

## 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 devices
6.16(124,155A)	Records

CHAPTER 7  
HOSPITAL PHARMACY PRACTICE

7.1(155A)	Purpose and scope
7.2(155A)	Pharmacist in charge
7.3(155A)	Reference library
7.4 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 8.30	Reserved
8.31(135,147A)	Opioid antagonist dispensing by pharmacists by standing order
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

Reserved

## CHAPTER 10

## CONTROLLED SUBSTANCES

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

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

## CHAPTER 11

## DRUGS IN EMERGENCY MEDICAL SERVICE PROGRAMS

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

CHAPTER 12  
PRECURSOR SUBSTANCES

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

CHAPTER 13  
TELEPHARMACY PRACTICE

13.1(155A)	Purpose and scope
13.2(155A)	Definitions
13.3(124,155A)	Written agreement
13.4(155A)	Responsible parties
13.5 to 13.7	Reserved
13.8(124,155A)	General requirements for telepharmacy site
13.9(155A)	General requirements for managing pharmacy
13.10(155A)	General requirements for verifying pharmacist
13.11(155A)	General requirements for telepharmacy technician
13.12 to 13.15	Reserved
13.16(124,155A)	Telepharmacy site—initial application
13.17(124,155A)	Changes to telepharmacy site or managing pharmacy
13.18(155A)	Opening of traditional pharmacy
13.19 and 13.20	Reserved
13.21(124,155A)	Policies and procedures
13.22(155A)	Reports to the board
13.23(124,155A)	Records

CHAPTER 14  
PUBLIC INFORMATION AND INSPECTION OF RECORDS

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

CHAPTER 15  
CORRECTIONAL PHARMACY PRACTICE

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

CHAPTER 16  
NUCLEAR PHARMACY PRACTICE

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

CHAPTER 17  
WHOLESALE 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)	Nonresident pharmacy license
19.3(155A)	Registered pharmacist in charge
19.4(124,155A)	Applicability of board rules
19.5 and 19.6	Reserved
19.7(155A)	Confidential data
19.8(124,155A)	Storage and shipment of drugs and devices
19.9(155A)	Patient record system, prospective drug use review, and patient counseling
19.10(155A)	Reporting discipline and criminal convictions
19.11(155A)	Discipline

## CHAPTER 20

## COMPOUNDING PRACTICES

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

## CHAPTER 21

## ELECTRONIC DATA AND AUTOMATED SYSTEMS IN PHARMACY PRACTICE

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

## CHAPTER 22

## UNIT DOSE, ALTERNATIVE PACKAGING, AND EMERGENCY BOXES

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

## CHAPTER 23

## 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)	Disposal of previously dispensed controlled substances

## CHAPTER 24

Reserved

## CHAPTER 25

## CHILD SUPPORT NONCOMPLIANCE

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

## CHAPTER 26

## PETITIONS FOR RULE MAKING

(Uniform Rules)

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

CHAPTER 27  
DECLARATORY ORDERS  
(Uniform Rules)

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

CHAPTER 28  
AGENCY PROCEDURE FOR RULE MAKING

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

CHAPTER 29  
SALES OF GOODS AND SERVICES

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

CHAPTER 30  
IOWA MONITORING PROGRAM FOR PHARMACY PROFESSIONALS

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

## CHAPTER 31

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 and providing notice
- 34.8 and 34.9 Reserved
- 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(17A) Applicability of Iowa Rules of Civil Procedure
- 35.5(17A,272C) Combined statement of charges and settlement agreement
- 35.6(17A,124B,126,147,155A,205,272C) Notice of hearing
- 35.7(17A,272C) Statement of charges
- 35.8(13,272C) Legal representation
- 35.9(17A,272C) Presiding officer in a disciplinary contested case
- 35.10(17A,272C) Presiding officer for nondisciplinary hearings
- 35.11(17A,124B,147,155A,272C) Waiver of procedures

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

#### CHAPTER 36 DISCIPLINE

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

#### 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

CHAPTER 41  
OUTSOURCING FACILITIES

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

CHAPTERS 42 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