approval or denial of a proposed project.*

a. Staff review. Upon receipt of a petition for approval of a project, board staff shall initially review the petition for completeness and appropriateness. If the petition is incomplete or inappropriate for board consideration, board staff shall return the petition to the requestor with a letter explaining the reason the petition is being returned. A petition that has been returned pursuant to this paragraph may be amended or supplemented as necessary and submitted for reconsideration.

b. Board review. Upon review by the board of a petition for approval of a project, the board shall either approve or deny the petition. If the board approves the petition, the approval:

(1) Shall be specific for the project requested;

(2) Shall approve the project for a specific time period; and

(3) May include conditions or qualifications applicable to the project.

c. Inspection. The project site and project documentation shall be available for inspection and review by the board or its representative at any time during the project review and the approval or denial processes and, if a project is approved, throughout the approved term of the project.

d. Documentation maintained. Project documentation shall be maintained and available for inspection, review, and copying by the board or its representative for at least two years following completion or termination of the project.

39.16(6) *Presentation of reports.* The pharmacist responsible for overseeing a project shall be responsible for submitting to the board any reports required as a condition of a project, including the final project report.

a. Final project report. The final project report shall include a written summary of the results of the project and the conclusions drawn from those results. The final project report shall be submitted to the board within three months after completion or termination of the project.

b. Board review. The board shall receive and review any report regarding the progress of a project and the final project report at a regularly scheduled meeting of the board. The report shall be an item on the open session agenda for the meeting.

[ARC 3858C, IAB 6/20/18, effective 7/25/18; ARC 6076C, IAB 12/15/21, effective 1/19/22]

These rules are intended to implement Iowa Code sections 135.190, 147.76, 147A.18, 155A.2, 155A.3, 155A.13, 155A.33, and 155A.44; and 2011 Iowa Acts, chapter 63, section 36, as amended by 2012 Iowa Acts, chapter 1113, section 31, and by 2013 Iowa Acts, chapter 138, section 128.

[Filed ARC 3858C (Notice ARC 3509C, IAB 12/20/17), IAB 6/20/18, effective 7/25/18]

[Filed ARC 4270C (Notice ARC 4096C, IAB 10/24/18), IAB 1/30/19, effective 3/6/19]¹
[Filed Emergency ARC 4387C, IAB 4/10/19, effective 4/5/19]
[Filed Emergency After Notice ARC 4555C (Notice ARC 4450C, IAB 5/22/19), IAB 7/17/19,
effective 7/1/19]
[Filed ARC 4583C (Notice ARC 4388C, IAB 4/10/19), IAB 7/31/19, effective 9/4/19]
[Filed ARC 5348C (Notice ARC 5113C, IAB 7/29/20), IAB 12/30/20, effective 2/3/21]
[Filed ARC 6076C (Notice ARC 5833C, IAB 8/11/21), IAB 12/15/21, effective 1/19/22]
[Filed ARC 6174C (Notice ARC 6012C, IAB 11/3/21), IAB 2/9/22, effective 3/16/22]
[Filed ARC 6953C (Notice ARC 6696C, IAB 11/30/22), IAB 3/22/23, effective 4/26/23]

¹ March 6, 2019, effective date of ARC 4270C [amendments to ch 39] delayed 70 days by the Administrative Rules Review Committee at its meeting held February 8, 2019; delay lifted at the meeting held April 5, 2019.

CHAPTER 102
ASSESSMENT PRACTICES AND EQUALIZATION

[Prior to 12/17/86, Revenue Department[730]]
[Prior to 11/2/22, see Revenue Department[701] Ch 71]

701—102.1(405,427A,428,441,499B) Classification of real estate.

102.1(1) Responsibility of assessors. All real estate subject to assessment by city and county assessors shall be classified as provided in this rule. It shall be the responsibility of city and county assessors to determine the proper classification of real estate. There can be only one classification per property under this rule, except as provided for in paragraph 71.1(4)“d.” An assessor shall not assign one classification to the land and a different classification to the building or separate classifications to the land or separate classifications to the building. A building or structure on leased land is considered a separate property and may be classified differently than the land upon which it is located. The determination shall be based upon the best judgment of the assessor following the guidelines set forth in this rule and the status of the real estate as of January 1 of the year in which the assessment is made. The assessor shall classify property according to its present use and not according to its highest and best use. See subrule 71.1(9) for an exception to the general rule that property is to be classified according to its use. The classification shall be utilized on the abstract of assessment submitted to the department of revenue pursuant to Iowa Code section 441.45. See rule 701—71.8(428,441).

102.1(2) Responsibility of boards of review, county auditors, and county treasurers. Whenever local boards of review, county auditors, and county treasurers exercise assessment functions allowed or required by law, they shall classify property as provided in this rule and adhere to the requirements of this rule.

102.1(3) Agricultural real estate.

a. Generally. Agricultural real estate shall include all tracts of land and the improvements and structures located on them which are in good faith used primarily for agricultural purposes except buildings which are primarily used or intended for human habitation as defined in subrule 71.1(4). Land and the nonresidential improvements and structures located on it shall be considered to be used primarily for agricultural purposes if its principal use is devoted to the raising and harvesting of crops or forest or fruit trees, the rearing, feeding, and management of livestock, or horticulture, all for intended profit. Agricultural real estate shall also include woodland, wasteland, and pastureland, but only if that land is held or operated in conjunction with agricultural real estate as defined in paragraph “a” or “b” of this subrule.

b. Vineyards. Beginning with valuations established on or after January 1, 2002, vineyards and any buildings located on a vineyard and used in connection with the vineyard shall be classified as agricultural real estate if the primary use of the land and buildings is an activity related to the production or sale of wine.

c. Algae cultivation and production. Beginning with valuations established on or after January 1, 2013, real estate used directly in the cultivation and production of algae for harvesting as a crop for animal feed, food, nutritionals, or biofuel production shall be classified as agricultural real estate if the real estate is an enclosed pond or land which contains a photobioreactor. Pursuant to 2013 Iowa Acts, House File 632, section 1, a photobioreactor is not attached to land upon which it sits and shall not be assessed and taxed as real property.

(1) Determining direct usage. To determine if real estate is used “directly” in the cultivation and production of algae, one must first ensure that the real estate is used to perform activities that cultivate and produce algae and is not used for activities that occur before or after the cultivation and production of algae. If the real estate is used to perform activities for the cultivation and production of algae, to be “directly” so used, the real estate must be used to perform activities that are integral and essential to the cultivation and production, as distinguished from activities that are incidental, merely convenient to, or remote from cultivation and production. The fact that real estate is used for activities that are essential or necessary to the cultivation and production of algae does not mean that the real estate is also “directly” used in production. Even if the real estate is used for activities that are essential or necessary

to the cultivation and production of algae, if the activities are far enough removed from the cultivation or production of algae, the real estate would not qualify for the agricultural designation.

(2) Examples. The following are nonexclusive examples of real estate which would not be directly used in the cultivation and production of algae:

1. Real estate that is used to store, assemble, or repair machinery and equipment that is used for cultivation and production of algae.
2. Real estate that is used in the management, administration, advertising, or selling of algae.
3. Real estate that is used in the management, administration, or planning of the cultivation and production of algae.
4. Real estate that is used for packaging of the algae which has been produced and cultivated.

102.1(4) Residential real estate.

a. Classification of residential real estate—in general. Residential real estate shall include all lands and buildings which are primarily used or intended for human habitation containing fewer than three dwelling units, including those buildings located on agricultural land. Buildings used primarily or intended for human habitation shall include the dwelling as well as structures and improvements used primarily as a part of, or in conjunction with, the dwelling. This includes but is not limited to garages, whether attached or detached, tennis courts, swimming pools, guest cottages, and storage sheds for household goods. “Used in conjunction with” means that the structure or improvement is located on the same parcel, on contiguous parcels, or on a parcel directly across a street or alley as the building or structure containing the dwelling and when marketed for sale would be sold as a unit. Residential real estate located on agricultural land shall include only buildings as defined in this subrule. Buildings for human habitation that are used as commercial ventures, including but not limited to hotels, motels, rest homes, and structures containing three or more separate living quarters, shall not be considered residential real estate. However, regardless of the number of separate living quarters, multiple housing cooperatives organized under Iowa Code chapter 499A and land and buildings owned and operated by organizations that have received tax-exempt status under Section 501(c)(3) of the Internal Revenue Code, if the rental income from the property is not taxed as unrelated business income under Iowa Code section 422.33(1A), shall be considered residential real estate.

b. Horizontal property regimes. An apartment in a horizontal property regime (condominium) referred to in Iowa Code chapter 499B which is used or intended for use for human habitation shall be classified as residential real estate regardless of who occupies the apartment. Existing structures shall not be converted to a horizontal property regime unless building code requirements have been met.

c. Classification of residential real estate on or after January 1, 2022. Beginning with valuations established on or after January 1, 2022, residential real estate shall also include:

- (1) Property primarily used or intended for human habitation containing two or fewer dwelling units.
- (2) Mobile home parks.
- (3) Manufactured home communities.
- (4) Land-leased communities.
- (5) Assisted living facilities.
- (6) A parcel primarily used or intended for human habitation containing three or more separate dwelling units. If a portion of such a parcel is used or intended for a purpose that, if the primary use would be classified as commercial property or industrial property, each such portion, including a proportionate share of the land included in the parcel, if applicable, shall be assigned the appropriate classification pursuant to Iowa Code section 441.21(14)“b” and paragraph 71.1(4)“d” below.

(7) For a parcel that is primarily used or intended for use as commercial property or industrial property, that portion of the parcel that is used or intended for human habitation, regardless of the number of dwelling units contained on the parcel, including a proportionate share of the land included in the parcel, if applicable. The portion of such a parcel used or intended for use as commercial property or industrial property, including a proportionate share of the land included in the parcel, if applicable, shall be assigned the appropriate classification pursuant to Iowa Code section 441.21(14)“b” and paragraph 71.1(4)“d” below.

d. Dual classification.

(1) For assessment years beginning January 1, 2022, and after, valuations of parcels for which a portion of the parcel satisfies the requirements for classification as residential property under Iowa Code section 441.21(14)“a”(6) or 441.21(14)“a”(7) and subparagraph 71.1(4)“c”(6) or 71.1(4)“c”(7), the assessor shall assign to that portion of the parcel the classification of residential property and to such other portions of the parcel the property classification for which such other portions qualify.

(2) The only permitted combinations of dual classifications are commercial and residential or industrial and residential. The assessor shall assign the classification of residential to that portion of the parcel that satisfies the requirements for the classification of residential property and to such other portions of the parcel the property classification for which such other portions qualify. The assessor shall maintain the valuation and assessment of property with a dual classification on one parcel record.

e. Section 42 housing. Property that is rented or leased to low-income individuals and families as authorized by Section 42 of the Internal Revenue Code, and that has not been withdrawn from Section 42 assessment procedures under Iowa Code section 441.21(2), shall not be classified as residential property.

f. Short-term leases. A hotel, motel, inn, or other building where rooms or dwelling units are usually rented for less than one month shall not be classified as residential property.

g. Definitions. For purposes of this subrule, the following definitions apply:

“*Assisted living facility*” means property for providing assisted living as defined in Iowa Code section 231C.2. “Assisted living facility” also includes a health care facility as defined in Iowa Code section 135C.1, an elder group home as defined in Iowa Code section 231B.1, a child foster care facility under Iowa Code chapter 237, or property used for a hospice program as defined in Iowa Code section 135J.1.

“*Dwelling unit*” means an apartment, group of rooms, or single room which is occupied as separate living quarters or, if vacant, is intended for occupancy as separate living quarters, in which a tenant can live and sleep separately from any other persons in the building.

“*Land-leased community*” means the same as defined in Iowa Code sections 335.30A and 414.28A.

“*Manufactured home community*” means the same as a land-leased community.

“*Mobile home park*” means the same as defined in Iowa Code section 435.1.

102.1(5) Reserved.

102.1(6) *Commercial real estate.* Commercial real estate shall include all lands and improvements and structures located thereon which are primarily used or intended as a place of business where goods, wares, services, or merchandise is stored or offered for sale at wholesale or retail. Commercial realty shall also include hotels, motels, and property that is rented or leased to low-income individuals and families as authorized by Section 42 of the Internal Revenue Code and has not been withdrawn from Section 42 assessment procedures under Iowa Code section 441.21(2). Commercial real estate shall also include data processing equipment as defined in Iowa Code section 427A.1(1)“j,” except data processing equipment used in the manufacturing process. However, regardless of the number of separate living quarters or any commercial use of the property, single- and two-family dwellings, multiple housing cooperatives organized under Iowa Code chapter 499A, and land and buildings used primarily for human habitation and owned and operated by organizations that have received tax-exempt status under Section 501(c)(3) of the Internal Revenue Code, if the rental income from the property is not taxed as unrelated business income under Iowa Code section 422.33(1A), shall be classified as residential real estate.

An apartment in a horizontal property regime (condominium) referred to in Iowa Code chapter 499B which is used or intended for use as a commercial venture, other than leased for human habitation, shall be classified as commercial real estate. Existing structures shall not be converted to a horizontal property regime unless building code requirements have been met.

102.1(7) *Industrial real estate.*

a. Land and buildings.

(1) Industrial real estate includes land, buildings, structures, and improvements used primarily as a manufacturing establishment. A manufacturing establishment is a business entity in which the primary activity consists of adding to the value of personal property by any process of manufacturing, refining, purifying, the packing of meats, or the combination of different materials with the intent of selling the

product for gain or profit. Industrial real estate includes land and buildings used for the storage of raw materials or finished products and which are an integral part of the manufacturing establishment, and also includes office space used as part of a manufacturing establishment.

(2) Whether property is used primarily as a manufacturing establishment and, therefore, assessed as industrial real estate depends upon the extent to which the property is used for the activities enumerated in subparagraph 71.1(7)“a”(1). Property in which the performance of these activities is only incidental to the property’s primary use for another purpose is not a manufacturing establishment. For example, a grocery store in which bakery goods are prepared would be assessed as commercial real estate since the primary use of the grocery store premises is for the sale of goods not manufactured by the grocery and the industrial activity, i.e., baking, is only incidental to the store premises’ primary use. However, property which is used primarily as a bakery would be assessed as industrial real estate even if baked goods are sold at retail on the premises since the bakery premises’ primary use would be for an industrial activity to which the retail sale of baked goods is merely incidental. See *Lichty v. Board of Review of Waterloo*, 230 Iowa 750, 298 N.W. 654 (1941).

Similarly, a facility which has as its primary use the mixing and blending of products to manufacture feed would be assessed as industrial real estate even though a portion of the facility is used solely for the storage of grain, if the use for storage is merely incidental to the property’s primary use as a manufacturing establishment. Conversely, a facility used primarily for the storage of grain would be assessed as commercial real estate even though a part of the facility is used to manufacture feed. In the latter situation, the industrial use of the property — the manufacture of feed — is merely incidental to the property’s primary use for commercial purposes — the storage of grain.

(3) Property used primarily for the extraction of rock or mineral substances from the earth is not a manufacturing establishment if the only processing performed on the substance is to change its size by crushing or pulverizing. See *River Products Company v. Board of Review of Washington County*, 332 N.W.2d 116 (Iowa Ct. App. 1982).

b. Machinery.

(1) Machinery includes equipment and devices, both automated and nonautomated, which is used in manufacturing as defined in Iowa Code section 428.20. See *Deere Manufacturing Co. v. Beiner*, 247 Iowa 1264, 78 N.W.2d 527 (1956).

(2) Machinery owned or used by a manufacturer but not used within the manufacturing establishment is not assessed as industrial real estate. For example, “X” operates a factory which manufactures building materials for sale. In addition, “X” uses some of these building materials in construction contracts. The machinery which “X” would primarily use at the construction site would not be used in a manufacturing establishment and, therefore, would not be assessed as industrial real estate.

(3) Machinery used in manufacturing but not used in or by a manufacturing establishment is not assessed as industrial real estate. See *Associated General Contractors of Iowa v. State Tax Commission*, 255 Iowa 673, 123 N.W.2d 922 (1963).

(4) Where the primary function of a manufacturing establishment is to manufacture personal property that is consumed by the manufacturer rather than sold, the machinery used in the manufacturing establishment is not assessed as industrial real estate. See *Associated General Contractors of Iowa v. State Tax Commission*, 255 Iowa 673, 123 N.W.2d 922 (1963).

