

## 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,147,155A,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
- 1.8(155A) Alternate board members

### CHAPTER 2

#### PHARMACIST LICENSES

- 2.1(147,155A) Purpose and scope
- 2.2(147,155A) Definitions
- 2.3(147,155A) License and criminal history record check required
- 2.4(147,155A) Licensure by examination
- 2.5(155A) Application for examination—requirements
- 2.6(155A) Internship requirements
- 2.7(147) Reexamination applications and fees
- 2.8(155A) Licensure by score transfer
- 2.9(147,155A) Licensure by license transfer
- 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(147,155A) Fees for additional license certificates and verification
- 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
- 2.18(147,155A) Temporary license

### 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
- 3.9(155A) Registration fee and term—technician trainee
- 3.10(155A) Registration fee, term, and renewal—certified pharmacy technician
- 3.11(155A) Verification fee
- 3.12 Reserved
- 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	Reserved
3.23(155A)	Functions a pharmacy technician shall not perform
3.24 to 3.27	Reserved
3.28(147,155A)	Unethical conduct or practice
3.29(155A)	Denial of registration
3.30(155A)	Reporting discipline and criminal convictions
3.31(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)	Notifications to the board
4.12(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, term, and renewal
5.10 to 5.12	Reserved
5.13(155A)	Registration verification
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 and 5.19	Reserved
5.20(155A)	Training and utilization of pharmacy support persons
5.21(155A)	Delegation of functions and 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) Reporting discipline and criminal convictions
- 5.27(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 Reserved
- 7.5(124,155A) Security
- 7.6(155A) Pharmacist absence
- 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) Definitions
- 8.3(155A) Responsible parties
- 8.4(155A) Pharmacist identification and staff logs
- 8.5(155A) Environment and equipment requirements
- 8.6 Reserved
- 8.7(155A) Procurement, storage, and recall of drugs and devices
- 8.8 Reserved
- 8.9(124,155A) Records storage
- 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 registered pharmacy staff
- 8.15(155A) Delivery of prescription drugs and devices

8.16(124,155A)	Confidential information
8.17	Reserved
8.18(124,155A)	Electronic prescription mandate
8.19(124,126,135,155A,280)	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(155A)	Notification of interchangeable biological product selection
8.23	Reserved
8.24(155A)	Documented verification
8.25	Reserved
8.26(155A)	Continuous quality improvement program
8.27 to 8.34	Reserved
8.35(155A)	Pharmacy license

## 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(124)	Who may register
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 of application or discipline of registration
10.11(124,147,155A)	Registration verification
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(124)	Schedule III through V accountability
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 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(155A)	General requirements for telepharmacy support person
13.13 to 13.15	Reserved
13.16(124,155A)	Telepharmacy site—initial application
13.17	Reserved
13.18(155A)	Opening of traditional pharmacy
13.19 to 13.22	Reserved
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 DISTRIBUTOR LICENSES

- 17.1(155A) Purpose and scope
- 17.2(155A) Definitions
- 17.3(155A) Wholesale distributor license
- 17.4(155A) Grounds for denial
- 17.5 and 17.6 Reserved
- 17.7(124,155A) Compliance with federal and state laws
- 17.8(124,155A) Written policies and procedures
- 17.9(155A) Facilities
- 17.10(124,155A) Security
- 17.11(155A) Storage and handling
- 17.12 to 17.16 Reserved
- 17.17(155A) Reporting discipline and criminal convictions
- 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
20.24(155A)	Annual reporting of interstate distribution of compounded preparations

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(124,155A)	Electronic prescription mandate and exemptions
21.9(124,155A)	Exemption from electronic prescription mandate—petition
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

