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

	PURPOSE AND ORGANIZATION	
1.1(17A)	Board mission	
1.2(17A,147,155A,	272C) Description and organization of board	
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	
	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	
2.1/155.4	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) 3.19 3.20(155A) 3.21(155A) 3.22 3.23(155A) 3.24 to 3.27 3.28(147,155A) 3.29(155A) 3.30(155A) 3.31(155A)	Identification of pharmacy technician Reserved Responsibility of supervising pharmacist Delegation of functions Reserved Functions a pharmacy technician shall not perform Reserved Unethical conduct or practice Denial of registration Reporting discipline and criminal convictions 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(155 A)	
6.1(155A)	Purpose and scope Pharmacist in charge
6.2(155A) 6.3(155A)	Reference library
6.4(155A)	Exemption from duplicate requirements
6.5 and 6.6	Reserved
6.7(124,155A)	
6.8(124,155A)	Security 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
	Patient counseling and instruction
6.14(155A) 6.15(124,126)	Return of drugs and devices
	Records
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)	
7.12(124,126,155A	•
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) 8.17 8.18(124,155A) 8.19(124,126,155A) 8.20(155A) 8.21(155A)	Confidential information Reserved Electronic prescription mandate) Manner of issuance of a prescription drug or medication order Valid prescriber/patient relationship Prospective drug use review
8.22(155A) 8.23(124,155A)	Notification of interchangeable biological product selection Individuals qualified to administer
8.24(155A) 8.25	Documented verification Reserved
8.26(155A)	Continuous quality improvement program
8.27 to 8.34	Reserved
8.35(155A)	Pharmacy license
	CHAPTER 9
	Reserved
	CHAPTER 10
10.1(124)	CONTROLLED SUBSTANCES
10.1(124) 10.2(124)	Purpose and scope 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 of application or discipline of registration
10.11(124,147,155)	
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(124,126,155	
10.25(124) 10.26(124)	Dispensing records 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
() -)	1

10.34(124)	Dispensing products containing ephedrine, pseudoephedrine, or phenylpropanolamine without a prescription	
10.35	Reserved	
10.36(124,155A)	Records	
10.30(124,133A)	Reserved	
10.38(124)	Revision of controlled substances schedules	
* *	Temporary designation of controlled substances	
10.39(124)	Excluded and exempt substances	
10.40(124)	Anabolic steroid defined	
10.41(124) 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,155		
11.2(124,147A,155		
11.3(124,147A,155	, ,	
11.4(124,147A,155		
11.5(124,147A,155	,	
11.6 and 11.7	Reserved	
11.8(124,147A,155	A) Identification	
11.9	Reserved	
	5A) Ownership of prescription drugs	
11.11(124,147A,15	5A) Policies and procedures	
11.12	Reserved	
11.13(124,147A,15		
11.14(124,147A,15	5A) Protocols	
11.15(124,147A,15	5A) Administration of drugs beyond the limits of a written protocol	
11.16(124,147A,15	5A) Administration of Schedule II controlled substances—pharmacy-based service program	
11.17 and 11.18	Reserved	
11.19(124,147A,15	5A) Patient care reports	
11.20(124,147A,15	5A) Prescription drugs in service programs	
11.21	Reserved	
11.22(124,147A,15	5A) Return of drugs	
11.23(124,147A,15	5A) Out-of-date drugs or devices	
11.24(124,147A,15	5A) Product recall	
11.25	Reserved	
11.26(124,147A,15	5A) Controlled substances records	
11.27(124,147A,15	5A) Ordering Schedule II controlled substances—medical director-based service programs	
11.28	Reserved	
11.29(124,147A,15	5A) Schedule II controlled substances perpetual inventory	
	5A) Controlled substances annual inventory	
11.31	Reserved	
	5A) Disposition of controlled substances	
	5A) Report of loss or theft of controlled substance	
11.34(124,147A,15	, .	

	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)	
14.7(22,124,155A)	
14.8(22,124,155A)	
14.9(22,124,155A)	* *
14.10(22,124,155A	, and the state of
14.11(22,124,155A	
14.12(22,124,155A	
14.13(22,124,155A	,
14.14(22,124,155A	
14.15(22,124,155A	·
14.16(22,124,155A	
` ' '	

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(124,126,155)	A) Policies and procedures	
	CHADTED 16	

