

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,147,272C) 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

CHAPTER 2

PHARMACIST LICENSES

- 2.1(147,155A) Licensure by examination
- 2.2(155A) Application for examination—requirements
- 2.3(147,155A) Examination fee
- 2.4(155A) Internship requirements
- 2.5(155A) College graduate certification
- 2.6(147) Reexamination applications and fees
- 2.7(147) Examination results
- 2.8(155A) Transfer of examination scores
- 2.9(147,155A) Licensure by license transfer/reciprocity
- 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(155A) Fees for additional license certificates
- 2.15(155A) Notifications to the board
- 2.16(235B,272C) Mandatory training for identifying and reporting abuse

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 form
- 3.9(155A) Registration term and renewal
- 3.10(155A) Registration fee
- 3.11(155A) Late applications and fees
- 3.12(155A) Registration certificates
- 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 technical functions
3.22(155A)	Technical functions
3.23(155A)	Tasks a pharmacy technician shall not perform
3.24(155A)	New prescription drug orders or medication orders
3.25(155A)	Delegation of nontechnical functions
3.26 and 3.27	Reserved
3.28(147,155A)	Unethical conduct or practice
3.29(155A)	Denial of registration
3.30(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)	Discipline of pharmacist-interns

CHAPTER 5 Reserved

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 other items
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 and 7.5	Reserved
7.6(124,155A)	Security
7.7(155A)	Verification by pharmacist when pharmacy is closed
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 Reserved
- 7.12(124,126,155A) Drugs dispensed to patients as a result of an emergency room visit
- 7.13(124,155A) Records

CHAPTER 8

UNIVERSAL PRACTICE STANDARDS

- 8.1(155A) Purpose and scope
- 8.2(155A) Pharmaceutical care
- 8.3(155A) Responsibility
- 8.4(155A) Pharmacist identification
- 8.5(155A) Environment and equipment requirements
- 8.6(155A) Health of personnel
- 8.7(155A) Procurement, storage, and recall of drugs and devices
- 8.8(124,155A) Out-of-date drugs or devices
- 8.9(124,155A) Records
- 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 pharmacy technicians
- 8.15(155A) Delivery of prescription drugs and devices
- 8.16(124,155A) Confidential information
- 8.17 and 8.18 Reserved
- 8.19(124,126,155A) 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 to 8.25 Reserved
- 8.26(155A) Continuous quality improvement program
- 8.27 to 8.31 Reserved
- 8.32(124,155A) Individuals qualified to administer
- 8.33(147,155A) Supervision of pharmacists who administer adult immunizations
- 8.34(155A) Collaborative drug therapy management
- 8.35(155A) Pharmacy license

CHAPTER 9

AUTOMATED MEDICATION DISTRIBUTION SYSTEMS AND TELEPHARMACY SERVICES

- 9.1(155A) Purpose and scope
- 9.2(147,155A) Definitions
- 9.3(147,155A) Pharmacist in charge responsibilities
- 9.4 Reserved
- 9.5(124,155A) General requirements for telepharmacy
- 9.6(155A) Duties of pharmacist in telepharmacy practice
- 9.7 to 9.9 Reserved
- 9.10(147,155A) Quality assurance and performance improvement
- 9.11(147,155A) Policies and procedures
- 9.12(147,155A) System, site, and process requirements
- 9.13(147,155A) Records
- 9.14 Reserved
- 9.15(147,155A) Decentralized unit dose AMDS

- 9.16(147,155A) Centralized unit dose AMDS
- 9.17(147,155A) Outpatient AMDS
- 9.18(124,155A) Remote dispensing site operations
- 9.19 Reserved
- 9.20(124,155A) Drugs at a remote dispensing site
- 9.21(124,155A) Record keeping