102.1(8) Point-of-sale equipment. As used in Iowa Code section 427A.1(1)“j,” the term “point-of-sale equipment” means input, output, and processing equipment used to consummate a sale and to record or process information pertaining to a sale transaction at the time the sale takes place and which is located at the counter, desk, or other specific point at which the transaction occurs. As used in this subrule, the term “sale” means the sale or rental of goods or services and includes both retail and wholesale transactions. Point-of-sale equipment does not include equipment used primarily for depositing or withdrawing funds from financial institution accounts.

102.1(9) Housing development property.

a. Ordinances adopted or amended on or after January 1, 2011.

(1) Adoption of ordinance by board of supervisors. A county board of supervisors may adopt an ordinance providing that property acquired and subdivided for development of housing on or after

January 1, 2011, shall continue to be assessed for taxation in the manner it was assessed prior to the acquisition. Each lot shall continue to be taxed in the manner it was taxed prior to acquisition for housing until the lot is sold for construction or occupancy of housing or 5 years from the date of subdivision, whichever occurs first.

(2) Amendments to ordinance by board of supervisors. On or after July 27, 2011, the board of supervisors of a county may amend an ordinance adopted or otherwise made effective under 2011 Iowa Code Supplement section 405.1(1) "a" to extend the 5-year time period for a period of time not to exceed 5 years beyond the end of the original 5-year period established under 2011 Iowa Code Supplement section 405.1(1). Thus, the maximum special assessment time for ordinances adopted on or subsequent to January 1, 2011, is 10 years. An extension of an ordinance under 2011 Iowa Code Supplement section 405.1(1) "a" may apply to all or a portion of the property that was subject to the original ordinance.

(3) Amendments to ordinance by city council. A city council may adopt an ordinance, affecting all or a portion of the property located within the incorporated area of the city subject to the county ordinance adopted under 2011 Iowa Code Supplement section 405.1(1) "a," extending the county ordinance not previously extended by the board of supervisors up to 5 years. An ordinance by a city council providing for an extension under 2011 Iowa Code Supplement section 405.1(3) shall be subject to the 5-year limitation under 2011 Iowa Code Supplement section 405.1(2). Thus, the maximum time to appeal an ordinance adopted on or subsequent to January 1, 2011, is 10 years if the city council amends an ordinance originally adopted by the county board of supervisors.

(4) Sale of lot; expiration of 5-year or extended period. Upon the sale of the lot for construction or occupancy for housing or upon the expiration of the 5-year or extended period, the property shall be assessed for taxation as residential or commercial multifamily property, whichever is applicable.

(5) Definition of "subdivide." As used in both paragraphs 71.1(9) "a" and "b," "subdivide" means to divide a tract of land into three or more lots.

b. Ordinances adopted on or after January 1, 2004, but prior to January 1, 2011.

(1) Ordinances adopted under 2011 Iowa Code Supplement sections 405.1(1) and 405.1(2), to the extent such ordinances affect the assessment of property subdivided for development of housing on or after January 1, 2004, but before January 1, 2011, shall remain in effect or otherwise be made effective, and such ordinances:

1. Adopted under 2011 Iowa Code Supplement section 405.1(1), applicable to counties with a population of less than 20,000, shall be extended, from a period of 5 years, to apply to a period of 10 years from the date of subdivision.

2. Adopted under 2011 Iowa Code Supplement section 405.1(2), applicable to counties with a population of 20,000 or more, shall be extended, from a period of 3 years, to apply to a period of 8 years from the date of subdivision.

Each lot shall continue to be taxed in the manner it was taxed prior to acquisition for housing until the lot is sold for construction or occupancy of housing, or 10 years pursuant to paragraph "1" above or 8 years pursuant to paragraph "2" above (or the extended period, if applicable) from the date of subdivision, whichever occurs first.

(2) Amendments to ordinance by board of supervisors. On or after July 27, 2011, the board of supervisors of a county may amend an ordinance adopted under 2011 Iowa Code Supplement section 405.1(1) or 405.1(2) to extend the 10- and 8-year periods, respectively, for a period of time not to exceed 5 years beyond the end of the 10- and 8-year periods established under 2011 Iowa Code Supplement section 405.1(1) "b." Thus, the maximum special assessment time for ordinances adopted on or after January 1, 2004, but prior to January 1, 2011, for counties with a population of less than 20,000 shall be 15 years. For counties with a population of 20,000 or more, the maximum shall be 13 years.

(3) Amendments to ordinance by city council. A city council may adopt an ordinance, affecting all or a portion of the property located within the incorporated area of the city subject to the county ordinance adopted under 2011 Iowa Code Supplement sections 405.1(1) and 405.1(2), extending the county ordinances not previously extended by the board of supervisors up to 5 years. An ordinance by a city council providing for an extension under 2011 Iowa Code Supplement section 405.1(3) shall be subject to the 5-year limitation under 2011 Iowa Code Supplement section 405.1(2). Thus, the maximum

time to appeal an ordinance adopted on or after January 1, 2004, but prior to January 1, 2011, for counties with a population of less than 20,000 shall be 15 years if the city council amends an ordinance originally adopted by the board of supervisors. For counties with a population of 20,000 or more, the maximum special assessment time shall be 13 years.

(4) Sale of lot. Upon the sale of the lot for construction or occupancy for housing or upon the expiration of the 10- or 8-year or extended period, the property shall be assessed for taxation as residential or commercial multifamily property, whichever is applicable.

102.1(10) Assessment of platted lots.

a. When a subdivision plat is recorded pursuant to Iowa Code chapter 354 on or after January 1, 2011, the individual lots within the subdivision plat shall not be assessed, in the aggregate, in excess of the total assessment of the land as acreage or unimproved property for 5 years after the recording of the plat or until the lot is actually improved with permanent construction, whichever occurs first. When an individual lot has been improved with permanent construction, the lot shall be assessed for taxation purposes as provided in Iowa Code chapters 428 and 441.

b. For subdivision plats recorded pursuant to Iowa Code chapter 354 (relating to division and subdivision of land) on or after January 1, 2004, but before January 1, 2011, the individual lots within the subdivision plat shall not be assessed, in the aggregate, in excess of the total assessment of the land as acreage or unimproved property for 8 years after the recording of the plat or until the lot is actually improved with permanent construction, whichever occurs first. When an individual lot has been improved with permanent construction, the lot shall be assessed for taxation purposes as provided in Iowa Code chapters 428 and 441.

c. 2011 Iowa Code Supplement section 441.72 does not apply to special assessment levies.

This rule is intended to implement Iowa Code sections 405.1, 427A.1, 428.4 and 441.22 and chapter 499B and Iowa Code Supplement section 441.21 as amended by 2002 Iowa Acts, House File 2584.

[ARC 8559B, IAB 3/10/10, effective 4/14/10; ARC 0400C, IAB 10/17/12, effective 11/21/12; ARC 1196C, IAB 11/27/13, effective 1/1/14; ARC 1765C, IAB 12/10/14, effective 1/14/15; ARC 2146C, IAB 9/16/15, effective 10/21/15; ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.2(421,428,441) Assessment and valuation of real estate.

102.2(1) Responsibility of assessor. The valuation of real estate as established by city and county assessors shall be the actual value of the real estate as of January 1 of the year in which the assessment is made. New parcels of real estate created by the division of existing parcels of real estate shall be assessed separately as of January 1 of the year following the division of the existing parcel of real estate.

102.2(2) Responsibility of other assessing officials. Whenever local boards of review, county auditors, and county treasurers exercise assessment functions allowed or required by law, they shall follow the provisions of subrule 71.2(1) and rules 701—71.3(421,428,441) to 701—71.7(421,427A,428,441).

This rule is intended to implement Iowa Code sections 421.17, 428.4 and 441.21.

[Editorial change: IAC Supplement 11/2/22]

701—102.3(421,428,441) Valuation of agricultural real estate. Agricultural real estate shall be assessed at its actual value as defined in Iowa Code section 441.21 by giving exclusive consideration to its productivity and net earning capacity. In determining the actual value of agricultural real estate, city and county assessors shall use the Iowa Real Property Appraisal Manual and any other guidelines issued by the department of revenue pursuant to Iowa Code section 421.17(17).

102.3(1) Productivity.

a. In determining the productivity and net earning capacity of agricultural real estate, the assessor shall also use available data from Iowa State University, the United States Department of Agriculture (USDA) National Agricultural Statistics Service (NASS), the USDA Farm Service Agency (FSA), the Iowa department of revenue, or other reliable sources. The assessor shall also consider the results of a modern soil survey, if completed. The assessor shall determine the actual valuation of agricultural real estate within the assessing jurisdiction and distribute such valuation throughout the jurisdiction so that each parcel of real estate is assessed at its actual value as defined in Iowa Code section 441.21.

b. In distributing such valuation to each parcel under paragraph 71.3(1) “*a*,” the assessor shall adjust non-cropland. The adjustment shall be applied to non-cropland with a corn suitability rating (CSR) that is greater than 50 percent of the average CSR for cropland for the county. The adjustment shall be determined for each county based upon the five-year average difference in cash rent between non-irrigated cropland and pasture land as published by NASS. The assessor may utilize the USDA FSA-published Common Land Unit digital data or other reliable sources in determining non-cropland. Counties shall implement the adjustments under this paragraph on or before the 2017 assessment year. The department of revenue may, in a case involving hardship, extend the implementation of the adjustments required under this paragraph to the 2019 assessment year. No extension of time shall be granted unless the county makes a written request to the department of revenue for such action.

c. A taxpayer may apply to the county for the adjustment to non-cropland under paragraph 71.3(1) “*b*” beginning with the 2014 assessment and until the county’s full implementation of this subrule. Upon application, and subsequent approval by the assessor, the county assessor shall adjust non-cropland as provided in paragraph 71.3(1) “*b*.” Once a taxpayer applies for the adjustment, and upon approval, the assessor shall make the adjustment to the assessment year for which the application was submitted and until the county’s full implementation of this subrule, without the need to reapply for the adjustment.

d. EXAMPLE. The following is an example of the calculation used to compute adjustment on land determined to be non-cropland with a CSR that is greater than 50 percent of the average CSR for cropland for the county:

Average county CSR rating for cropland	80 CSR
50% of average cropland CSR	40 CSR
Example of non-cropland soil 11b CSR rating	58 CSR
Non-cropland CSR points to be adjusted	$58 - 40 = 18$ CSR points
5-year average rent for non-irrigated cropland	\$163.60
5-year average rent for pasture land	\$48.30
Percent difference (rounded)	$1 - (\$48.30/\$163.60) = 70\%$
Apply the percent difference to points to be adjusted	$18 \text{ CSR points} \times (1 - .70) = 5.40$ adjusted CSR points
Adjusted CSR non-cropland	$40 + 5.40 = 45.40$ adjusted CSR points

102.3(2) Agricultural factor. In order to determine a productivity value for agricultural buildings and structures, assessors must make an agricultural adjustment to the market value of these buildings and structures by developing an “agricultural factor” for the assessors’ jurisdictions. The agricultural factor for each jurisdiction is the product of the ratio of the productivity and net earning capacity value per acre as determined under subrule 71.12(1) over the market value of agricultural land within the assessing jurisdiction. The resulting ratio is then applied to the actual value of the agricultural buildings and structures as determined under the Iowa Real Property Appraisal Manual prepared by the department. The agricultural factor must be applied uniformly to all agricultural buildings and structures in the assessing jurisdiction. As an example, if a building’s actual value is \$500,000 and the agricultural factor is 30 percent, the productivity value of that building is \$150,000. See *H & R Partnership v. Davis County Board of Review*, 654 N.W.2d 521 (Iowa 2002). The 2007, 2008, and 2009 average of the market value of land will be used in determining the agricultural factor for assessment year 2011. A five-year market value average of land for years used to determine the productivity formula will be used to determine the agricultural factor for assessment year 2013 and subsequent assessment years.

102.3(3) Classification. Land classified as agricultural real estate includes the land beneath any dwelling and appurtenant structures located on that land and shall be valued by the assessor pursuant to rule 701—71.3(421,428,441). An assessor shall not value a part of the land as agricultural real estate and a part of the land as if it is residential real estate.

This rule is intended to implement Iowa Code sections 421.17, 428.4 and 441.21.
 [ARC 8542B, IAB 2/24/10, effective 3/31/10; ARC 9478B, IAB 4/20/11, effective 5/25/11; ARC 0770C, IAB 5/29/13, effective 7/3/13; ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.4(421,428,441) Valuation of residential real estate. Residential real estate shall be assessed at its actual value as defined in Iowa Code section 441.21.

In determining the actual value of residential real estate, city and county assessors shall use the appraisal manual issued by the department of revenue pursuant to Iowa Code section 421.17(17) as well as a locally conducted assessment/sales ratio study, an analysis of sales of comparable properties, and any other relevant data available.

This rule is intended to implement Iowa Code sections 421.17, 428.4 and 441.21.
[ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.5(421,428,441) Valuation of commercial real estate. Commercial real estate shall be assessed at its actual value as defined in Iowa Code section 441.21. In determining the actual value of commercial real estate, city and county assessors shall use the appraisal manual issued by the department of revenue pursuant to Iowa Code section 421.17(17) as well as a locally conducted assessment/sales ratio study, an analysis of sales of comparable properties, and any other relevant data available. In cases involving the valuation of owner-occupied commercial property, the data relating to the financial performance of the owner or the owner's business, including but not limited to its sales, revenue, expenses, or profits, shall not be considered relevant in determining the property's actual value.

102.5(1) Property of long distance telephone companies. The director of revenue shall assess the property of long distance telephone companies as defined in Iowa Code section 476.1D(10) which property is first assessed for taxation on or after January 1, 1996, in the same manner as commercial real estate.

102.5(2) Low-income housing subject to Section 42 of the Internal Revenue Code.

a. Productive and earning capacity. In assessing property that is rented or leased to low-income individuals and families as authorized by Section 42 of the Internal Revenue Code which limits the amount that the individual or family pays for the rental or lease of units in the property, the assessor shall use the productive and earning capacity from the actual rents received as a method of appraisal and shall take into account the extent to which that use and limitation reduces the market value of the property.

b. Direct capitalization method. The income approach to valuation shall be applied using the direct capitalization method. The assessor may use the discounted cash flow method as a test of the reasonableness of the results produced by the direct capitalization method. The direct capitalization method of the income approach involves dividing the Net Operating Income (NOI) on a cash basis by an overall capitalization rate to derive an indication of the value of the property for the assessment year.

In applying the direct capitalization method, the assessor shall develop a normalized measure of annual NOI based on the productive and earning capacity of the development utilizing (1) the actual rent schedule applicable for each of the available units as of January 1 of the year of assessment indicating the actual rent to be paid by the resident plus any Section 8 rental assistance or other direct cash rental subsidy provided to the resident by federal, state or local rent subsidy programs as limited pursuant to Section 42 of the Internal Revenue Code, (2) a normal vacancy/collection allowance, (3) the prior year's actual and current year's projected annual operating expenses associated with the property, excluding noncash items such as depreciation and amortization, but including property taxes and those actual costs expected to be incurred and paid as required by Internal Revenue Code Section 42 regulations, provisions, and restrictions as applicable to the assessment year, and (4) an appropriate provision for replacement reserves.

