

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
- 2.17(272C) Continuing professional development portfolio

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(155A) Extension of deadline for national certification
- 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 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 to 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 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
5.10(155A)	Registration renewal
5.11(155A)	Late application
5.12	Reserved
5.13(155A)	Registration certificates
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(155A)	Nontechnical pharmacy support tasks
5.19	Reserved
5.20(155A)	Training and utilization of pharmacy support persons
5.21(155A)	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) 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 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 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) Pharmaceutical care
- 8.3(155A) Responsibility
- 8.4(155A) Pharmacist identification and staff logs
- 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 or pharmacy support persons
- 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(155A)	Vaccine administration by pharmacists
8.34(155A)	Collaborative drug therapy management
8.35(155A)	Pharmacy license
8.36 to 8.39	Reserved
8.40(155A,84GA,ch63)	Pharmacy pilot or demonstration research projects

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(124)	Schedule II—issuing multiple prescriptions
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 without a prescription
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
10.41(124A)	Designation of imitation controlled substances

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)	Written agreement
11.4(124,147A,155A)	Termination of services
11.5	Reserved
11.6(124,147A,155A)	Registration required
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 the written protocol
11.16(124,147A,155A)	Administration of Schedule II controlled substances—pharmacy-based service
11.17 and 11.18	Reserved
11.19(124,147A,155A)	Patient care reports
11.20(124,147A,155A)	Prescription drugs in EMS 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
- 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) Destruction or disposal 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 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(155A) Record requirements
- 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 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 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) 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
 16.7(155A) Training and utilization of pharmacy support persons

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 prepared or electronically or fax 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) Electronic 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) Facsimile transmission of a prescription 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

PHARMACY INTERNET SITES

24.1(155A)	Purpose and scope
24.2(155A)	Definitions
24.3(155A)	General requirements for Internet pharmacy
24.4 and 24.5	Reserved
24.6(155A)	Prescription requirements
24.7(155A)	Internet site registration
24.8(155A)	Internet site information
24.9 and 24.10	Reserved
24.11(155A)	Records
24.12(155A)	Pharmacy liability
24.13(155A)	Application denial
24.14(155A)	Discipline

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

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

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

CHAPTER 37 IOWA PRESCRIPTION MONITORING PROGRAM

- 37.1(124) Purpose
- 37.2(124) Definitions
- 37.3(124) Requirements for the PMP
- 37.4(124) Access to database information
- 37.5(124) Fees
- 37.6(124) PMP information retained
- 37.7(124) Information errors
- 37.8(124) Dispenser and practitioner records
- 37.9(124) Prohibited acts

CHAPTERS 38 and 39 Reserved

CHAPTER 40 TECH-CHECK-TECH PROGRAMS

- 40.1(155A) Purpose and scope
- 40.2(155A) Definitions
- 40.3(155A) General requirements
- 40.4(155A) TCT program requirements

CHAPTERS 41 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