CHAPTER 10 CONTROLLED SUBSTANCES

- 10.1(124) Who shall register
- 10.2(124) Application forms
- 10.3(124) Registration and renewal
- 10.4(124) Exemptions—registration fee
- 10.5(124) Separate registration for independent activities; coincident activities
- 10.6(124) Separate registrations for separate locations; exemption from registration
- 10.7 to 10.9 Reserved
- 10.10(124,147,155A) Inspection
- 10.11(124) Modification or termination of registration
- 10.12(124) Denial, modification, suspension, or revocation of registration
- 10.13 and 10.14 Reserved
- 10.15(124,155A) Security requirements
- 10.16(124) Report of theft or loss
- 10.17(124) Accountability of stock supply
- 10.18(124) Disposal
- 10.19 and 10.20 Reserved
- 10.21(124,126,155A) Prescription requirements
- 10.22(124) Schedule II emergency prescriptions
- 10.23(124) Schedule II prescriptions—partial filling
- 10.24(124) Schedule II medication order
- 10.25 and 10.26 Reserved
- 10.27(124,155A) Facsimile transmission of a controlled substance prescription
- 10.28(124,155A) Schedule III, IV, or V refills
- 10.29(124,155A) Schedule III, IV, or V partial fills
- 10.30(124,155A) Schedule III, IV, and V medication order
- 10.31(124,155A) Dispensing Schedule V controlled substances without a prescription
- 10.32(124,155A) Dispensing products containing ephedrine, pseudoephedrine, or phenylpropanolamine
- 10.33(124,155A) Schedule II perpetual inventory in pharmacy
- 10.34(124,155A) Records
- 10.35(124,155A) Physical count and record of inventory
- 10.36(124) Samples and other complimentary packages—records
- 10.37(124,126) Revision of controlled substances schedules
- 10.38(124) Temporary designation of controlled substances
- 10.39(124,126) Excluded substances
- 10.40(124,126) Anabolic steroid defined

CHAPTER 11 DRUGS IN EMERGENCY MEDICAL SERVICE PROGRAMS

- 11.1(124,147A,155A) Definitions
- 11.2(124,147A,155A) Ownership of drugs—options
- 11.3(124,147A,155A) General requirements
- 11.4(124,147A,155A) Procurement and storage

- 11.5(124,147A,155A) Records
- 11.6(124,147A,155A) Inspections
- 11.7(124,147A,155A) Security and control

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
STERILE COMPOUNDING PRACTICES

- 13.1(124,126,155A) Purpose and scope
- 13.2(124,126,155A) Definitions
- 13.3(155A) Responsibilities
- 13.4 Reserved
- 13.5(155A) References required
- 13.6(126,155A) Policies and procedures
- 13.7(126,155A) Labeling requirements
- 13.8 and 13.9 Reserved
- 13.10(126,155A) Microbial contamination risk levels
- 13.11(155A) Low-risk preparations and low-risk preparations with 12-hour or less beyond-use date
- 13.12(155A) Medium-risk preparations
- 13.13(155A) High-risk preparations
- 13.14(155A) Immediate-use preparations
- 13.15(155A) Utilization of single-dose and multiple-dose containers
- 13.16(155A) Utilization of proprietary bag and vial systems
- 13.17 to 13.19 Reserved
- 13.20(124,155A) Sterile preparation of hazardous drugs
- 13.21 and 13.22 Reserved
- 13.23(124,155A) Verification of compounding accuracy and sterility
- 13.24(124,155A) Sterilization methods
- 13.25(155A) Media-fill testing by personnel
- 13.26 Reserved
- 13.27(124,126,155A) Physical environment requirements
- 13.28(155A) Cleaning, maintenance, and supplies
- 13.29(126,155A) Environmental monitoring requirements
- 13.30 Reserved
- 13.31(155A) Quality assurance (QA)
- 13.32(155A) Patient or caregiver education and training
- 13.33(124,155A) Storage and delivery of sterile preparations

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 FACILITY PHARMACY PRACTICE

15.1(155A)	Purpose and scope
15.2(126,155A)	Definitions
15.3(155A)	Pharmacist in charge
15.4(155A)	Reference library
15.5(124,155A)	Security
15.6 and 15.7	Reserved
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)	General requirements for qualified nuclear pharmacist
16.4(155A)	General requirements for pharmacies providing radiopharmaceutical services
16.5(155A)	Library
16.6(155A)	Minimum equipment requirements