If no separate line item is included for reserves for replacement in the historic income and expense data, then the maintenance and repair categories of the historic expense data must be itemized. For properties that have attained a normalized operating history, the NOI results of the prior three years (as represented in the statements variously named as the Income and Loss Statement, the Profit and Loss Statement, the Income Statement, the Actual to Budget Comparison Statement, Balance Sheet, or some name variation of these) may be used to provide the basis for determining the normalized NOI used for purposes of applying the direct capitalization method for the year of assessment, provided an appropriate replacement reserve is included in the NOI determination and provided any additional costs required as a result of Section 42 regulation or compliance changes for the assessment year are included

as an operating expense in the NOI determination. In addition, the assessor may utilize the current year operating budget to develop a measure of NOI for the assessment year. The assessor, in developing the measure of annual NOI on a cash basis, shall not consider as income any potential rental income differential that could otherwise be received from the property if the rents were not limited pursuant to Section 42 of the Internal Revenue Code, any tax credit equity, any tax credit value, or other subsidized financing.

c. *Filing of reports.* It shall be the responsibility of the property owner to file income and expense data with the local assessor by March 1 of each year. The assessor may require the filing of additional information if deemed necessary.

d. *Capitalization rate.* The overall capitalization rate to be used in applying the direct capitalization method for a Section 42 property is developed through the band-of-investment technique. The capitalization rate will be calculated annually by the Iowa department of revenue and distributed to all Iowa assessors by March 1. The capitalization rate is a composite rate weighted by the proportions of total property investment represented by debt and equity. The capital structure weights equity at 80 percent and debt at 20 percent unless actual market capital structure can be verified to the assessor. The yield, or market rate of return, for equity is calculated using the capital asset pricing model (CAPM). The yield for debt is equivalent to the average yield on 25-year Treasury bonds referred to as the Treasury long-term average rate. An example of the band-of-investment technique to be utilized is as follows:

	<u>% to Total</u>	<u>Yield</u>	<u>Composite</u>
Equity	80%	11.05%	8.84%
Debt	20%	5.94%	1.19%
	<u>100%</u>		<u>10.03%</u>

e. *Capital asset pricing model.* The capital asset pricing model (CAPM) is utilized to develop the equity rate. The formula is:

$$Re = B (Rm - Rf) + Rf$$

Where:

- Re = return on equity
- B = beta
- Rm = return on the market
- Rf = risk-free rate of return
- Rm - Rf = market-risk premium

The beta is assumed to be 1 which indicates the risk level to be consistent with the market as a whole. The risk-free rate is calculated by finding the average of the three-month and six-month Treasury bill. The return on the market is calculated by taking the average of the return on the market for the Merrill Lynch Universe and Standard and Poor’s 500 or by reference to other published secondary sources.

f. *Properties under construction.* For Section 42 properties under construction, the assessor may value the property by applying the percentage of completion to the replacement cost new (RCN) as calculated from the Iowa Real Property Appraisal Manual and adding the fair market value of the land. Alternatively, projected income and expense data may be utilized if available.

g. *Negative or minimal NOI.* If the Section 42 property shows a negative or minimal net operating income (NOI), the indicator of value as set forth in these rules shall not be utilized.

h. Eligibility withdrawn. The property owner shall notify the assessor when property is withdrawn from Section 42 eligibility under the Internal Revenue Code. The notification must be provided by March 1 of the assessment year or the owner is subject to a penalty of \$500.

This rule is intended to implement Iowa Code sections 421.17, 428.4, 441.21 as amended by 2004 Iowa Acts, Senate File 2296, and 476.1D(10).

[ARC 3107C, IAB 6/7/17, effective 7/12/17; ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.6(421,428,441) Valuation of industrial land and buildings. Industrial real estate shall be assessed at its actual value as defined in Iowa Code section 441.21.

In determining the actual value of industrial land and buildings, city and county assessors shall use the appraisal manual issued by the department of revenue pursuant to Iowa Code section 421.17(17), and any other relevant data available.

This rule is intended to implement Iowa Code sections 421.17, 428.4 and 441.21.

[ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.7(421,427A,428,441) Valuation of industrial machinery. Industrial machinery as referred to in Iowa Code section 427A.1(1) “e” shall include all machinery used in manufacturing establishments and shall be assessed as real estate even though such machinery might be assessed as personal property if not used in a manufacturing establishment.

In determining the actual value of industrial machinery assessed as real estate, the assessor shall give consideration to the “Industrial Machinery and Equipment Valuation Guide” issued by the department of revenue and any other relevant data available.

This rule is intended to implement Iowa Code sections 421.17, 427A.1, 428.4 and 441.21.

[Editorial change: IAC Supplement 11/2/22]

701—102.8(428,441) Abstract of assessment. Each city and county assessor shall submit annually to the department of revenue at the times specified in Iowa Code section 441.45 an abstract of assessment for the current year. The assessor shall use the form of abstract prescribed and furnished by the department and shall enter on the abstract all information required by the department. However, the department may approve the use of a computer-prepared abstract if the data is in essentially the same format as on the form prescribed by the department. The information entered on the abstract of assessment shall be reviewed and considered by the department in equalizing the valuations of classes of properties.

This rule is intended to implement Iowa Code sections 428.4 and 441.45.

[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.9(428,441) Reconciliation report. The assessor’s report of any revaluation required by Iowa Code section 428.4 shall be made on the reconciliation report prescribed and furnished by the department of revenue. The assessor shall enter on the report all information required by the department. The reconciliation report shall be a part of the abstract of assessment required by Iowa Code section 441.45 and shall be reviewed and considered by the department in equalizing valuations of classes of property.

This rule is intended to implement Iowa Code sections 428.4 and 441.45.

[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.10(421) Assessment/sales ratio study.

102.10(1) Basic data. Basic data shall be that submitted to the department of revenue by county recorders and city and county assessors on forms prescribed and provided by the department, information furnished by parties to real estate transactions, and information obtained by field investigations made by the department of revenue.

102.10(2) Responsibility of recorders and assessors. County recorders and city and county assessors shall complete the prescribed forms as required by Iowa Code subsection 421.17(6) and rule 701—79.3(428A) in accordance with instructions issued by the department. Assessed values entered on the prescribed form shall be those established as of January 1 of the year in which the sale takes place.

102.10(3) Normal sales. All real estate transfers shall be considered by the department of revenue to be normal sales unless there exists definite information which would indicate the transfer was not an arms-length transaction or is of an excludable nature as provided in Iowa Code section 441.21.

This rule is intended to implement Iowa Code section 421.17.

[Editorial change: IAC Supplement 11/2/22]

701—102.11(441) Equalization of assessments by class of property.

102.11(1) Commencing in 1977 and every two years thereafter, the department of revenue shall order the equalization of the levels of assessment of each class of property as provided in rule 701—71.12(441) by adding to or deducting from the valuation of each class of property, as reported to the department on the abstract of assessment and reconciliation report that is a part of the abstract, the percentage in each case as may be necessary to bring the level of assessment to its actual value as defined in Iowa Code section 441.21. Valuation adjustments shall be ordered if the department determines that the aggregate valuation of a class of property as reported on the abstract of assessment submitted by the assessor is at least 5 percent above or below the aggregate valuation for that class of property as determined by the department pursuant to rule 701—71.12(441). Equalization orders of the department shall be restricted to equalizing the aggregate valuations of entire classes of property among the several assessing jurisdictions. All classifications of real estate shall be applied uniformly throughout the state of Iowa.

102.11(2) Equalization percentage adjustments determined for residential realty located outside incorporated areas and not located on agricultural land shall apply to buildings located on agricultural land outside incorporated areas, which are primarily used or intended for human habitation, as defined in subrule 71.1(4).

Equalization percentage adjustments determined for residential realty located within incorporated cities and not located on agricultural land shall apply to buildings located on agricultural land within incorporated cities that are primarily used or intended for human habitation as defined in subrule 71.1(4).

This rule is intended to implement Iowa Code sections 441.21, 441.47, 441.48 and 441.49.

[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.12(441) Determination of aggregate actual values.

102.12(1) Agricultural real estate.

a. Use of income capitalization study. The equalized valuation of agricultural realty shall be based upon its productivity and net earning capacity and shall be determined in accordance with the provisions of this subrule. Data used shall pertain to crops harvested during the five-year period ending with the calendar year in which assessments were last equalized. The equalized valuation of agricultural realty shall be determined for each county as follows:

(1) Computation of county acres. This information shall be obtained from the USDA National Agricultural Statistics Service.

1. Total acres in farms: Total acreage used for agricultural purposes.
2. Corn acres: Sum of corn acres harvested including silage, popcorn and acres planted for sorghum.
3. Oats and wheat acres: Sum of oats and wheat acres harvested.
4. Soybean acres: Soybean acres harvested.
5. Hay acres: All hay acres harvested.
6. Pasture acres: All pasture acres. Total pasture acres shall be determined by multiplying the total acres in farms reported by the USDA National Agricultural Statistics Service by the percentage which total pasture land as reported in the most recent U.S. Census of Agriculture bears to the total acreage in farmland also reported in the most recent U.S. Census of Agriculture. The amount of tillable and nontillable pasture acres shall be determined as follows:

1.	From the most recent U.S. Census of Agriculture obtain the following:		
	Cropland used only for pasture and grazing	_____	acres
	Woodland pasture	_____	acres
	Pasture land and rangeland (other than cropland and woodland pasture)	_____	acres
	TOTAL PASTURE LAND (total of above):	_____	acres
2.	Determine what percentage of the total pasture land is cropland used only for pasture:	_____	%
3.	Apply the percentage in "2" above to the 5-year average total acres of pasture as determined above to determine the pasture acres to be classified as tillable pasture. The remainder of the 5-year average shall be classified as nontillable pasture land.	_____	acres

7. Government programs: Determine the 5-year average acres participating in applicable government programs. Obtain data from the USDA Farm Service Agency, including but not limited to acreage devoted to the Payment-In-Kind (PIK), diverted and deficiency programs.

8. Other acres: The difference between the total acreage for land uses listed above and the total of all land in farms. Add the total of the corn, oats, soybeans, hay, tillable and nontillable pasture and diverted acres. Subtract this total from total acres in farms. The residual is classified as other acres.

(2) Computation of county yields. This information shall be obtained for each county from the USDA National Agricultural Statistics Service.

1. Corn yield (including silage): Number of bushels of corn harvested for grain per acre.
2. Oat yield (including wheat): Number of bushels of oats harvested per acre.
3. Soybean yield: Number of bushels per acre harvested.
4. Hay yield in tons: Number of tons per acre harvested.

(3) Computation of county gross income.

1. Corn: One-half of the 5-year average production multiplied by the 5-year average price received for corn.

2. Silage: One-half of the 5-year average number of acres devoted to the production of silage multiplied by the 5-year average production per acre for corn. The amount of production so determined shall be added to the 5-year average production for corn and included in the determination of the gross income for corn.

3. Soybeans: One-half of the 5-year average production multiplied by the 5-year average price received.

4. Oats: One-half of the 5-year average production of oats and wheat multiplied by the 5-year average price received for oats.

5. Price adjustment: For corn, soybeans, hay, and oats, the prices used shall be as obtained from the USDA National Agricultural Statistics Service and shall be adjusted to reflect any individual county price conditions prior to the 2007 crop year. For the 2007 crop year and later, the USDA National Agricultural Statistics Service district prices shall be used and shall be adjusted to reflect any individual county price conditions.

6. Government programs: Gross income shall be one-half of the 5-year average amount of cash payments or equivalent (such as PIK bushels) including but not limited to diverted, deficiency and PIK programs as reported by the USDA Farm Service Agency.

7. Hay: Gross income shall be a cash rent amount determined by multiplying the 5-year average number of acres devoted to hay by the product obtained by multiplying one-fourth of the 5-year average hay yield by the 5-year average price received for all types of hay.

8. Tillable pasture: Gross income shall be a cash rent amount determined by multiplying the 5-year average number of acres devoted to tillable pasture by the product obtained in “hay” above.

9. Nontillable pasture: Gross income shall be a cash rent amount determined by multiplying the 5-year average number of acres devoted to nontillable pasture by one-half the product obtained in “hay” above.

10. Other acres: Income shall be the product of the number of other acres multiplied by 17 percent of the net income per acre for all other land uses.

(4) Computation of county production costs. The following data and procedures shall be used to determine specific county production costs.

1. Basic average landlord production costs. Landlord production costs for corn, soybeans, oats, diverted acres, hay, tillable pasture, nontillable pasture, fertilizer costs, and facilities’ costs shall be obtained for each year from Iowa State University.

2. Production cost adjustment. The production costs for corn, soybeans, oats, and hay are adjusted for each county by multiplying the difference between the 5-year state average yield per acre and the 5-year county average yield per acre by the 5-year average facilities’ costs. If a county’s yield exceeds the state yield, production costs are increased by this amount. If a county’s yield is less than the state yield, production costs are reduced by this amount.

3. Fertilizer cost adjustment. The adjustment for fertilizer costs is determined as follows: Multiply the difference between the 5-year state average corn yield per acre and the 5-year county average corn yield per acre obtained from the USDA National Agricultural Statistics Service by the fertilizer cost amount per bushel determined by dividing the statewide average cost of landlord’s share of fertilizer cost per acre from Iowa State University by the statewide average corn yield per acre to produce the corn fertilizer cost per bushel adjustment. This amount is then multiplied by the 5-year county average corn acres determined in (2) above.

4. Expense adjustments. If a county’s 5-year average corn yield is greater than the state 5-year average corn yield, this amount is allowed as an additional expense. If the county’s average is less than the state average, this amount is an expense reduction.

5. Liability insurance cost adjustment. The 5-year average per acre cost of obtaining tort liability insurance shall be determined.

(5) Computation of county net income. From the total gross income, subtract the total expenses. Divide the resulting total by the total number of acres.

(6) Computation of dwelling adjustment factor. The amount determined in (5) above shall be reduced by 10.6 percent.

(7) Computation of county tax adjustment. Subtract the 5-year average per acre real estate taxes levied for land and structures including drainage and levee district taxes but excluding those levied against agricultural dwellings from the amount determined in (6) above. Taxes shall be the tax levied for collection during the 5-year period as reported by county auditors, and reduced by the amount of the agricultural land tax credit.

(8) Calculation of county valuation per acre. Divide the net income per acre ((7) above) for each county as determined above by the capitalization rate specified in Iowa Code section 441.21. The quotient shall be the actual per acre equalized valuation of agricultural land and structures for the current equalization year.

b. Use of other relevant data. The department of revenue may also consider other relevant data, including field investigations conducted by representatives of the department, to determine the level of assessment of agricultural real estate.

c. Determination of value. The aggregate actual value of agricultural real estate in each county shall be determined by multiplying the equalized per acre value by the number of acres of agricultural real estate reported on the abstract of assessment for the current year, adjusted where necessary by the results of any field investigations conducted by the department of revenue and any other relevant data available.

102.12(2) Residential real estate outside and within incorporated cities.

a. Use of assessment/sales ratio study.

(1) Basic data shall be that set forth in rule 701—71.10(421) refined by eliminating any sales determined to be abnormal or by adjusting the sales to eliminate the effects of factors that resulted in the determination that the sales were abnormal. The basic data used shall be the assessment/sales ratio study conducted for sales taking place during the calendar year immediately preceding the year in which the equalization order is issued. The department of revenue may also supplement the assessment/sales ratio study with appraisals made by department appraisal personnel for the year immediately preceding the year in which the equalization order is issued. The assessment/sales ratio study including relevant appraisals, if any, shall be used to determine the aggregate actual valuation of residential real estate in each assessing jurisdiction. The department may consider sales and appraisal data for prior years if it is determined the use of the sales and appraisal data for the year immediately preceding the year in which the equalization order is issued is insufficient to determine market value. If such sales and appraisal data for prior years is used, consideration shall be given for any subsequent changes in either assessed value or market value.

(2) Assessors shall provide any known facts or circumstances regarding reported sales transactions and department appraisals that would indicate abnormal or unusual conditions or reporting discrepancies that would necessitate exclusion or adjustment of sales or appraisals from the determination of aggregate actual values. Assessors shall provide those facts within 45 days of receipt from the department of information concerning sales and appraisal data proposed for assessment/sales ratio and equalization purposes.

b. Use of other relevant data. The department of revenue may also consider other relevant data, including field investigations conducted by representatives of the department, to determine the level of assessment of residential real estate.

c. Equalization appraisal selection procedures for residential real estate. Residential properties to be appraised by department of revenue personnel for use in supplementing the assessment/sales ratio study shall be selected for each jurisdiction in the following manner:

(1) The department appraiser assigned to the jurisdiction shall determine a systematic random sequence of numbers equal to the number of appraisals required and document the following steps.

1. The department appraiser assigned to the jurisdiction shall compute the interval number by dividing the total number of improved properties in the classification to be sampled by the number of appraisals to be performed.

EXAMPLE: In this example, ten appraisals are needed with a total of 1,397 improved residential units. Dividing 1,397 by 10, 139.7 is arrived at, which is rounded down to 139. This is the interval number.

2. The selection of the first sequence number shall be accomplished by having an available disinterested person randomly select a number from one through the interval number.

EXAMPLE: In this example a number from 1 to 139 is to be selected. The person randomly selected number 20.

3. The department appraiser shall develop a systematic sequence of numbers equal to the number of appraisals required. Starting with the randomly selected number previously picked by the disinterested person, add the interval number to this number and to each resulting number until a systematic sequence of numbers is obtained.

EXAMPLE: In this example ten appraisals are needed, so a sequence of ten numbers must be developed. Starting with number 20 and adding the interval number of 139 to it, each resulting number provides the following systematic sequence: 20, 159, 298, 437, 576, 715, 854, 993, 1,132, 1,271.

(2) Number of improved properties.