## CARE FACILITY PHARMACY PRACTICE

23.1(155A)	Purpose and scope
23.2(155A)	Definitions
23.3(124,155A)	Freedom of choice
23.4(124,155A)	Responsibilities
23.5(124,155A)	Emergency drugs
23.6	Reserved
23.7(124,155A)	Policies and procedures
23.8	Reserved
23.9(124,155A)	Medication orders
23.10	Reserved
23.11(124,155A)	Drugs dispensed—general requirements
23.12	Reserved
23.13(124,155A)	Labeling drugs under special circumstances
23.14(124,155A)	Provision of drugs to a facility for immunization or screening programs
23.15(124,155A)	Return and reuse of drugs and devices
23.16	Reserved
23.17(124,155A)	Accountability of controlled substances
23.18	Reserved
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  
CRIMINAL CONVICTIONS

- 31.1(272C) Purpose and scope
- 31.2(272C) Definitions
- 31.3(272C) License application
- 31.4(272C) Eligibility determination
- 31.5(272C) Appeal
- 31.6(272C) Future petitions or applications

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(272C) Definitions
- 33.2(272C) Military education, training, and service credit
- 33.3(272C) Veteran and spouse licensure or registration
- 33.4(272C) Request for contested case

CHAPTER 34  
WAIVERS

- 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
- 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) Submission of waiver information
- 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
- 36.11(88GA,SF304) Prohibited grounds for discipline

#### CHAPTER 37 IOWA PRESCRIPTION MONITORING PROGRAM

- 37.1(124) Purpose and scope
- 37.2(124) Definitions
- 37.3(124) Registration
- 37.4(124) Prescription monitoring program advisory council
- 37.5 Reserved
- 37.6(124) Security of PMP credentials
- 37.7(124) PMP reporting—exemptions

37.8(124)	PMP reporting—dispensing prescribers
37.9(124)	PMP reporting—pharmacies
37.10 and 37.11	Reserved
37.12(124)	Reporting requirements
37.13(124)	Opioid antagonist administration by first responders
37.14 and 37.15	Reserved
37.16(124)	Access to PMP information
37.17(124)	Integrated systems
37.18(124)	PMP administrator access
37.19(124)	Prescriber activity reports
37.20(124)	Proactive notifications
37.21(124)	Record retention
37.22(124)	Information errors
37.23(124)	Discipline

## CHAPTER 38

Reserved

## CHAPTER 39

## EXPANDED PRACTICE STANDARDS

39.1(155A)	Purpose and scope
39.2 and 39.3	Reserved
39.4(155A)	Pharmaceutical care
39.5	Reserved
39.6(155A)	Statewide protocols
39.7(135,147A)	Opioid antagonist dispensing by pharmacist—standing order
39.8(155A)	Medications administered via prescription
39.9 to 39.12	Reserved
39.13(155A)	Collaborative pharmacy practice
39.14 and 39.15	Reserved
39.16(155A)	Pharmacy pilot or demonstration research projects

## CHAPTER 40

## TECHNOLOGY-ASSISTED TECHNICIAN PRODUCT VERIFICATION PROGRAMS

40.1(155A)	Purpose and scope
40.2(155A)	Definitions
40.3(155A)	TPV program requirements
40.4(155A)	Checking technician requirements
40.5 and 40.6	Reserved
40.7(155A)	Policies and procedures
40.8(155A)	TPV program quality assurance
40.9 and 40.10	Reserved
40.11(155A)	TPV program records

## 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

CHAPTER 42  
LIMITED DISTRIBUTOR LICENSES

42.1(155A)	Purpose and scope
42.2(155A)	Definitions
42.3(155A)	Limited distributor license
42.4 and 42.5	Reserved
42.6(155A)	Grounds for denial
42.7(155A)	Policies and procedures
42.8 and 42.9	Reserved
42.10(155A)	Requirements
42.11	Reserved
42.12(155A)	Records
42.13	Reserved
42.14(155A)	Reporting discipline and criminal convictions
42.15(155A)	Discipline

CHAPTER 43  
THIRD-PARTY LOGISTICS PROVIDER LICENSES

43.1(155A)	Purpose and scope
43.2(155A)	Definitions
43.3(155A)	3PL license
43.4	Reserved
43.5(155A)	Compliance with federal and state laws
43.6(155A)	Policies and procedures
43.7 and 43.8	Reserved
43.9(155A)	Reporting discipline and criminal conviction
43.10(155A)	Discipline

CHAPTERS 44 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