CHAPTER 17

WHOLESALE DRUG LICENSES

17.1(155A)	Definitions
17.2	Reserved
17.3(155A)	Wholesale drug license
17.4(155A)	Minimum qualifications
17.5(155A)	Personnel
17.6(155A)	Responsibility for conduct
17.7(124,155A)	Distribution to authorized licensees
17.8(124,155A)	Written policies and procedures
17.9(155A)	Facilities
17.10(124,155A)	Security
17.11(155A)	Storage
17.12	Reserved
17.13(155A)	Drugs in possession of representatives
17.14(155A)	Examination of materials

- 17.15(155A) Returned, damaged, and outdated prescription drugs
- 17.16(124,155A) Record keeping
- 17.17(124,155A) Compliance with federal, state, and local laws
- 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) Application and license requirements
- 19.3(124,155A) Applicability of board rules
- 19.4 to 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) Discipline

CHAPTER 20

PHARMACY COMPOUNDING PRACTICES

- 20.1(124,126,155A) Purpose and scope
- 20.2(124,126,155A) Definitions
- 20.3(124,126,155A) General requirements
- 20.4(126,155A) Organization and personnel
- 20.5(126,155A) Drug compounding facilities
- 20.6(126,155A) Sterile products and radiopharmaceuticals
- 20.7 Reserved
- 20.8(126,155A) Equipment
- 20.9(126,155A) Control of bulk drug substances, components, containers, and closures
- 20.10(124,126,155A) Drug compounding controls
- 20.11(126) Bulk compounding
- 20.12(124,126,155A) Records

CHAPTER 21

ELECTRONIC DATA IN PHARMACY PRACTICE

- 21.1(124,155A) Definitions
- 21.2(124,155A) System security and safeguards
- 21.3(124,155A) Verifying authenticity of an electronically transmitted prescription
- 21.4(124,155A) Automated data processing system
- 21.5(124,155A) Pharmacist verification of controlled substance refills—daily printout or logbook
- 21.6 Reserved
- 21.7(124,155A) Electronically prepared prescriptions
- 21.8(124,155A) Computer-to-computer transmission of a prescription

- 21.9(124,155A) Facsimile transmission (fax) of a prescription
- 21.10 and 21.11 Reserved
- 21.12(124,155A) Prescription drug orders for Schedule II controlled substances
- 21.13(124,155A) Prescription drug orders for Schedule II controlled substances—emergency situations
- 21.14(124,155A) Facsimile transmission of a prescription for Schedule II narcotic substances—parenteral
- 21.15(124,155A) Facsimile transmission of Schedule II controlled substances—long-term care facility patients
- 21.16(124,155A) Facsimile transmission of Schedule II controlled substances—hospice patients

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

LONG-TERM CARE PHARMACY PRACTICE

- 23.1(155A) Definitions
- 23.2(124,155A) Applicability of rules
- 23.3(124,155A) Freedom of choice
- 23.4(124,155A) Pharmacy responsibilities
- 23.5(124,155A) Emergency drugs
- 23.6(124,155A) Space, equipment, and supplies
- 23.7(124,155A) Policies and procedures
- 23.8 Reserved
- 23.9(124,155A) Medication orders
- 23.10(124,155A) Stop orders
- 23.11(124,155A) Drugs dispensed—general requirements
- 23.12 Reserved
- 23.13(124,155A) Labeling drugs under special circumstances
- 23.14(124,155A) Labeling of biologicals and other injectables supplied to a facility
- 23.15(124,155A) Return and reuse of drugs and devices
- 23.16(124,155A) Destruction of outdated and improperly labeled drugs
- 23.17(124,155A) Accountability of controlled substances
- 23.18(124,155A) Schedule II orders
- 23.19(124,155A) Dispensing Schedule II controlled substances
- 23.20(124,155A) Partial filling of Schedule II controlled substances
- 23.21(124,155A) Destruction of 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
(Uniform Rules)

28.1(17A)	Applicability
28.2(17A)	Advice on possible rules before notice of proposed rule adoption
28.3(17A)	Public rule-making docket
28.4(17A)	Notice of proposed rule making
28.5(17A)	Public participation
28.6(17A)	Regulatory analysis
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)	Contents, style, and form of rule
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)	General statements of policy
28.17(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
IMPAIRED PHARMACY PROFESSIONAL
AND TECHNICIAN RECOVERY PROGRAM

