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)	
1.5(17A,21)	Meetings
1.6(124,147,155A)	•
	155A) Overpayment of fees
, , , ,	, 1 3
	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
	CHA PETER A
	CHAPTER 3
2.1(1.55.1)	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) 3.22(155A) 3.23(155A) 3.24(155A) 3.25 to 3.27 3.28(147,155A)	Delegation of functions Technical functions Tasks a pharmacy technician shall not perform New prescription drug orders or medication orders Reserved Unethical conduct or practice
3.29(155A) 3.30(155A)	Denial of registration 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 5.12(155A)	Reserved
5.13(155A) 5.14(155A)	Registration certificates Notifications to the board
5.14(155A) 5.15(155A)	Identification of pharmacy support person
5.15(133A) 5.16	Reserved
5.17(155A)	Tasks a pharmacy support person shall not perform
5.18(155A)	Nontechnical pharmacy support tasks
5.19	Reserved
5.20(155A)	Training and utilization of pharmacy support persons
5.21(155A)	Responsibility of supervising pharmacist
5.22(155A)	Delegation of nontechnical functions
5.23	Reserved
5.24(155A)	Denial of registration
5.25(147,155A)	Unethical conduct or practice
5.26(155A)	Discipline of pharmacy support persons

CHAPTER 6 GENERAL PHARMACY PRACTICE

6.1(155A)	Purpose and scope
6.2(155A)	Pharmacist in charge
6.3(155A)	Reference library
6.4(155A)	Exemption from duplicate requirements
6.5 and 6.6	Reserved
6.7(124,155A)	Security
6.8(124,155A)	Prescription processing documentation
6.9(124,155A)	Transfer of prescription
6.10(126,155A)	Prescription label requirements
6.11 and 6.12	Reserved
6.13(155A)	Patient record system
6.14(155A)	Patient counseling and instruction
6.15(124,126)	Return of drugs and other items
6.16(124,155A)	Records

CHAPTER 7

HOSPITAL PHARMACY PRACTICE

7.1(155A)	Purpose and scope
7.2(155A)	Pharmacist in charge
7.3(155A)	Reference library
7.4 and 7.5	Reserved
7.6(124,155A)	Security
7.7(155A)	Verification by 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	Reserved
7.12(124,126,155A) Drugs dispensed to patients as a result of an emergency room visit	
7.13(124,155A)	Records

