

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 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 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)	Responsible parties
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)	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

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
Reserved

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) 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
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

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)	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
MILITARY SERVICE AND VETERAN RECIPROCITY

33.1(85GA,ch1116)	Definitions
33.2(85GA,ch1116)	Military education, training, and service credit
33.3(85GA,ch1116)	Veteran licensure or registration
33.4(85GA,ch1116)	Request for contested case

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