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
- Responsibilities 1.3(17A,272C)
- 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(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 technic	eians

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

	FIIAKMACT SUFFORT FERSONS
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

	UENERAL FHARMACT F
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(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(147,155A)	Supervision of pharmacists who administer adult immunizations
8.34(155A)	Collaborative drug therapy management
0.05(1.55.4)	

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
0.21(124.155A)	Description of the second seco

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(124,147,1554	A) 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

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

	Controlled substances annual inventory
11.31	Reserved
	Destruction or disposal of controlled substances
	Report of loss or theft of controlled substance
11.34(124,147A,155A)	Records

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(124,126,155)	
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

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

- 15.2(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

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

ELECTRONIC DATA IN PHARMACY PRACTICE

21.1(124.155A)	
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
01 4(104 155 4)	
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

PHARMACY INTERNET SITES

Purpose and scope 24.1(155A) 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 Internet site information 24.8(155A) 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

AGENCY PROCEDURE FOR RULE MAKING

(Uniform Rules)

- 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

Reserved

CHAPTER 34

RULES FOR WAIVERS AND VARIANCES

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
- 25.25(17A) Final desistant
- 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