County jurisdictions—Put the name of each city or township having improved units in the classification to be sampled into a hat. Draw each one out of the hat and record its name in the order of its draw. Likewise, record the respective number of improved units for each. Then consecutively number all the improved units and document the procedure.

EXAMPLE:

City or Township	Number of Improved Residential Units	Code Numbers
Franklin Twp.	57	1-57
Pleasant View	160	58-217
Jackson Twp.	56	218-273
Johnston	300	274-573
Polk Twp.	110	574-683
Washington Twp.	114	684-797
Maryville	306	798-1103
Camden Twp.	110	1104-1213
Salem	184	1214-1397
Total	<u>1,397</u>	

(3) Determine the location of the improved properties selected for appraisal and document the procedure.

EXAMPLE:

City or Township	Number of Improved Residential Units	Code Numbers	Sequence Number	Entry on Rolls
Franklin Twp.	57	1-57	20	20
Pleasant View	160	58-217	159	102
Jackson Twp.	56	218-273		
Johnston	300	274-573	298,437	25,164
Polk Twp.	110	574-683	576	3
Washington Twp.	114	684-797	715	32
Maryville	306	798-1103	854,993	57,196
Camden Twp.	110	1104-1213	1132	29
Salem	184	1214-1397	1271	58
Total	<u>1,397</u>			

1. The department appraiser shall locate the property to be appraised by finding the relationship between the sequence numbers and the code numbers and identify the property.

EXAMPLE: The first sequence number is 20. Since the improved residential properties in Franklin Township have been assigned code numbers 1 to 57, sequence number 20 is in that location.

To identify this property, examine the Franklin Township assessment roll book and stop at the twentieth improved residential entry.

Document the parcel number, owner’s name, and legal description of this property.

2. The department appraiser shall appraise the property selected unless it is ineligible because of any of the following restrictions:

- Current year sale
- Partial assessment
- Prior equalization appraisal
- Tax-exempt
- Value established by court action
- Value is not more than \$10,000
- Building on leased land

3. The department appraiser shall determine a substitute property if the originally selected one is ineligible. In ascending order, select code numbers until an eligible property is found.

EXAMPLE: If code number 20 is ineligible, use code number 21 as a substitute. If code number 21 is ineligible, use code number 22, etc., until an eligible property is found.

If the procedure described in 71.12(2)“c”(3)“3” moves the substitute property to another city or township, select substitute code numbers in descending order until an eligible property is found.

If the procedure described in the previous paragraph moves the substitute property to a preceding city or township, go back to the procedure of 71.12(2)“c”(3)“3” even if it moves the substitute property to a subsequent city or township.

4. Select an alternate property for the originally selected property which also would be eligible. This is necessary because at the time of appraisal the property may be found to be ineligible due to one of the restrictions in 71.12(2)“c”(3)“2.” Alternate properties are selected by using the same procedure described in 71.12(2)“c”(3)“3.”

5. Follow procedures 71.12(2)“c”(3), items “1” to “4,” for each of the other originally selected sequence numbers.

102.12(3) Reserved.

102.12(4) *Commercial real estate.*

a. Use of assessment/sales ratio study.

(1) Basic data shall be that set forth in rule 701—71.10(421), refined by eliminating any sales determined to be abnormal or by adjusting same to eliminate the effects of factors that resulted in the determination that the sales were abnormal. The basic data used shall be the assessment/sales ratio study conducted for sales taking place during the calendar year immediately preceding the year in which the equalization order is issued. The department of revenue may also supplement the assessment/sales ratio study with appraisals made by department appraisal personnel for the year immediately preceding the year in which the equalization order is issued. The assessment/sales ratio study including relevant appraisals, if any, shall be used to determine the aggregate actual valuation of commercial real estate in each assessing jurisdiction. The department may consider sales and appraisal data for prior years if it is determined the use of sales and appraisal data for the year immediately preceding the year in which the equalization order is issued is insufficient to determine market value. If such sales and appraisal data for prior years are used, consideration shall be given for any subsequent changes in either assessed value or market value. Properties receiving a dual classification with the primary use being commercial shall be included.

(2) Assessors shall provide any known facts or circumstances regarding reported sales transactions and department appraisals that would indicate abnormal or unusual conditions or reporting discrepancies that would necessitate exclusion or adjustment of sales or appraisals from the determination of aggregate actual values. Assessors shall provide those facts within 45 days of receipt from the department of information concerning sales and appraisal data proposed for assessment/sales ratio and equalization purposes.

b. Use of other relevant data. The department of revenue may also consider other relevant data and statistical measures, including field investigations conducted by representatives of the department, to determine the level of assessment of commercial real estate. The diverse nature of commercial real estate precludes the use of a countywide or citywide income capitalization study.

c. Equalization appraisal selection procedures for commercial real estate. Commercial properties to be appraised by department of revenue personnel for use in supplementing the assessment/sales ratio study shall be selected for each jurisdiction in the manner outlined below. Properties receiving a dual classification with the primary use being commercial shall be included.

(1) The department appraiser assigned to the jurisdiction shall determine a systematic random sequence of numbers equal to the number of appraisals required and document the following steps.

1. The department appraiser shall compute the interval number by dividing the total number of improved properties in the classification to be sampled by the number of appraisals to be performed.

EXAMPLE: In this example, ten appraisals are needed with a total of 397 improved commercial units. Dividing 397 by 10, 39.7 is arrived at, which is rounded down to 39. This is the interval number.

2. The selection of the first sequence number shall be accomplished by having an available disinterested person randomly select a number from one through the interval number.

EXAMPLE: In this example a number from 1 to 39 is to be selected. The person randomly selected number 2.

3. The department appraiser shall develop a systematic sequence of numbers equal to the number of appraisals required. Starting with the randomly selected number previously picked by the disinterested person, add the interval number to this number and to each resulting number until a systematic sequence of numbers is obtained.

EXAMPLE: In this example ten appraisals are needed, so a sequence of ten numbers must be developed. Starting with number 2 and adding the interval number of 39 to it, each resulting number provides the following systematic sequence: 2, 41, 80, 119, 158, 197, 236, 275, 314, 353.

(2) Number of improved properties.

1. City jurisdictions—Utilizing the assessment book or a computer printout which follows the same order as the assessment book, consecutively number all the improved units and document the procedure.

2. County jurisdictions—Put the name of each city or township having improved units in the classification to be sampled into a hat. Draw each one out of the hat and record its name in the order of its draw. Likewise, record the respective number of improved units for each. Then consecutively number all the improved units and document the procedure.

EXAMPLE:

City or Township	Number of Improved Commercial Units	Code Numbers
Franklin Twp.	4	1-4
Pleasant View	60	5-64
Jackson Twp.	9	65-73
Johnston	100	74-173
Polk Twp.	10	174-183
Washington Twp.	14	184-197
Maryville	106	198-303
Camden Twp.	10	304-313
Salem	84	314-397
Total	<u>397</u>	

(3) The department appraiser shall determine the location of the improved properties selected for appraisal and document the procedure.

EXAMPLE:

<u>City or Township</u>	<u>Number of Improved Commercial Units</u>	<u>Code Numbers</u>	<u>Sequence Number</u>	<u>Entry on Rolls</u>
Franklin Twp.	4	1-4	2	2
Pleasant View	60	5-64	41	37
Jackson Twp.	9	65-73		
Johnston	100	74-173	80,119,158	7,46,85
Polk Twp.	10	174-183		
Washington Twp.	14	184-197	197	14
Maryville	106	198-303	236,275	39,78
Camden Twp.	10	304-313		
Salem	84	314-397	314,353	1,40
Total	<u>397</u>			

1. The department appraiser shall locate the property to be appraised by finding the relationship between the sequence numbers and the code numbers and identify the property.

EXAMPLE: The first sequence number is 2. Since the improved commercial properties in Franklin Township have been assigned code numbers 1 to 4, sequence number 2 is in that location.

To identify this property, examine the Franklin Township assessment roll book and stop at the second improved commercial entry.

The department appraiser shall document the parcel number, owner's name, and legal description of this property.

2. The department appraiser shall appraise the property selected unless it is ineligible because of any of the following restrictions:

Vacant building

Current-year sale

Partial assessment

Prior equalization appraisal

Tax-exempt

Only one portion of a total property unit (example—a parking lot of a grocery store)

Value established by court action

Value is not more than \$10,000

Building on leased land

3. The department appraiser shall determine a substitute property if the originally selected one is ineligible. In ascending order, select code numbers until an eligible property is found.

EXAMPLE: If code number 2 is ineligible, use code number 3 as a substitute. If code number 3 is ineligible, use code number 4, etc., until an eligible property is found.

If the procedure described in 71.12(4) "c"(3)"3" moves the substitute property to a city or township, select substitute code numbers in descending order until an eligible property is found.

If the procedure described in the previous paragraph moves the substitute property to a preceding city or township, go back to the procedure of 71.12(4) "c"(3)"3" even if it moves the substitute property to a subsequent city or township.

4. Select an alternate property for the originally selected property which also would be eligible. This is necessary because at the time of appraisal the property may be found to be ineligible due to one of the restrictions in 71.12(4) "c"(3)"2." Alternate properties are selected by using the same procedure described in 71.12(4) "c"(3)"3."

5. Follow procedures 71.12(4) "c"(3), items "1" to "4," for each of the other originally selected sequence numbers.

102.12(5) Industrial real estate. It is not possible to determine the level of assessment of industrial real estate by using accepted equalization methods. The lack of sales data precludes the use of an

assessment/sales ratio study, the diverse nature of industrial real estate precludes the use of a countywide or citywide income capitalization study, and the limited number of industrial properties precludes the use of sample appraisals. The level of assessment of industrial real estate can only be determined by the valuation of individual parcels of industrial real estate. Any attempt to equalize industrial valuations by using accepted equalization methods would create an arbitrary result. However, under the circumstances set forth in Iowa Code subsection 421.17(10), the department may correct any errors in such assessments that are brought to the attention of the department, including errors related to property with a dual classification if the primary use of the property is from the industrial portions.

102.12(6) Centrally assessed property. Property assessed by the department of revenue pursuant to Iowa Code chapters 428 and 433 to 438, inclusive, is equalized internally by the department in the making of the assessments. Further, the assessments are equalized with the aggregate valuations of other classes of property as a result of actions taken by the department pursuant to rule 701—71.11(441).

102.12(7) Miscellaneous real estate. Since it is not possible to use accepted equalization methods to determine the level of assessment of mineral rights and interstate railroad and toll bridges, these classes of property shall not be subject to equalization by the department of revenue. However, under the circumstances set forth in Iowa Code section 421.17(10), the department may correct any errors in assessments which are brought to the attention of the department.

This rule is intended to implement Iowa Code sections 441.21, 441.47, 441.48 and 441.49.
[ARC 7726B, IAB 4/22/09, effective 5/27/09; ARC 9478B, IAB 4/20/11, effective 5/25/11; ARC 1765C, IAB 12/10/14, effective 1/14/15; ARC 2657C, IAB 8/3/16, effective 9/7/16; ARC 4170C, IAB 12/5/18, effective 1/9/19; ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.13(441) Tentative equalization notices. Prior to the issuance of the final equalization order to each county auditor, a tentative equalization notice providing for proposed percentage adjustments to the aggregate valuations of classes of property as set forth in rule 701—71.12(441) shall be mailed to the county auditor whose valuations are proposed to be adjusted. The tentative equalization notice constitutes the ten days' notice required by Iowa Code section 441.48.

This rule is intended to implement Iowa Code sections 441.47 and 441.48.
[Editorial change: IAC Supplement 11/2/22]

701—102.14(441) Hearings before the department.

102.14(1) Protests. Written or oral protest against the proposed percentage adjustments as set forth in the tentative equalization notice issued by the department of revenue shall be made only on behalf of the affected assessing jurisdiction. The protests shall be made only by officials of the assessing jurisdiction, including, but not limited to, an assessing jurisdiction's city council or board of supervisors, assessor, or city or county attorney. An assessing jurisdiction may submit a written protest in lieu of making an oral presentation before the department, or may submit an oral protest supported by written documentation. Protests against the adjustments in valuation contained in the tentative equalization notices shall be limited to a statement of the error or errors complained of and shall include such facts as might lead to their correction. No other factors shall be considered by the department in reviewing the protests. Protests and hearings on tentative equalization notices before the department are excluded from the provisions of the Iowa Administrative Procedure Act governing contested case proceedings.

102.14(2) Conduct of hearing. The department shall schedule each hearing so as to allow the same amount of time within which each assessing jurisdiction can make its presentation. During the hearing each assessing jurisdiction shall be afforded the opportunity to present evidence relevant to its protest. The division administrator for the local government services division shall act as the department's representative. The department's representative shall preside at the hearing, which shall be held at the time and place designated by the department or such other time and place as may be mutually agreed upon by the department and the protesting assessing jurisdiction.

This rule is intended to implement Iowa Code section 441.48.
[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22; ARC 6958C, IAB 3/22/23, effective 4/26/23]

701—102.15(441) Final equalization order and appeals.

102.15(1) *Issuance of final equalization order.* After the tentative equalization notice has been issued and an opportunity for a hearing described in rule 701—71.14(441) has been afforded, the department of revenue shall issue a final equalization order by mail to the county auditor. The order shall specify any percentage adjustments in the aggregate valuations of any class of property to be made effective for the county as of January 1 of the year in which the order is issued. The final equalization order shall be issued on or before October 1 unless for good cause it cannot be issued until after October 1. The final equalization order shall be implemented by the county auditor.

102.15(2) *Appeal of final equalization order.* The city or county officials of the affected county or assessing jurisdiction may appeal a final equalization order to the director of revenue by filing a notice of appeal with the clerk of the hearings section of the department of revenue. The notice of appeal must be filed or postmarked not later than ten days after the date the final equalization order is issued.

a. Form of appeal. The notice of appeal shall be in writing and in the same format as provided in 701—subrule 7.8(6).

(1) The notice of appeal shall substantially state in separate numbered paragraphs the following:

1. The county or assessing jurisdiction;
2. The date on which the final equalization order was issued;
3. The portion of the equalization order being appealed;
4. A clear and concise assignment of each and every error;
5. A clear and concise statement of the facts upon which the affected county or assessing jurisdiction relies as sustaining the assignment of error;
6. The relief requested;
7. The signature of the city or county officials bringing the appeal, or their representative, along with the address to which all subsequent correspondence, notice or papers shall be served or mailed.

(2) A county or assessing jurisdiction may amend its notice of appeal at any time prior to the commencement of the evidentiary hearing. The department may request that the county or assessing jurisdiction amend the notice of appeal for clarification.

b. Filing of notice of appeal. The notice of appeal must either be delivered to the department by electronic means or by United States Postal Service or a common carrier, by ordinary, certified, or registered mail, directed to the attention of the clerk of the hearings section at P.O. Box 14457, Des Moines, Iowa 50319, or be personally delivered to the clerk of the hearings section or served on the clerk of the hearings section by personal service during business hours. For the purpose of mailing, a notice of appeal is considered filed on the date of the postmark. If a postmark date is not present on the mailed article, then the date of receipt of protest will be considered the date of mailing. Any document, including a notice of appeal, is considered filed on the date personal service or personal delivery to the office of the clerk of the hearings section is made. See Iowa Code section 622.105 for the evidence necessary to establish proof of mailing.

c. Answer. The department of revenue shall file an answer with the clerk of the hearings section within 30 days after the filing of the pleading responded to, unless attacked by motion as provided in 701—subrule 7.17(5), and then the answer shall be filed within 30 days after the date on which the fact finder issues a ruling on the motion. The department may amend its answer at any time prior to the commencement of the evidentiary hearing.

d. Docketing. Appeals shall be assigned a docket number as provided in rule 701—7.10(17A). Records consisting of the case name and the corresponding docket number assigned to the case must be maintained by the clerk of the hearings section. The records of each case shall also include each action and each act done, with the proper dates as follows:

- (1) The title of the appeal;
- (2) Brief statement of the date of the final equalization order, the property tax classification affected, and the relief sought;
- (3) The manner and time of service of notice of appeal;
- (4) The appearance of all parties;
- (5) Notice of hearing, together with manner and time of service; and

(6) The decision of the director or administrative law judge or other disposition of the case and the date.

e. Hearing. Rules 701—7.14(17A) through 701—7.22(17A) shall apply to any hearing or proceeding regarding the appeal of a final equalization order to the director of revenue.

This rule is intended to implement Iowa Code chapter 17A and sections 441.48 and 441.49.
[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.16(441) Alternative method of implementing equalization orders.