- 30.1(155A) Definitions
- 30.2(155A) Purpose, function, and responsibilities
- 30.3(155A) Program committee and personnel; confidentiality; liability
- 30.4(155A) Identification and referral of impaired professionals and technicians
- 30.5(155A) Recovery contract requirements
- 30.6(155A) Program provider contract
- 30.7(155A) Disclosure of information
- 30.8(155A) Program funds

CHAPTER 31
STUDENT LOAN DEFAULT OR NONCOMPLIANCE
WITH AGREEMENT FOR PAYMENT OF OBLIGATION

- 31.1(261) Definitions
- 31.2(261) Issuance or renewal of a license—denial
- 31.3(261) Suspension or revocation of a license
- 31.4(17A,22,261) Share information

CHAPTERS 32 and 33
Reserved

CHAPTER 34
RULES FOR WAIVERS AND VARIANCES

- 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 or variance
- 34.5(17A,124,126,147,155A,205,272C) Filing of petition
- 34.6(17A) Content of petition
- 34.7(17A) Additional information
- 34.8(17A) Notice
- 34.9(17A) Hearing procedures
- 34.10(17A) Ruling
- 34.11(17A,22) Public availability
- 34.12(17A) Summary reports
- 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 Reserved
- 35.5(17A,124B,126,147,155A,205,272C) Notice of hearing
- 35.6(17A,272C) Presiding officer for nondisciplinary hearings
- 35.7(17A,124B,147,155A,272C) Waiver of procedures
- 35.8(17A,272C) Telephone or network proceedings
- 35.9(17A) Disqualification
- 35.10(17A,272C) Consolidation—severance
- 35.11(17A,272C) Service and filing of pleadings and other papers
- 35.12(17A,272C) Discovery
- 35.13(17A,272C) Subpoenas
- 35.14(17A,272C) Motions
- 35.15(17A,272C) Prehearing conference
- 35.16(17A,272C) Continuances
- 35.17(17A) Withdrawals
- 35.18 Reserved
- 35.19(17A,124B,126,147,155A,205,272C) Hearing procedures in contested cases
- 35.20(17A,272C) Evidence
- 35.21(17A,272C) Default
- 35.22(17A,272C) Ex parte communication
- 35.23(17A,272C) Recording costs
- 35.24(17A,272C) Interlocutory appeals
- 35.25(17A) Final decision
- 35.26(17A,124B,126,147,155A,205,272C) Appeals and review
- 35.27(17A,124B,126,147,155A,205,272C) Applications for rehearing
- 35.28(17A,272C) Stays of board actions
- 35.29(17A,272C) No factual dispute contested cases
- 35.30(17A,124B,126,147,155A,205,272C) Emergency adjudicative proceedings

CHAPTER 36
DISCIPLINE

- 36.1(147,155A,272C) Authority and grounds for discipline
- 36.2(155A,272C) Investigations
- 36.3(147,272C) Peer review committees
- 36.4(17A,124,124B,126,147,155A,272C) Disciplinary proceedings
- 36.5(17A,124,124B,126,147,155A,272C) Notice of disciplinary hearing
- 36.6(17A,124B,147,155A,272C) Informal settlement
- 36.7(272C) Appearance
- 36.8(17A,124B,147,155A,272C) Order of proceedings
- 36.9(272C) Confidentiality
- 36.10(17A,272C) Notification of decision
- 36.11(272C) Board decision
- 36.12(17A,272C) Publication of decisions
- 36.13(17A,124B,147,155A,272C) Reinstatement
- 36.14(17A,124B,147,155A,272C) Informal reinstatement conference
- 36.15(17A,124B,147,155A,272C) Voluntary surrender of a license, permit, or registration
- 36.16(17A,124B,147,155A,272C) License, permit, or registration denial

36.17(155A,272C) Order for mental or physical examination
36.18(272C) Disciplinary hearings—fees and costs