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

	WITO EESTIEE DISTING CIGIC ETCE
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 PIPTION FILLING AND PROCESSING

	CENTRALIZED PRESCRIPTION FILLING AND
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) Del	ayed compliance
20.6(126,155A) Con	mpounding standards for outsourcing facilities
20.7 and 20.8 Res	served
20.9(124,155A) Pre	scriber/patient/pharmacist relationship
20.10(126,155A) Ant	ticipatory compounding
20.11(126,155A) Pro	hibition on resale of compounded preparations
20.12(126,155A) Con	mpounding copies of an approved drug
20.13(124,126,155A)	Use of flavoring agents
20.14 Res	served
20.15(124,126,155A)	Compounding for office use
20.16(124,126,155A)	Compounding for hospital use
20.17 and 20.18 Res	served
20.19(124,126,155A)	Labeling
20.20(126,155A) Lab	peling for batch preparation compounding
20.21 and 20.22 Res	served
20.23(124,126,155A)	Records

CHAPTER 21

ELECTRONIC DATA AND AUTOMATED SYSTEMS IN PHARMACY PRACTICE

EEEE TROTTE DIMITING HETOTRINED STOTEMS IN TIMMEMIET TRATETIEE	
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)

(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) 28.16(17A)	Review by board of rules	
28.10(1/A)		
	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	
29.2(68B) 29.3(68B)	Authorized sales	
29.4(68B)	Application for consent Limitation of consent	
29.5(68B)		
IOW	CHAPTER 30 A MONITORING PROGRAM FOR PHARMACY PROFESSIONALS	
30.1(272C)	Iowa monitoring program for pharmacy professionals committee Definitions	
30.2(272C)		
30.3(272C)	Organization of the committee	
30.4(272C)	Eligibility Towns of participation	
30.5(272C)	Terms of participation	
30.6(272C)	Confidentiality Authority for 28E agreements	
30.7(28E)	Authority for 28E agreements	

CHAPTER 31 CRIMINAL CONVICTIONS

	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
22 1(272D)	NONPAYMENT OF STATE DEBT
32.1(272D)	Definitions Issuance or renewal of a license—denial
32.2(272D)	Suspension or revocation of a license
32.3(272D)	-
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 licensure or registration
33.4(272C)	Request for contested case
	CHAPTER 34
	WAIVERS
34.1(17A)	Definition
	47,155A,205,272C) Scope of chapter
	47,155A,205,272C) Applicability of chapter
34.4(17A)	Criteria for waiver
` /	47,155A,205,272C) Filing of petition
34.6(17A)	Content of petition
34.7(17A)	Additional information and providing notice
	Reserved
34.10(17A)	Ruling
34.11(17A,22)	
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

	17,155A,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,12	26,147,155A,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
	26,147,155A,205,272C) Applications for rehearing
	Stays of board actions
	No factual dispute contested cases
	26,147,155A,205,272C) Emergency adjudicative proceedings
	C) Application for reinstatement
35.37(17A,22,272C	,
35.38(17A)	Judicial review
33.30(1711)	
	CHAPTER 36
	DISCIPLINE
36.1(147,155A,272	C) Authority
36.2(147,155A,272	C) Definitions
36.3(147,155A,272	C) Complaints, investigations, and board action
36.4(17A,147,152,2	272C) Issuance of investigatory subpoenas
36.5(147,272C)	Peer review committee
36.6(147,155A,272	
36.7(147,155A,272	•
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	
30.11(00071,51307	Tromoted 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 and 37.5	Reserved
37.6(124)	Security of PMP credentials
37.7(124)	PMP reporting—exemptions
37.8(124)	PMP reporting—dispensing prescribers
2,10(121)	The stand ambanemed branching

37.9(124) 37.10 and 37.11 37.12(124) 37.13(124) 37.14 and 37.15 37.16(124) 37.17(124) 37.18(124)	PMP reporting—pharmacies Reserved Reporting requirements Opioid antagonist administration by first responders Reserved Access to PMP information Integrated systems PMP administrator access
37.19(124) 37.20(124)	Prescriber activity reports 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) 39.7(135,147A)	Statewide protocols Opioid antagonist dispensing by pharmacist—standing order
39.8(155A)	Statewide protocol—naloxone
39.9(155A)	Statewide protocol—nicotine replacement tobacco cessation products
39.10(155A)	Vaccine administration by pharmacists—physician-approved protocol
39.11(155A)	Vaccine administration by pharmacists—statewide protocol
39.12	Reserved
39.13(155A)	Collaborative drug therapy management
39.14 and 39.15	Reserved
39.16(155A)	Pharmacy pilot or demonstration research projects
	CHAPTER 40
	OGY-ASSISTED TECHNICIAN PRODUCT VERIFICATION PROGRAMS
	Purpose and scope
40.2(155A)	Definitions TRY are grown as expressed to
40.3(155A) 40.4(155A)	TPV program requirements 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
	CILLA DETERMAN
	CHAPTER 42
40.1(155.4)	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
	CILLA DETERRIO AAA AA AA
	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