102.16(1) Application for permission to use an alternative method.

a. A request by an assessing jurisdiction for permission to use an alternative method of applying the final equalization order must be made in writing to the department of revenue within ten days from the date the county auditor receives the final equalization order. The written request shall include the following information:

(1) Facts evidencing the need to use an alternative method of implementing the final equalization order. Such facts shall clearly show that the proposed method is essential to ensure compliance with the provisions of Iowa Code section 441.21.

(2) The exact methods to be employed in implementing the requested alternative method for each class of property.

(3) The specific method of notifying affected property owners of the valuation changes.

(4) Evidence that the alternative method will result in an aggregate property class valuation adjustment equivalent to that prescribed in the department's final equalization order.

b. The department of revenue shall review each written request for an alternative method and shall notify the assessing jurisdiction of acceptance or rejection of the proposed method by October 15. The assessing jurisdiction shall immediately inform the county auditor of the department's decision. The county auditor shall include a description of any approved alternative method in the required newspaper publication of the final equalization order. In those instances where the approved alternative method includes individual property owner notification, the publication shall not be considered proper notice to the affected property owners.

102.16(2) Implementation of alternative method. If an alternative method is approved by the department of revenue, any individual notification of property owners shall be completed by the assessor by not later than October 25.

102.16(3) Appeal by property owners. If an alternative method is approved by the department of revenue, the special session of the local board of review to hear equalization protests shall be extended to November 30. In such instances, protests may be filed up to and including November 4.

This rule is intended to implement Iowa Code section 441.49.
[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.17(441) Special session of boards of review.

102.17(1) Grounds for protest. The only ground for protesting to the local board of review reconvened in special session pursuant to Iowa Code section 441.49 is that the application of the department's final equalization order results in a value greater than that permitted under Iowa Code section 441.21.

102.17(2) Authority of board of review. When in special session to hear protests resulting from equalization adjustments, the local board of review shall only act upon protests for those properties for which valuations have been increased as a result of the application of the department of revenue's final equalization order.

The local board of review may adjust valuations of those properties it deems warranted, but under no circumstance shall the adjustment result in a value less than that which existed prior to the application of the department's equalization order. The local board of review shall not adjust the valuation of properties for which no protests have been filed.

102.17(3) Report of board of review. In the report to the department of revenue of action taken by the local board of review in special session, the board of review shall report the aggregate valuation

adjustments by class of property as well as all other information required by the department of revenue to determine if such actions may have substantially altered the equalization order.

102.17(4) Meetings of board of review. If the final equalization order does not increase the valuation of any class of property, the board of review is not required to meet during the special session. If the final equalization order increases the valuation of one or more classes of property but no protests are filed by the times specified in Iowa Code section 441.49, the board of review is not required to meet during the special session.

This rule is intended to implement Iowa Code sections 421.17(10) and 441.49.
[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.18(441) Judgment of assessors and local boards of review. Nothing stated in these rules should be construed as prohibiting the exercise of honest judgment, as provided by law, by the assessors and local boards of review in matters pertaining to valuing and assessing of individual properties within their respective jurisdictions.

This rule is intended to implement Iowa Code sections 441.17 and 441.35.
[Editorial change: IAC Supplement 11/2/22]

701—102.19(441) Conference boards.

102.19(1) Establishment and abolition of office.

a. As referred to in Iowa Code section 441.1, the term “federal census” includes any special census conducted by the Bureau of the Census of the U.S. Department of Commerce as well as the Bureau’s decennial census.

b. Within 60 days of receiving the certified results of a federal census indicating the population of a city having its own assessor has fallen below 10,000, the city council of the city shall repeal the ordinance providing for its own assessor.

c. Whenever the office of city assessor is abolished, all moneys in the assessment expense fund and the special appraiser fund shall be transferred to the appropriate accounts in the county assessor’s office, and all equipment and supplies shall be transferred to the county assessor’s office. Employees of the city assessor’s office may, at the discretion of the county assessor, become employees of the county assessor. However, any deputy assessor of the city may not be appointed a deputy county assessor unless certified as eligible for appointment pursuant to Iowa Code sections 441.5 and 441.10.

102.19(2) Membership.

a. County conference boards. A county conference board consists of the county board of supervisors, the mayor of each incorporated city in the county whose property is assessed by the county assessor, and one member of the board of directors of each high school district in the county, provided the member is a resident of the county. Members representing school districts serve one-year terms, and the board of directors each year must notify the clerk of the conference board of its representative on the conference board. A member of the board of directors of a school district may serve on the county conference board even though the member lives in a city having its own assessor (1978 O.A.G. 466).

b. City conference boards. A city conference board consists of the county board of supervisors, the city council, and the entire board of directors of each school district whose property is assessed by the city assessor.

102.19(3) Voting.

a. Votes on matters before a conference board shall be by units as provided in Iowa Code section 441.2. At least two members of each voting unit must be present in order for the unit to cast a vote (1960 O.A.G. 226). In the event the vote of the members of a voting unit ends in a tie, that unit shall not cast a vote on the particular matter before the conference board.

b. If a member of a conference board is absent from a meeting, the member’s vote may not be cast by another person, except that a mayor pro tem as provided in Iowa Code section 372.14(3) may vote for the mayor when the mayor is absent from or unable to perform official duties.

This rule is intended to implement Iowa Code section 441.2.
[Editorial change: IAC Supplement 11/2/22]

701—102.20(441) Board of review.**102.20(1) Membership.**

a. Occupation of members. One member of the county board of review must be actively engaged in farming as that member's primary occupation. However, it is not necessary for a board of review to have as a member one licensed real estate broker and one licensed architect or person experienced in the building and construction field if the person cannot be located after a good-faith effort to do so has been made by the conference board (1966 O.A.G. 416). In determining eligibility for membership on a board of review, a retired person is not considered to be employed in the occupation pursued prior to retirement, unless that person remains in reasonable contact with the former occupation, including some participation in matters associated with that occupation.

b. Residency of members. A person must be a resident of the assessor jurisdiction served to qualify for appointment as a member of the board of review. However, a member changing assessing jurisdiction residency after appointment to the board may continue to serve on the board until the member's current term of office expires.

c. Term of office. The term of office of members of boards of review shall be for six years and shall be staggered as provided in Iowa Code section 441.31. In the event of the death, resignation, or removal from office of a member of a board of review, the conference board or city council shall appoint a successor to serve the unexpired term of the previous incumbent.

d. Membership on other boards. A member of a board of review shall not at the same time serve on either the conference board or the examining board, or be an employee of the assessor's office (1948 O.A.G. 120, 1960 O.A.G. 226).

e. Number of members. A conference board or city council may at any time change the composition of a board of review to either three or five members. To reduce membership from five members to three members, the conference board or city council shall not appoint successors to fill the next two vacancies which occur (1970 O.A.G. 342). To increase membership from three members to five members, the conference board or city council shall appoint two additional members whose initial terms shall expire at such times so that no two board members' terms expire at the end of the same year. Also, the conference board or city council may increase the membership of the board of review by an additional two members if it determines that a large number of protests warrant the emergency appointments. If the board of review has ten members, not more than four additional members may be appointed by the conference board. The terms of the emergency members will not exceed two years.

f. Removal from office. A member of a board of review may be removed from office by the conference board or city council but only after specific charges have been filed by the conference board or city council.

g. Appointment of members. Members of a county board of review shall be appointed by the county conference board. Members of a city board of review shall be appointed by the city conference board in cities with an assessor or by the city council in cities without an assessor. A city without an assessor can only have a board of review if the population of the city is 75,000 or more. A city with a population of more than 125,000 may appoint a city board of review or request the county conference board to appoint a ten-member county board of review.

102.20(2) Sessions of boards of review.

a. It is mandatory that a board of review convene on May 1 and adjourn no later than May 31 of each year. However, if either date falls on a Saturday, Sunday, or legal holiday, the board of review shall convene or adjourn on the following Monday.

b. Extended session. If a board of review determines it will be unable to complete its work by May 31, it may request that the director of revenue extend its session up to July 15. The request must be signed by a majority of the membership of the board of review and must contain the reasons the board of review cannot complete its work by May 31. During the extended session, a board of review may perform the same functions as during its regular session unless specifically limited by the director of revenue.

c. Special session. If a board of review is reconvened by the director of revenue pursuant to Iowa Code section 421.17, the board of review shall perform those functions specified in the order of the director of revenue and shall perform no other functions.

102.20(3) Actions initiated by boards of review.

a. Internal equalization of assessments. A board of review in reassessment years as provided in Iowa Code section 428.4 has the power to equalize individual assessments as established by the assessor, but cannot make percentage adjustments in the aggregate valuations of classes of property (1966 O.A.G. 416). In nonreassessment years, a board of review can adjust the valuation of an entire class of property by adjusting all assessment by a uniform percentage. Nothing contained in this rule shall restrict the director from exercising the responsibilities set forth in Iowa Code section 421.17.

b. Omitted assessments. A board of review may assess for taxation any property which was not assessed by the assessor, including property which the assessor determines erroneously is not subject to taxation by virtue of enjoying an exempt status (*Talley v. Brown*, 146 Iowa 360, 125 N.W. 248 (1910)).

c. Notice to taxpayers. If the value of any property is increased by a board of review or a board of review assesses property not previously assessed by the assessor, the person to whom the property is assessed shall be notified by regular mail of the board's action. The notification shall state that the taxpayer may protest the action by filing a written protest with the board of review within five days of the date of the notice. After at least five days have passed since notifying the taxpayer, the board of review shall meet to take final action on the matter, including the consideration of any protest filed. However, if the valuations of all properties within a class of property are raised or lowered by a uniform percentage in a nonreassessment year, notice to taxpayers shall be provided by newspaper publication as described in Iowa Code section 441.35 and in the manner specified in Iowa Code section 441.36.

102.20(4) Appeals to boards of review.

a. Jurisdiction. A board of review may act only upon written protests which have been filed with the board of review in compliance with Iowa Code section 441.37(1) "a."

(1) Protests must be filed between April 2 and April 30, inclusive. In the event April 30 falls on a Saturday or Sunday, protests filed the following Monday shall be considered to have been timely filed. Protests postmarked by April 30 or the following Monday if April 30 falls on a Saturday or Sunday shall also be considered to have been timely filed.

(2) The protest must identify one or more grounds for protest under Iowa Code section 441.37.

(3) All protests must be in writing, on forms prescribed by the director of revenue, and signed by the protester or the protester's authorized agent. A protest shall not be rejected for the sole reason that the protest was not filed using the prescribed form if the protest otherwise complies with Iowa Code section 441.37(1) "a." A written request for an oral hearing must be made at the time of filing the protest. The protester may combine on one form assessment protests on parcels separately assessed if the same grounds are relied upon as the basis for protesting each separate assessment. If an oral hearing is requested on more than one of the protests, the person making the combined protests may request that the oral hearings be held consecutively.

(4) A board of review may allow protests to be filed in electronic format. Protests transmitted electronically are subject to the same deadlines as written protests.

b. Grounds for protest. Taxpayers may protest to a board of review on one or more of the grounds specified in Iowa Code section 441.37. The grounds for protest and procedures for considering protests are as follows:

(1) The assessment is not equitable when compared with those of similar properties in the same taxing district. If this ground is a basis for the protest, the protester may identify comparable properties to support the claim. In considering a protest based upon this ground, the board of review should examine carefully all information used to determine the assessment of the subject property, consider any comparable properties, and determine whether the evidence demonstrates the subject property is inequitably assessed.

(2) The property is assessed at more than the value authorized by law. If this ground is the basis for a protest, the protester may indicate the amount considered to be the actual value of the property.

(3) The property is not assessable, is exempt from taxes, or is misclassified. If this ground is the basis for a protest, the protester may indicate why the property is exempt, misclassified, or not assessable.

(4) There is an error in the assessment. An error may include, but is not limited to, listing errors, assessment of subject property for less than authorized by law, or erroneous mathematical calculations. If this ground is the basis for a protest, the protester must indicate the alleged error.

A board of review must determine:

1. If an error exists, and
2. How the error might be corrected.

(5) There is fraud or misconduct in the assessment. If this ground of protest is used, the protester must state the specific fraud or misconduct alleged, and the board of review must first determine if there is validity to the protester's allegation. If it is determined that there is fraud in the assessment or that there has been misconduct by the assessor, the board of review shall take action to correct the assessment and report the matter to the director of revenue. For purposes of this subrule, "misconduct" means the same as defined in 2017 Iowa Code section 441.9.

(6) Protests may be filed for previous years if the protester discovers that a mathematical or clerical error was made in the assessment, provided the taxes have not been fully paid or otherwise legally discharged.

c. Disposition of protests. After reaching a decision on a protest, the board of review shall give the taxpayer written notice of its decision. The decision shall be mailed no later than three days after the board of review's adjournment. The notice shall contain the following information:

(1) The valuation and classification of the property as determined by the board of review.

(2) If the protest was based on the ground the property was not assessable, the notice shall state whether the exemption is allowed and the value at which the property would be assessed in the absence of the exemption.

(3) The specific reasons for the board's decision with respect to the protest.

(4) That the board of review's decision may be appealed to either the property assessment appeal board or district court within 20 days of the board's adjournment or May 31, whichever date is later. If the adjournment date is known, the date shall be stated on the notice. If the adjournment date is not known, the notice shall state the date will be no earlier than May 31.

1. Appeal to property assessment appeal board. An appeal from the board of review to the property assessment appeal board may be made pursuant to the provisions of Iowa Code section 441.37A and rule 701—126.1(421,441).

2. Appeal to district court. An appeal from the board of review to the district court may be made pursuant to the provisions of Iowa Code section 441.38. The appeal shall be filed in the county where the property is located. Notice of the appeal shall be served on the chairperson, presiding officer, or clerk of the board of review after the written notice of appeal has been filed with the clerk of district court.

This rule is intended to implement Iowa Code sections 441.31 to 441.37 and Iowa Code Supplement section 441.38 as amended by 2006 Iowa Acts, House File 2794.

[ARC 2707C, IAB 9/14/16, effective 10/19/16; ARC 3312C, IAB 9/13/17, effective 10/18/17; ARC 3771C, IAB 4/25/18, effective 5/30/18; Editorial change: IAC Supplement 11/2/22]

701—102.21(421,17A) Property assessment appeal board. Rescinded ARC 6858C, IAB 2/8/23, effective 3/15/23.

701—102.22(428,441) Assessors.

102.22(1) Conflict of interest. An assessor shall not act as a private appraiser, or as a real estate broker or option agent in the jurisdiction in which serving as assessor (1976 O.A.G. 744).

102.22(2) Listing of property.

a. Forms. Assessors may design and use their own forms in lieu of those prescribed by the department of revenue provided that the forms contain all information contained on the prescribed form, are not substantially different from the prescribed form, and are approved by the director of revenue.

b. Assessment rolls. Assessment rolls must be prepared in duplicate for each property in a reassessment year as defined in Iowa Code section 428.4. However, the copy of the roll does not have to

be issued to a taxpayer unless there is a change in the assessment or the taxpayer requests the issuance of the duplicate copy.

c. Whenever a date specified in Iowa Code chapter 441 falls on a Saturday, Sunday, or legal holiday, the action required to be completed on or before that date shall be considered to have been timely completed if performed on or before the following day which is not a Saturday, Sunday, or holiday.

d. Buildings erected or improvements made by a person other than the owner of the land on which they are located are to be assessed to the owner of the buildings or improvements. Unpaid taxes are a lien on the buildings or improvements and not a lien on the land on which they are located.

102.22(3) Notice of protest. If a protest or appeal is filed with the board of review, property assessment appeal board, or district court against the assessment of property valued at \$5 million or more, the assessor shall provide notice to the school district in which the property is located within ten days of the filing of the protest or the appeal, as applicable.

This rule is intended to implement Iowa Code chapter 428 and Iowa Code chapter 441 as amended by 2006 Iowa Acts, House File 2797.

[Editorial change: IAC Supplement 11/2/22]

701—102.23 Reserved.

701—102.24(421,428,441) Valuation of dual classification property. Real estate with a dual classification of commercial/residential or industrial/residential shall be assessed at its actual value as defined in Iowa Code section 441.21.

102.24(1) Allocation of dual classification values. The assessor shall value as a whole properties that have portions classified as residential and portions classified as commercial or industrial. After the assessor has assigned a value to the property, the value shall be allocated between the two classes of property based on the appropriate appraisal methodology. The assessor shall allocate land value proportionately by class.

102.24(2) Notice of valuation. The valuation notice issued pursuant to Iowa Code section 441.23 shall include a breakdown of the valuation by class for the current year and the prior year.

102.24(3) Protest of assessment. The valuation and assessment of property with a dual classification shall be considered one assessment, and any protest of assessment brought under Iowa Code section 441.37 or subsequent appeal must be made on the entire assessment. Protests of assessments on the valuation of only one class of property are not permitted. The board of review shall review the valuation in total as both classifications are subject to the board's adjustment in any review proceeding. Likewise, any tribunal or court reviewing the board's decision shall base its review on the entire assessment.

This rule is intended to implement Iowa Code sections 421.17, 428.4 and 441.21 as amended by 2013 Iowa Acts, Senate File 295.

[ARC 1765C, IAB 12/10/14, effective 1/14/15; ARC 6096C, IAB 12/15/21, effective 1/19/22; Editorial change: IAC Supplement 11/2/22]

701—102.25(441,443) Omitted assessments.

102.25(1) Property subject to omitted assessment.

a. Land and buildings. An omitted assessment can be made only if land or buildings were not listed and assessed by the assessor. The failure to list and assess an entire building is an omission for which an omitted assessment can be made even if the land upon which the building is located has been listed and assessed. See *Okland v. Bilyeu*, 359 N.W.2d 412 (Iowa 1984). However, the failure to consider the value added as a result of an improvement made does not constitute an omission for which an omitted assessment can be made if the building or land to which the improvement was made has been listed and assessed.

b. Previously exempt property. Property which has been erroneously determined to be exempt from taxation may be restored to taxation by the making of an omitted assessment. See *Talley v. Brown*, 146 Iowa 360, 125 N.W. 243 (1910). An omitted assessment is also made to restore to taxation previously exempt property which ceases to be eligible for an exemption.

102.25(2) Officials authorized to make an omitted assessment.

a. Local board of review. A local board of review may make an omitted assessment of property during its regular session only if the property was not listed and assessed as of January 1 of the current assessment year. For example, during its regular session which begins May 1, 1986, a local board of review may make an omitted assessment only of property that was not assessed by the assessor as of January 1, 1986. During that session, the board of review could not make an omitted assessment for an assessment year prior to 1986.

b. County auditor and local assessor. The county auditor and local assessor may make an omitted assessment. However, no omitted assessment can be made by the county auditor or local assessor if taxes based on the assessment year in question have been paid or otherwise legally discharged. For example, if a tract of land was listed and assessed and taxes levied against that assessment have been paid or legally discharged, no omitted assessment can be made of a building located upon that tract of land even though the building was not listed and assessed at the time the land was listed and assessed. See *Okland v. Bilyeu*, 359 N.W.2d 412, 417 (Iowa 1984).

c. County treasurer. The county treasurer may make an omitted assessment within two years from the date the tax list which should have contained the assessment should have been delivered to the county treasurer. For example, for the 1999 assessment year, the tax list is to be delivered to the county treasurer on or before June 30, 2000. Thus, the county treasurer may make an omitted assessment for the 1999 assessment year at any time on or before June 30, 2002. The county treasurer may make an omitted assessment of a building even if taxes levied against the land upon which the building is located have been paid or legally discharged. See *Okland v. Bilyeu*, 359 N.W.2d 412, 417 (Iowa 1984). The county treasurer may not make an omitted assessment if the omitted property is no longer owned by the person who owned the property on January 1 of the year the original assessment should have been made.

d. Department of revenue. The department of revenue may make an omitted assessment of any property assessable by the department at any time within two years from the date the assessment should have been made.

This rule is intended to implement Iowa Code chapter 440 and sections 443.6 through 443.15 as amended by 1999 Iowa Acts, chapter 174.

[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.26(441) Assessor compliance.

102.26(1) The assessor shall determine the value of real property in accordance with rules adopted by the department of revenue and in accordance with forms and guidelines contained in the Iowa Real Property Appraisal Manual prepared by the department. The assessor may use an alternative manual to value property if it is a unique type of property not covered in the manual prepared by the department.

102.26(2) If the department finds that an assessor is not in compliance with the rules of the department relating to valuation of property or has disregarded the forms and guidelines contained in the real property appraisal manual, the department shall notify the assessor and each member of the conference board for that assessing jurisdiction. The notice shall be mailed by restricted certified mail and shall specify the areas of noncompliance and the steps necessary to achieve compliance. The notice shall also inform the assessor and conference board that if compliance is not achieved, a penalty may be imposed.

102.26(3) The conference board shall respond to the department within 30 days of receipt of the notice of noncompliance. The conference board may respond to the notice by asserting that the assessor is in compliance with the rules, guidelines, and forms of the department or by informing the department that the conference board intends to submit a plan of action to achieve compliance. If the conference board responds to the notification by asserting that the assessor is in compliance, a hearing before the director of revenue shall be held on the matter within 60 days of receipt of the notice of noncompliance. The director's decision is subject to judicial review in accordance with Iowa Code chapter 17A. If it is agreed that the assessor is not in compliance, the conference board shall submit a plan of action within 60 days of receipt of the notice of noncompliance.

102.26(4) The plan of action shall contain a time frame under which compliance shall be achieved, which shall be no later than January 1 of the following assessment year. The plan shall contain the

signature of the assessor and of the chairperson of the conference board. The department shall review the plan to determine whether the plan is sufficient to achieve compliance. Within 30 days of receipt of the plan, the department shall notify the assessor and the chairperson of the conference board that it has accepted the plan or that it is necessary to submit an amended plan of action.

102.26(5) By January 1 of the assessment year following the calendar year in which the plan of action was submitted to the department, the conference board shall submit a report to the department verifying that the plan was followed and compliance has been achieved. The department may conduct a field inspection to ensure that the assessor is in compliance. By January 31, the department shall notify the assessor and the conference board, by restricted certified mail, either that compliance has been achieved or that the assessor remains in noncompliance. If the department determines that the assessor remains in noncompliance, the department shall take steps to withhold up to 5 percent of the reimbursement payment authorized in Iowa Code section 425.1 until the department determines that the assessor is in compliance.

102.26(6) If the conference board disputes the determination of the department, the chairperson of the conference board may appeal the determination to the director of revenue under 701—Chapter 7.

This rule is intended to implement Iowa Code section 441.21.

[ARC 2657C, IAB 8/3/16, effective 9/7/16; Editorial change: IAC Supplement 11/2/22]

701—102.27(441) Assessor shall not assess own property.

102.27(1) *Assessor and deputy assessor prohibited from assessing own property.* An assessor or deputy assessor shall not personally assess a property if the assessor or deputy assessor owns the property, has a financial interest in the property, or has a financial interest in the entity that owns the property. The assessing jurisdiction shall pay all costs and expenses associated with the assessment of the above property.

102.27(2) *Certification to the department.*

a. Not later than January 1 of each year, assessors shall certify to the director that the assessor did not personally assess the following property in the previous assessment year:

- (1) Property owned by the assessor;
- (2) Property in which the assessor has a financial interest;
- (3) Property owned by an entity in which the assessor has a financial interest.

b. Not later than January 1 of each year, deputy assessors shall certify to the director that the deputy assessor did not personally assess the following property in the previous assessment year:

- (1) Property owned by the deputy assessor;
- (2) Property in which the deputy assessor has a financial interest;
- (3) Property owned by an entity in which the deputy assessor has a financial interest.

c. Assessors and deputy assessors shall use forms and procedures prescribed and provided by the director for the certifications described in paragraphs 71.27(2) “a” and “b.”

102.27(3) *Powers and duties of director.* The director shall have and assume all of the powers and duties under Iowa Code section 421.17 in administering this rule.

102.27(4) *Definitions.* For purposes of this rule, the following definitions shall govern.

“*Financial interest*” includes but is not limited to the holding of legal title to real property or any ownership interest in an entity that holds legal title to real property. Notwithstanding the preceding sentence, ownership interest in an entity shall not be deemed a “financial interest” when a person’s ownership interest equals less than 10 percent of the entity’s total ownership interest.

“*Personally assess*” means engaging in the listing, valuation, and classification of real property.

This rule is intended to implement Iowa Code section 441.17.

[ARC 5288C, IAB 11/18/20, effective 12/23/20; ARC 6025C, IAB 11/3/21, effective 12/8/21; Editorial change: IAC Supplement 11/2/22]

701—102.28(441) Special counsel.

102.28(1) Before the conference board may employ special counsel to assist the city legal department or county attorney under Iowa Code section 441.41, the city legal department in the case of cities having an assessor, or county attorney in the case of counties, shall first provide written approval

of the employment of special counsel for each matter in which the special counsel will be employed on a case-by-case basis.

102.28(2) In the event special counsel is employed, the assessor shall provide the department with written notice of said employment, including the matter being litigated, justification for the hiring of special counsel, and the special counsel's name and hourly rate, within ten days of the hiring. In the event that special counsel has been employed by the conference board as of December 23, 2020, the assessor shall provide the department with written notification of said employment, including the matter being litigated, justification for the hiring of special counsel, and the special counsel's name and hourly rate, within ten days of December 23, 2020, for each case. On or before January 1 of each year, the assessor shall submit to the director, on forms prescribed by the director, a report of all matters litigated by special counsel in the previous 12-month period and the cost of said litigation for each case.

This rule is intended to implement Iowa Code section 441.41 as amended by 2020 Iowa Acts, House File 2641.

[ARC 5288C, IAB 11/18/20, effective 12/23/20; Editorial change: IAC Supplement 11/2/22]

701—102.29(441) Application of two-tier assessment limitation.

102.29(1) Following receipt of the certification of assessment limitations described in Iowa Code section 441.21(9), the county auditor shall determine the assessed values of property by applying the assessment limitations as required under Iowa Code section 441.21(9).

102.29(2) When a property unit of commercial property, industrial property, or property valued by the department pursuant to Iowa Code chapter 434 is comprised of more than one parcel, the county auditor shall apply the assessment limitations described in Iowa Code sections 441.21(5) "b"(2)(a) and 441.21(5) "c"(2)(a), as applicable, to each parcel within the property unit by dividing 150,000 by the value of the entire property unit and multiplying the quotient by the value of each parcel within the property unit. Any remaining value of each parcel within the property unit shall receive the assessment limitations described in Iowa Code sections 441.21(5) "b"(2)(b) and 441.21(5) "c"(2)(b), as applicable. The assessment limitations shall be applied as whole numbers.

EXAMPLE A: Parcels 1, 2, and 3 comprise one property unit of commercial, industrial, or railway property valued at \$300,000 total.

Parcel 1 is assessed at \$100,000.

Parcel 2 is assessed at \$100,000.

Parcel 3 is assessed at \$100,000.

The first \$50,000 of value of each parcel receives the assessment limitation applicable to residential property. The additional value of each parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) "b"(2)(b) and 441.21(5) "c"(2)(b).

EXAMPLE B: Parcels 1, 2, 3, and 4 comprise one property unit of commercial, industrial, or railway property valued at \$850,000 total.

Parcel 1 is assessed at \$500,000.

Parcel 2 is assessed at \$200,000.

Parcel 3 is assessed at \$100,000.

Parcel 4 is assessed at \$50,000.

The first \$88,235 of value of Parcel 1 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) "b"(2)(b) and 441.21(5) "c"(2)(b).

The first \$35,294 of value of Parcel 2 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) "b"(2)(b) and 441.21(5) "c"(2)(b).

The first \$17,647 of value of Parcel 3 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial,

industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) “b”(2)(b) and 441.21(5) “c”(2)(b).

The first \$8,824 of value of Parcel 4 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) “b”(2)(b) and 441.21(5) “c”(2)(b).

EXAMPLE C: Parcels 1 and 2 comprise one property unit of commercial, industrial, or railway property valued at \$500,000 total.

Parcel 1 is assessed at \$400,000.

Parcel 2 is assessed at \$100,000.

The first \$120,000 of value of Parcel 1 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) “b”(2)(b) and 441.21(5) “c”(2)(b).

The first \$30,000 of value of Parcel 2 receives the assessment limitation applicable to residential property. The additional value of the parcel receives the applicable assessment limitation for commercial, industrial, or railway property assessed under Iowa Code chapter 434 described in Iowa Code sections 441.21(5) “b”(2)(b) and 441.21(5) “c”(2)(b).

This rule is intended to implement Iowa Code sections 441.21(5) and 441.21(9) as amended by 2022 Iowa Acts, House File 2552.

[ARC 6525C, IAB 9/21/22, effective 10/26/22; Editorial change: IAC Supplement 11/2/22]

[Filed 5/11/71; amended 8/16/73]

[Filed 6/21/77, Notice 4/6/77—published 7/13/77, effective 8/17/77]

[Filed emergency 7/21/77—published 8/10/77, effective 7/21/77]

[Filed emergency 8/3/79—published 8/22/79, effective 8/3/79]

[Filed emergency 8/1/80—published 8/20/80, effective 8/1/80]

[Filed 3/25/81, Notice 2/18/81—published 4/15/81, effective 5/20/81]

[Filed 5/8/81, Notice 4/1/81—published 5/27/81, effective 7/1/81]

[Filed 3/25/83, Notice 2/16/83—published 4/13/83, effective 5/18/83]

[Filed 7/27/84, Notice 6/20/84—published 8/15/84, effective 9/19/84]

[Filed emergency 8/13/84—published 8/29/84, effective 8/13/84]

[Filed 8/10/84, Notice 7/4/84—published 8/29/84, effective 10/3/84]

[Filed 4/5/85, Notice 1/16/85—published 4/24/85, effective 5/29/85]

[Filed 5/31/85, Notice 4/24/85—published 6/19/85, effective 7/24/85]

[Filed 1/10/86, Notice 12/4/85—published 1/29/86, effective 3/5/86]

[Filed 3/21/86, Notice 2/12/86—published 4/9/86, effective 5/14/86]

[Filed 8/22/86, Notice 7/16/86—published 9/10/86, effective 10/15/86]

[Filed emergency 11/14/86—published 12/17/86, effective 11/14/86]

[Filed 5/15/87, Notice 3/25/87—published 6/3/87, effective 7/8/87]

[Filed 9/18/87, Notice 8/12/87—published 10/7/87, effective 11/11/87]

[Filed 6/10/88, Notice 5/4/88—published 6/29/88, effective 8/3/88]

[Filed 9/2/88, Notice 7/27/88—published 9/21/88, effective 10/26/88]

[Filed 12/7/90, Notice 10/17/90—published 12/26/90, effective 1/30/91]

[Filed 11/18/94, Notice 10/12/94—published 12/7/94, effective 1/11/95]

[Filed 10/6/95, Notice 8/30/95—published 10/25/95, effective 11/29/95]

[Filed 11/15/96, Notice 10/9/96—published 12/4/96, effective 1/8/97]

[Filed 10/17/97, Notice 9/10/97—published 11/5/97, effective 12/10/97]

[Filed 2/12/99, Notice 9/23/98—published 3/10/99, effective 4/14/99]¹

[Filed 1/7/00, Notice 12/1/99—published 1/26/00, effective 3/1/00]

[Filed 9/15/00, Notice 8/9/00—published 10/4/00, effective 11/8/00]

[Filed 12/19/01, Notice 11/14/01—published 1/9/02, effective 2/13/02]

[Filed emergency 2/14/02—published 3/6/02, effective 2/15/02]

[Filed 10/25/02, Notice 9/4/02—published 11/13/02, effective 12/18/02]
[Filed 10/25/02, Notice 9/18/02—published 11/13/02, effective 12/18/02]
[Filed 9/10/04, Notice 8/4/04—published 9/29/04, effective 11/3/04]
[Filed 12/30/05, Notice 11/9/05—published 1/18/06, effective 2/22/06]
[Filed 10/5/06, Notice 8/30/06—published 10/25/06, effective 11/29/06]
[Filed 1/11/07, Notice 11/22/06—published 1/31/07, effective 3/7/07]
[Filed 5/4/07, Notice 3/28/07—published 5/23/07, effective 6/27/07]
[Filed 10/19/07, Notice 9/12/07—published 11/7/07, effective 12/12/07]
[Filed 5/29/08, Notice 4/23/08—published 6/18/08, effective 7/23/08]
[Filed ARC 7726B (Notice ARC 7592B, IAB 2/25/09), IAB 4/22/09, effective 5/27/09]
[Filed ARC 8542B (Notice ARC 8428B, IAB 12/30/09), IAB 2/24/10, effective 3/31/10]
[Filed ARC 8559B (Notice ARC 8352B, IAB 12/2/09), IAB 3/10/10, effective 4/14/10]
[Filed ARC 9478B (Notice ARC 9113B, IAB 10/6/10), IAB 4/20/11, effective 5/25/11]
[Filed ARC 9877B (Notice ARC 9761B, IAB 10/5/11), IAB 11/30/11, effective 1/4/12]
[Filed ARC 0400C (Notice ARC 0286C, IAB 8/22/12), IAB 10/17/12, effective 11/21/12]
[Filed ARC 0770C (Notice ARC 0653C, IAB 3/20/13; Amended Notice ARC 0659C, IAB 4/3/13),
IAB 5/29/13, effective 7/3/13]
[Filed ARC 1196C (Notice ARC 1042C, IAB 10/2/13), IAB 11/27/13, effective 1/1/14]
[Filed ARC 1306C (Notice ARC 1238C, IAB 12/11/13), IAB 2/5/14, effective 3/12/14]
[Filed Emergency ARC 1496C, IAB 6/11/14, effective 5/20/14]
[Filed ARC 1765C (Notice ARC 1593C, IAB 8/20/14), IAB 12/10/14, effective 1/14/15]
[Filed ARC 2108C (Notice ARC 2047C, IAB 6/24/15), IAB 8/19/15, effective 9/23/15]
[Filed ARC 2146C (Notice ARC 2060C, IAB 7/22/15), IAB 9/16/15, effective 10/21/15]
[Filed ARC 2657C (Notice ARC 2519C, IAB 4/27/16), IAB 8/3/16, effective 9/7/16]
[Filed ARC 2707C (Notice ARC 2520C, IAB 4/27/16), IAB 9/14/16, effective 10/19/16]
[Filed ARC 3107C (Notice ARC 2990C, IAB 3/29/17), IAB 6/7/17, effective 7/12/17]
[Filed ARC 3312C (Notice ARC 3203C, IAB 7/19/17), IAB 9/13/17, effective 10/18/17]
[Filed ARC 3771C (Notice ARC 3620C, IAB 2/14/18), IAB 4/25/18, effective 5/30/18]
[Filed ARC 4170C (Notice ARC 4042C, IAB 10/10/18), IAB 12/5/18, effective 1/9/19]
[Filed ARC 5288C (Notice ARC 5182C, IAB 9/23/20), IAB 11/18/20, effective 12/23/20]
[Filed ARC 6025C (Notice ARC 5887C, IAB 9/8/21), IAB 11/3/21, effective 12/8/21]
[Filed ARC 6096C (Notice ARC 5985C, IAB 10/20/21), IAB 12/15/21, effective 1/19/22]
[Filed ARC 6525C (Notice ARC 6429C, IAB 7/27/22), IAB 9/21/22, effective 10/26/22]
[Editorial change: IAC Supplement 11/2/22]
[Filed ARC 6858C (Notice ARC 6601C, IAB 10/19/22), IAB 2/8/23, effective 3/15/23]
[Filed ARC 6958C (Notice ARC 6494C, IAB 9/7/22), IAB 3/22/23, effective 4/26/23]

¹ Amendments nullified by 2000 Iowa Acts, SJR 2005, editorially removed IAC Supplement 7/12/00 pursuant to Iowa Code section 17A.6(3).

CHAPTER 103
EXAMINATION AND CERTIFICATION OF ASSESSORS AND DEPUTY ASSESSORS

[Prior to 12/17/86, Revenue Department[730]]
[Prior to 11/2/22, see Revenue Department[701] Ch 72]

701—103.1(441) Application for examination.

103.1(1) The application for the examination shall be made on a form prescribed by the director and shall constitute an integral part of the examination. The application form shall require information as to the education, training, and experience of the applicant, including evidence of successful completion of the preliminary education requirements required in subrule 72.3(2), and such other information as the director may deem pertinent. Applications must be received by the department at least three days prior to the date of the examination. Applications filed after February 9, 1976, shall be considered public records pursuant to Iowa Code chapter 22 (*City of Dubuque v. Telegraph Herald, Inc.*, 297 N.W.2d 523 (Iowa 1980); 1982 O.A.G. 3).

103.1(2) Upon receipt of a properly filed application, the department shall issue to the applicant a card granting the applicant admission to the examination. No applicant shall be admitted to the examination without presenting the admission card to the examination monitor.

103.1(3) Whenever there occurs a vacancy in the office of assessor, the director shall, upon the written request of the examining board or conference board, forward to the board a copy of any applications requested by either board. When a vacancy occurs in the office of deputy assessor, the director shall, upon the written request of the assessor, forward to the assessor a copy of any applications requested by the assessor.

This rule is intended to implement Iowa Code section 441.5.
[ARC 3838C, IAB 6/6/18, effective 7/11/18; Editorial change: IAC Supplement 11/2/22]

701—103.2(441) Examinations.

103.2(1) Examination questions. Examination questions and answers shall not be made available to persons other than employees of the department authorized by the director to have access to them. Persons who take the examination shall not discuss with anyone the specific questions contained in the examination, nor shall they reveal any specific examination question to another person. This shall not restrict persons who have taken the examination from discussing the general subject matter of the examination.

103.2(2) Materials and supplies. All examination materials shall be furnished by the department and must be returned to the monitor prior to the applicants' leaving the examination room site. During the examination, applicants may be permitted to use their own slide rules or electronic calculators as long as their use does not disturb other applicants. Applicants shall not be permitted to bring any other materials into the examination room, nor shall they be permitted to take any materials from the examination room except their own slide rules or electronic calculators.

103.2(3) Personal conduct during examination. To preserve the integrity of the examinations and the assessing profession, each person taking an examination shall exhibit behavior which is not disruptive to other applicants and no person shall cheat or attempt to cheat on an examination in any manner.

103.2(4) Monitors. The director shall, prior to the examination, provide all applicants with a copy of subrules 72.2(1), 72.2(2), and 72.2(3). Examination monitors shall have the authority to enforce these rules in accordance with subrule 72.2(5).

103.2(5) Violations. Any person who intentionally violates any of the provisions of subrule 72.2(1), 72.2(2), or 72.2(3) shall be subject to the penalties specified in this subrule. If an infraction of subrule 72.2(1), 72.2(2), or 72.2(3) occurs and is detected and confirmed during the examination, the examination of the person committing the infraction shall be confiscated by the monitor and shall not be scored. If the infraction is detected and confirmed after the examination of the person committing the infraction has been scored, the score resulting from that examination shall be reduced to a failing grade and, if necessary, the list of candidates eligible for the position of city or county assessor or deputy assessor shall be adjusted accordingly.

103.2(6) Reserved.

103.2(7) Assessor examination scores. The scores of persons who take the assessor or deputy assessor examination shall be considered public records pursuant to Iowa Code chapter 22.

103.2(8) Reserved.

103.2(9) Length of examination. The director shall determine the appropriate amount of time in which persons may take the examination. Any person who arrives at the examination site after the examination has begun shall not be permitted to complete the examination after the time scheduled for its completion.

103.2(10) Retaking examination. A person who takes the examination for the position of city or county assessor shall not be eligible to take the examination again for a period of at least 30 days following the date the examination was taken, subject also to the restrictions contained in subrule 72.2(5).

103.2(11) Frequency of examination. At the discretion of the director, statewide examinations for the positions of assessor or deputy assessor may be held more than twice a year in Des Moines.

103.2(12) Make-up examination prohibited. Special make-up examinations shall not be held for persons who applied to take the examination for the position of assessor or deputy assessor but who did not for any reason appear at the scheduled examination site.

This rule is intended to implement Iowa Code section 441.5.

[ARC 7726B, IAB 4/22/09, effective 5/27/09; ARC 3313C, IAB 9/13/17, effective 10/18/17; Editorial change: IAC Supplement 11/2/22]

701—103.3(441) Eligibility requirements to take the examination.

103.3(1) High school diploma or its equivalent. Only persons who possess a high school diploma or its equivalent are eligible to take the examination. The equivalent of high school diploma shall consist of a high school equivalency diploma issued by the department of education pursuant to Iowa Code chapter 259A, a similar document issued by the U.S. armed forces, or a similar document issued by another state.

103.3(2) Preliminary education requirements.

a. Only persons who have successfully completed the preliminary education requirements are eligible to take the examination. These requirements may be met by achieving one of the following:

(1) Successful completion of a department-approved course on Iowa assessment and taxation that includes coursework on Iowa laws within the time frame defined in paragraph 72.3(2) “*b*”;

(2) Successful completion of a department-approved course on general appraisal and assessment practice in addition to a department-approved course on Iowa laws. Both courses must be successfully completed within the time frame defined in paragraph 72.3(2) “*b*”; or

(3) Receipt of a currently active department-approved professional appraisal designation from a recognized appraisal organization in conjunction with successful completion of a department-approved course on Iowa laws within the time frame defined in paragraph 72.3(2) “*b*” if the appraisal designation is not already specific to Iowa.

b. All required coursework must be completed within five years prior to the date of the examination.

c. For the purposes of this subrule, “successful completion” shall mean answering a minimum of 70 percent of questions correctly on the test given at the completion of the course.

d. The department will publish a list of approved courses and professional designations on its official website.

This rule is intended to implement Iowa Code section 441.5.

[ARC 3838C, IAB 6/6/18, effective 7/11/18; Editorial change: IAC Supplement 11/2/22]

701—103.4(441) Appraisal-related experience. Appraisal-related experience shall include only such experience as may have been obtained through full-time paid employment consisting of the actual appraisal and valuation of property. The experience shall have included the physical inspection of property as part of the appraisal process and the setting of values for parcels of property.

This rule is intended to implement Iowa Code section 441.5.

[Editorial change: IAC Supplement 11/2/22]

701—103.5(441) Regular certification.

103.5(1) To obtain regular certification, a person must pass the examination and (a) possess two years' appraisal-related experience at the time of passing the examination, or (b) have obtained temporary certification, received a provisional appointment as assessor, and successfully completed the course of study prescribed by the director as provided in Iowa Code section 441.5.

103.5(2) If subsequent to passing the examination a person who has not received a provisional appointment as assessor attains two years' appraisal-related experience, the person must again pass the examination to obtain regular certification.

103.5(3) A regular certificate expires two years after the most recent date certification is granted by the director. However, the regular certificate of a person who receives an appointment as assessor remains valid until the person's resignation or removal from the position of assessor, even though more than two years may have expired since certification was last granted.

103.5(4) A regular certificate may at any time be renewed if the person possessing such a certificate passes the assessor examination. A regular certificate so renewed shall remain valid for a period of two years from the date certification was last granted, except as provided in subrule 72.5(3).

This rule is intended to implement Iowa Code section 441.5.

[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.6(441) Temporary certification.

103.6(1) To obtain temporary certification, a person who does not possess two years' appraisal-related experience must pass the examination for the position of assessor.

103.6(2) The temporary certificate of a person who does not receive a provisional appointment as assessor shall expire two years after the date the certification is granted by the director.

103.6(3) The temporary certificate of a person who does not receive a provisional appointment as assessor may be renewed if the person retakes and passes the assessor examination. A temporary certificate so renewed shall remain valid for a period of two years from the date temporary certification was last granted.

103.6(4) The temporary certificate of a person who receives a provisional appointment as assessor shall expire upon the person's successful completion of the course of study provided in Iowa Code section 441.5 and the granting of regular certification by the director.

103.6(5) The director shall revoke the temporary certificate of a person who receives a provisional appointment as assessor and who does not complete the course of study provided in Iowa Code section 441.5 within 18 months of the person's appointment as assessor. Upon the revocation of an assessor's temporary certificate, the director shall notify the person of the revocation and shall notify the appropriate conference board of the revocation and that the assessor whose temporary certificate has been revoked is no longer eligible to hold the position of assessor.

This rule is intended to implement Iowa Code section 441.5.

[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.7 Reserved.

701—103.8(441) Deputy assessors—regular certification.

103.8(1) A person who passes the examination for assessor or deputy assessor shall be granted regular deputy assessor certification by the director and shall be eligible for appointment to a deputy assessor position.

103.8(2) A deputy assessor regular certificate shall expire two years after the most recent date certification is granted, except as provided in subrule 72.8(3).

103.8(3) The deputy assessor regular certificate of a person who is appointed deputy assessor shall remain valid until the person's resignation or removal from the position of deputy assessor, or until the death, resignation, or removal of the assessor who appointed the person as deputy assessor. However, in the event of the death, resignation, or removal of the assessor, the deputy assessor certificate of the chief deputy shall remain valid until a new assessor is appointed. Nothing contained in this rule shall be construed to relieve a deputy assessor holding a restricted certificate of the continuing education requirements for the retention of the deputy assessor's position as provided in Iowa Code section 441.8.

103.8(4) A deputy assessor regular certificate may at any time be renewed if the person possessing such a certificate passes the assessor or deputy assessor examination. A deputy assessor certificate so renewed shall remain valid for a period of two years from the date certification was last granted, except as provided in subrule 72.8(3).

This rule is intended to implement Iowa Code section 441.5.
[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.9 Reserved.

701—103.10(441) Appointment of deputy assessors.

103.10(1) The appointments of deputy assessors holding regular certificates shall expire upon the death, resignation, or removal of the assessor, except that the appointment of the chief deputy assessor shall not expire until the appointment of a new assessor, nor shall the restricted certificate of a deputy assessor expire at that time.

103.10(2) After the appointment of a new assessor, the assessor may appoint one or more deputy assessors from the registers of persons certified as eligible for appointment as assessor or deputy assessor. The assessor shall notify the director immediately of persons appointed as deputy assessors, the vacating of office by a deputy assessor, or a change in a deputy assessor's legal name.

This rule is intended to implement Iowa Code sections 441.5, 441.10 and 441.11.
[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.11(441) Special examinations. The conference board of the city or county in which a special examination is held shall reimburse the department for all expenses incurred in the administration of the examination. In determining the amount of reimbursement, the director shall take into consideration the costs of traveling to and from the examination site, meals and lodging, if any, for the monitor administering the examination, the costs of preparing and grading the examinations, and the salary of the monitor during the time expended on the examination.

This rule is intended to implement Iowa Code sections 441.5 to 441.7.
[Editorial change: IAC Supplement 11/2/22]

701—103.12(441) Register of eligible candidates.

103.12(1) Assessor and deputy assessor register.

a. Following the administration and grading of an examination for assessor or deputy assessor, the director shall establish updated registers containing the names, in alphabetical order, and addresses of all persons eligible for appointment. The registers shall not contain test scores, but the scores shall be given to the city or county conference board upon request. Eligible candidates shall remain on the register for two years following the date of certification by the director after which time the person must successfully retake the examination to be placed on the register. However, assessors and deputy assessors with six years of consecutive service shall be placed on the register without further testing being required. "Consecutive service" means service in which there was not more than 30 days' break in service. Assessor and deputy assessor service cannot be combined to meet the six-year consecutive service requirement. Assessors and deputy assessors are responsible for maintaining current contact information with the department, including mailing address, email address, and telephone number.

b. In maintaining the register, the department shall indicate which assessors and deputy assessors have retired from the profession. An assessor or deputy assessor may request to no longer be indicated as retired on the register.

c. Deceased assessors and deputy assessors shall be removed from the register.

103.12(2) Continuing education requirements. Assessors and deputy assessors must complete the continuing education requirements provided in Iowa Code sections 441.5 and 441.10 to be reappointed to their present position or appointed to the same position in a different assessing jurisdiction. This provision does not apply to persons not presently serving as an assessor or deputy assessor. It shall be the duty of the conference board in the case of assessor appointments and the duty of the assessor in the case of deputy assessor appointments to receive written verification from the director of continuing

education requirement compliance. An assessor or deputy assessor appointed as such without having complied with continuing education requirements shall be removed from office on order of the director. No continuing education requirements need be met for an assessor to be appointed a deputy assessor nor for a deputy assessor to be appointed an assessor.

This rule is intended to implement Iowa Code sections 441.5 and 441.10.

[Editorial change: IAC Supplement 11/2/22; ARC 6871C, IAB 2/8/23, effective 3/15/23]

701—103.13(441) Course of study for provisional appointees. A person who possesses temporary certification and receives a provisional appointment as assessor shall within 18 months of the appointment complete a course of study prescribed and administered by the department of revenue. The course of study shall include the following: (1) attendance of at least one basic assessment school conducted by the department of revenue; (2) field instruction by appraisal personnel of the department of revenue; (3) the actual appraisal of representative properties in each class of real estate; and (4) attendance at the annual school of instruction sponsored by the department of revenue and the Iowa State Association of Assessors. In the event a person is unable to attend the annual school of instruction due to circumstances beyond the person's control, the director may, upon the request of the person, substitute comparable instruction for the fulfillment of this requirement. At three-month intervals following the appointment of the assessor, department of revenue appraisal personnel shall complete a review of the assessor's performance and discuss the review with the assessor. If the review indicates unsatisfactory progress is being made toward developing a working knowledge of appraisal principles, the assessor shall be informed as to how the assessor's performance could be improved. Not less than 60 nor more than 90 days before the expiration of the 18-month period, the director of revenue shall inform the assessor and the conference board of the assessor's jurisdiction of the director's determination as to whether the assessor satisfactorily completed the course. If the assessor satisfactorily completes the course, the assessor shall be granted regular certification. If the assessor does not satisfactorily complete the course, the director shall revoke the assessor's temporary certificate and notify the assessor and the conference board of the revocation and that the person is no longer eligible to hold the position of assessor.

This rule is intended to implement Iowa Code section 441.5.

[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.14(441) Examining board.

103.14(1) Membership. Each voting unit of the conference board shall appoint a member of the examining board. Members of the examining board shall not be members of the conference board, a body which selects a member of the conference board, or the local board of review (1960 O.A.G. 226). A person must be a resident of the assessing jurisdiction served to qualify for appointment as a member of the examining board. A member changing assessing jurisdiction residency after appointment to the board may continue to serve on the board until the member's current term of office expires.

103.14(2) Terms of members. Members of the examining board shall be appointed for terms of six years. In the event of death, resignation, or removal from office of a member of the examining board, the appropriate voting unit of the conference board shall appoint a successor to serve the unexpired term of the previous incumbent.

103.14(3) Removal of member. A member of an examining board may be removed from office only after specific charges have been filed against the member and a public hearing has been held if requested by the member.

103.14(4) Duties. The examining board may, at its discretion, contact all or some of the persons on the register of candidates eligible for appointment as assessor. The examining board may conduct interviews with interested persons and may administer such further examinations as may enable the board to submit a recommendation to the conference board. In arriving at its recommendation, the examining board may set other professional standards including, but not limited to, examination scores, education, and experience.

103.14(5) Report to conference board. The report to the conference board required pursuant to Iowa Code section 441.6 should contain a complete description of the examining board's investigations and

activities. The report may, at the discretion of the examining board, contain recommendations to the conference board.

103.14(6) *Time for action.* The examining board shall take all steps necessary to comply with the time frames set forth in Iowa Code section 441.6.

This rule is intended to implement Iowa Code sections 441.2, 441.3, 441.4, and 441.6.
[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22]

701—103.15(441) Appointment of assessor.

103.15(1) *Meeting of the conference board.* At the time specified in Iowa Code section 441.6, the conference board shall hold a meeting and take action to appoint an assessor or request permission to hold a special examination. Within ten days of this meeting, the conference board shall notify the director of the appointment or request a special examination. The notice shall include a statement by the conference board stating whether there have been any charges or evidence of any misconduct, nonfeasance, malfeasance, or misfeasance against the appointee. If there have been charges or evidence of any misconduct, nonfeasance, malfeasance, or misfeasance against the appointee, the notice shall include a summary of the misconduct, nonfeasance, malfeasance, or misfeasance and any action taken regarding the misconduct, nonfeasance, malfeasance, or misfeasance. For purposes of this rule, “misconduct” means the same as defined in Iowa Code section 441.9.

103.15(2) *Time for action.* A conference board shall adhere to the time frames specified in Iowa Code section 441.6 in appointing an assessor to fill a vacant position.

103.15(3) *Special examination.* A request for a special examination shall be made only after the conference board has made a good-faith attempt to appoint an assessor from the current register of eligible candidates. The request shall state the reason or reasons the conference board feels the director of revenue should grant permission to hold the special examination.

103.15(4) *Confirmation by the director of revenue.*

a. The appointee selected by the conference board shall not assume the office of city or county assessor until such appointment is confirmed by the director of revenue. In considering whether to confirm the appointment, the director shall consider any charges or evidence of misconduct, nonfeasance, malfeasance, or misfeasance by the appointee. For purposes of this rule, “misconduct” means the same as defined in Iowa Code section 441.9. Within 30 days of receiving the notice contemplated in subrule 72.15(1), the director shall notify the conference board and assessor of the acceptance or rejection of the appointment. An appeal of the director’s decision under this subrule may be made under rule 701—7.37(441).

b. Immediately following selection by the conference board, the appointee assessor shall submit information to the director as required for the director or designee to conduct a background check. The director or designee may review the department’s records and other records in considering whether to confirm the appointment of an assessor.

This rule is intended to implement Iowa Code section 441.6.
[ARC 5288C, IAB 11/18/20, effective 12/23/20; Editorial change: IAC Supplement 11/2/22]

701—103.16(441) Reappointment of assessor.

103.16(1) *Time for reappointment.* A conference board must decide whether to reappoint an incumbent assessor at least 90 days before the expiration of the incumbent’s term. If the incumbent is not to be reappointed, the conference board shall so notify the incumbent in writing at least 90 days before the expiration of the incumbent’s term. Failure of the conference board to provide timely notification of the decision not to reappoint the assessor shall result in the assessor being reappointed. In no case may an incumbent assessor be reappointed earlier than 180 days before the expiration of the incumbent’s term. Within ten days of reappointment or notification of expiration of the incumbent’s term, the conference board shall notify the director of the reappointment or notification of expiration of the incumbent’s term. If the conference board reappoints an incumbent assessor, the notice shall include a statement by the conference board stating whether there have been any charges or evidence of any misconduct, nonfeasance, malfeasance, or misfeasance against the appointee. If there have been charges or evidence of any misconduct, nonfeasance, malfeasance, or misfeasance against the appointee, the

notice shall include a summary of the misconduct, nonfeasance, malfeasance, or misfeasance and any action taken regarding the misconduct, nonfeasance, malfeasance, or misfeasance. For purposes of this rule, “misconduct” means the same as defined in Iowa Code section 441.9.

103.16(2) Continuing education. A conference board shall not reappoint an incumbent assessor if the board has not received from the assessor education advisory committee certification that the incumbent assessor has satisfied all continuing education requirements.

103.16(3) Confirmation by the director of revenue.

a. An assessor reappointed by the conference board shall not assume the office of city or county assessor in the subsequent term until such reappointment is confirmed by the director of revenue. In considering whether to confirm the reappointment, the director shall consider any charges or evidence of misconduct, nonfeasance, malfeasance, or misfeasance by the appointee. For purposes of this rule, “misconduct” means the same as defined in Iowa Code section 441.9. Within 30 days of receiving notice of reappointment by the conference board, the director shall notify the conference board and assessor of the acceptance or rejection of the reappointment. An appeal of the director’s decision under this subrule may be made under rule 701—7.37(441).

b. Immediately following selection by the conference board, the appointee assessor shall submit information to the director as required for the director or designee to conduct a background check. The director or designee may review the department’s records and other records in considering whether to confirm the reappointment of an assessor.

This rule is intended to implement Iowa Code section 441.6 as amended by 2020 Iowa Acts, House File 2641, section 106, and Iowa Code section 441.8.

[ARC 5288C, IAB 11/18/20, effective 12/23/20; Editorial change: IAC Supplement 11/2/22]

701—103.17(441) Removal of assessor. An assessor may be removed from office for the reasons stated in Iowa Code section 441.9, but only after the charges have been substantiated.

This rule is intended to implement Iowa Code section 441.9.

[Editorial change: IAC Supplement 11/2/22]

701—103.18(421,441) Courses offered by the department of revenue.

103.18(1) Class size. The director may determine the maximum number of students for a particular class in order to maintain a suitable learning environment. Applications to take a course shall be accepted in the order in which they are received by the department. If the number of applications received as of a specific mail delivery results in the receipt of more applications than there are spaces for the class, those applications received in that mail delivery shall be subject to a drawing by lot to determine those which shall be accepted for the class. However, persons who are not currently serving as assessors or deputy assessors shall not be admitted to a course ahead of persons serving as assessors or deputy assessors, regardless of the date on which their applications were received.

103.18(2) Examinations during the course. Examination questions and answers shall not be made available to persons other than employees of the department authorized by the director to have access to such information. Persons who take the examination shall not discuss with anyone the specific questions contained in the examination, nor shall they reveal any specific examination question to another person. This shall not restrict persons who have taken a course examination from discussing the general subject matter of the examination.

103.18(3) Materials and supplies. All examination materials shall be furnished by the department and must be returned to the monitor prior to the students leaving the examination. During the examination, students may be permitted to use their own slide rules or electronic calculators as long as their use does not disturb other students. Students shall not be permitted to bring any other materials into the examination room, nor shall they be permitted to take any materials from the examination room except their own slide rules or electronic calculators.

103.18(4) Personal conduct during course and examination. To preserve the integrity of the examinations and the assessing profession, each person taking an examination shall not exhibit behavior which is disruptive to other persons taking the examination, nor shall a person cheat or attempt to cheat on an examination in any manner.

103.18(5) *Violations.* Any person who intentionally violates any of the provisions of subrule 72.18(2), 72.18(3), or 72.18(4) shall be subject to the penalties specified in this subrule. If an infraction of subrule 72.18(2), 72.18(3), or 72.18(4) occurs and is detected and confirmed during the examination, the examination of the person committing the infraction shall be confiscated by the instructor and shall not be scored. If the infraction is detected and confirmed after the examination of the person committing the infraction has been scored, the score resulting from that examination shall be reduced to a failing grade and the director shall notify the assessor education advisory committee of the action taken. If the infraction is detected and confirmed during the course, the instructor shall expel the student from the classroom, and the student shall not be permitted to take the examination for the course.

103.18(6) *Instructors.* Course instructors shall inform all students of the provisions of subrules 72.18(2), 72.18(3), and 72.18(4). The instructors shall have the authority to enforce these rules in accordance with subrule 72.18(5).

103.18(7) *Retaking examination.* A person who receives a failing score on the examination for a course may retake the examination by submitting a request to the director within ten days of the date the director notifies the person of the examination score. The examination shall be retaken at the office of the department in Des Moines or at the site of any scheduled course examination, and shall be retaken within 30 days of the date the original examination was taken. A person who retakes an examination may not again take that particular course for credit until at least 30 days have passed from the date the examination was retaken. A special examination may be taken only once for a particular course, regardless of the number of times a student takes the course. A special examination shall be given only if the student took and failed the examination given at the end of a course taken for credit.

103.18(8) *Review of examination.* Persons who have taken a course examination may, after presenting proper identification, review their examinations in the office of the department's local government services division within 60 days after the date the examination has been administered. The review shall consist only of examining the person's own answer sheet and the question book. Persons reviewing their examinations shall not be permitted to take notes or otherwise transcribe information during this review, nor shall they have access to the answers to questions contained in the examination. Persons who review their examinations shall be permitted to do so only once, and shall not be eligible to take the same examination for a period of at least 30 days following the date of the review of the examinations.

103.18(9) *Length of examination.* The director shall determine the appropriate amount of time in which persons may take each examination. Any person who arrives at the examination site after the examination has begun shall not be permitted to complete the examination after the time scheduled for completion.

This rule is intended to implement Iowa Code section 441.8.

[ARC 7726B, IAB 4/22/09, effective 5/27/09; Editorial change: IAC Supplement 11/2/22; ARC 6958C, IAB 3/22/23, effective 4/26/23]

[Filed emergency 1/23/76—published 2/9/76, effective 2/9/76]

[Filed 3/15/76, Notice 2/9/76—published 4/5/76, effective 5/10/76]

[Filed 4/14/78, Notice 2/22/78—published 5/3/78, effective 6/7/78]

[Filed emergency 8/1/80—published 8/20/80, effective 8/1/80]

[Filed 1/30/81, Notice 12/24/80—published 2/18/81, effective 3/25/81]

[Filed 9/23/82, Notice 8/18/82—published 10/13/82, effective 11/17/82]

[Filed 2/8/85, Notice 11/21/84—published 2/27/85, effective 4/3/85]

[Filed emergency 11/14/86—published 12/17/86, effective 11/14/86]

[Filed 9/2/88, Notice 7/27/88—published 9/21/88, effective 10/26/88]

[Filed 10/27/89, Notice 9/20/89—published 11/15/89, effective 12/20/89]

[Filed 12/19/01, Notice 11/14/01—published 1/9/02, effective 2/13/02]

[Filed 12/30/05, Notice 11/9/05—published 1/18/06, effective 2/22/06]

[Filed ARC 7726B (Notice ARC 7592B, IAB 2/25/09), IAB 4/22/09, effective 5/27/09]

[Filed ARC 3313C (Notice ARC 3206C, IAB 7/19/17), IAB 9/13/17, effective 10/18/17]

[Filed ARC 3838C (Notice ARC 3725C, IAB 4/11/18), IAB 6/6/18, effective 7/11/18]

[Filed ARC 5288C (Notice ARC 5182C, IAB 9/23/20), IAB 11/18/20, effective 12/23/20]

[Editorial change: IAC Supplement 11/2/22]

[Filed ARC 6871C (Notice ARC 6747C, IAB 12/14/22), IAB 2/8/23, effective 3/15/23]

[Filed ARC 6958C (Notice ARC 6494C, IAB 9/7/22), IAB 3/22/23, effective 4/26/23]

CHAPTER 111
ADMINISTRATION

[Prior to 12/17/86, Revenue Department[730]]
[Prior to 11/2/22, see Revenue Department[701] Ch 122]

701—111.1(441) Establishment. Iowa Code section 441.8 established a program of continuing education to be developed and administered by the director of revenue, hereinafter referred to as the director. To administer the program, the director has established an assessor education advisory committee, hereinafter referred to as the committee.

This rule is intended to implement Iowa Code section 441.8.
[Editorial change: IAC Supplement 11/2/22]

701—111.2(441) General operation. The chairperson of the committee shall be the director. The director shall appoint to the committee a representative of the local government services division of the department of revenue and two assessor representatives. The assessor representatives shall serve four-year staggered terms. To initiate the staggered-term policy, one assessor shall serve through December 31, 2009, and the other assessor shall serve through December 31, 2011. The committee will meet at least once each year.

This rule is intended to implement Iowa Code section 441.8.
[Editorial change: IAC Supplement 11/2/22; ARC 6958C, IAB 3/22/23, effective 4/26/23]

701—111.3(441) Location. Persons may obtain information about the committee and its activities at the Department of Revenue in the Hoover State Office Building, Des Moines, Iowa 50319. Persons wishing to obtain information or make submissions should address their correspondence to that address.

This rule is intended to implement Iowa Code section 441.8.
[Editorial change: IAC Supplement 11/2/22]

701—111.4(441) Purpose. The committee is established to assist the director in developing and administering a program of continuing education for Iowa assessors and deputy assessors. The program will emphasize assessment and appraisal procedures, assessment laws, rights and responsibilities of taxpayers and property owners related to the assessment of property for taxation, duties of assessors and deputy assessors, and other matters related to the positions of assessor and deputy assessor. The director, with the assistance of the committee, will designate the courses to be offered in the program, the content of the courses, and the number of hours of classroom instruction for each course. An evaluation of the program will be conducted at least annually with any necessary changes made.

The director shall certify those assessors and deputy assessors who have received sufficient credit to be eligible for reappointment to their present position.

This rule is intended to implement Iowa Code section 441.8.
[Editorial change: IAC Supplement 11/2/22]

[Filed 5/25/79, Notice 4/18/79—published 6/13/79, effective 7/18/79]

[Filed 9/14/79, Notice 8/8/79—published 10/3/79, effective 11/7/79]

[Filed emergency 8/1/80—published 8/20/80, effective 8/1/80]

[Filed 11/7/80, Notice 10/1/80—published 11/26/80, effective 12/31/80]

[Filed emergency 11/14/86—published 12/17/86, effective 11/14/86]

[Filed 6/22/90, Notice 5/16/90—published 7/11/90, effective 8/15/90]

[Filed 12/19/01, Notice 11/14/01—published 1/9/02, effective 2/13/02]

[Filed 12/30/05, Notice 11/9/05—published 1/18/06, effective 2/22/06]

[Filed 2/8/08, Notice 1/2/08—published 2/27/08, effective 4/2/08]

[Editorial change: IAC Supplement 11/2/22]

[Filed ARC 6958C (Notice ARC 6494C, IAB 9/7/22), IAB 3/22/23, effective 4/26/23]