CHAPTER 8

UNIVERSAL PRACTICE STANDARDS

	CIVITY ENGINE THE ICTION OF TH
8.1(155A)	Purpose and scope
8.2(155A)	Pharmaceutical care
8.3(155A)	Responsibility
8.4(155A)	Pharmacist identification
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
8.35(155A)	Pharmacy license
	CHAPTER 9
A	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 Records
9.13(147,155A) 9.14	Reserved
9.14 9.15(147,155A)	Decentralized unit dose AMDS
9.15(147,155A) 9.16(147,155A)	Centralized unit dose AMDS
9.17(147,155A)	Outpatient AMDS
9.18(124,155A)	Remote dispensing site operations
9.19	Reserved
9.20(124,155A)	Drugs at a remote dispensing site
9.21(124,155A)	Record keeping
	CHAPTER 10
	CHAPTER 10 CONTROLLED SUBSTANCES
10.1(124)	Who shall register
10.1(124)	Application forms
10.2(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(124,147,155	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(124,126,155	A) 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
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
10.10(121,120)	Thubble steroid defined
	CHAPTER 11
	DRUGS IN EMERGENCY MEDICAL SERVICE PROGRAMS
11.1(124,147A,155	
11.2(124,147A,155.	
11.3(124,147A,155.	A) General requirements
11.4(124,147A,155.	,
11.5(124,147A,155.	A) Records
11.6(124,147A,155.	A) Inspections
11.7(124,147A,155.	A) Security and control
	CHAPTER 12
	PRECURSOR SUBSTANCES
12.1(124B)	Precursor substance identified
12.1(124B) 12.2(124B)	Reports required
12.2(124B) 12.3(124B)	Form of reports
,	1
12.4(124B) 12.5(124B)	Monthly reporting option Exemptions
12.5(124B) 12.6(124B)	Identification of purchaser or other recipient
12.7(124B)	Permits
12.8(124B)	Denial, modification, suspension, or revocation of permit
12.0(124D)	Demar, modification, suspension, or revocation of permit
	CHAPTER 13
	STERILE COMPOUNDING PRACTICES
13.1(124,126,155A)	
13.2(124,126,155A)	
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
	CHAPTER 14
	PUBLIC INFORMATION AND INSPECTION OF RECORDS
14.1(22,124,155A)	
14.2(22,124,155A)	
14.3(22,124,155A)	
14.4(22,124,155A)	•
14.5(22,124,155A)	
14.3(22,124,133A)	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)	
14.9(22,124,155A)	••
14.10(22,124,155)	<u> </u>
14.11(22,124,155	
14.12(22,124,155)	
14.13(22,124,155)	
14.14(22,124,155)	
14.15(22,124,155)	
14.16(22,124,155	
	•
	CHAPTER 15
15 1/155 1	CORRECTIONAL PHARMACY PRACTICE
15.1(155A)	Purpose and scope
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(124,126,155)	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	
15 1 (155 1)	WHOLESALE DRUG LICENSES	
17.1(155A)	Definitions	
17.2	Reserved	
17.3(155A)	Wholesale drug license	
17.4(155A)	Minimum qualifications Personnel	
17.5(155A)		
17.6(155A) 17.7(124,155A)	Responsibility for conduct Distribution to authorized licensees	
17.7(124,133A) 17.8(124,155A)	Written policies and procedures	
17.9(155A)	Facilities	
17.10(124,155A)	Security	
17.11(155A)	Storage	
17.11	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	
C	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(124,1	*	Storage and shipment of drugs and devices
19.9(155A	.)	Patient record system, prospective drug use review, and patient counseling
19.10(155	*	Discipline
		CILLA DEED AO
		CHAPTER 20
20 1/124 1	26 155 4	PHARMACY COMPOUNDING PRACTICES
20.1(124,1 20.2(124,1		
20.2(124,1		,
20.3(124,1		A) General requirements Organization and personnel
20.4(126,1	/	Drug compounding facilities
20.5(126,1		Sterile products and radiopharmaceuticals
20.0(120,1	33A)	Reserved
20.8(126,1	554)	Equipment
20.8(126,1		Control of bulk drug substances, components, containers, and closures
20.10(124,		
20.11(124)		Bulk compounding
		(A) Records
20.12(121,	,120,133	71) 1000103
		CHAPTER 21
		ELECTRONIC DATA IN PHARMACY PRACTICE
21.1(124,1	/	Definitions
21.2(124,1		System security and safeguards
21.3(124,1		Verifying authenticity of an electronically transmitted prescription
21.4(124,1		Automated data processing system
21.5(124,1	55A)	Pharmacist verification of controlled substance refills—daily printout or logbook
21.6		Reserved
21.7(124,1		Electronically prepared prescriptions
21.8(124,1		Computer-to-computer transmission of a prescription
21.9(124,1		Facsimile transmission (fax) of a prescription
21.10 and		Reserved
21.12(124,	,	Prescription drug orders for Schedule II controlled substances
21.13(124,	,133A)	Prescription drug orders for Schedule II controlled substances—emergency situations
21 14(124	1551)	Facsimile transmission of a prescription for Schedule II narcotic
21.14(124,	,133A)	substances—parenteral
21.15(124,	1554)	Facsimile transmission of Schedule II controlled substances—long-term care
21.13(124,	,133A)	facility patients
21.16(124,	155A)	Facsimile transmission of Schedule II controlled substances—hospice patients
21.10(121,	,13311)	ruestimic transmission of senedate if controlled substances mospiec 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(126,1	55A)	Patient med paks
22.6		Reserved
22.7(124,1	55A)	Emergency/first dose drug supply
22.8	,	Reserved
22.9(155A	.)	Home health agency/hospice emergency drugs

CHAPTER 23 LONG-TERM CARE PHARMACY PRACTICE

	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
23.17(124,155A) 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
23.21(124,133A)	Destruction of 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
	GV L DEPTH A
	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) 27.9(17A)	Refusal to issue order
41.7(1/A)	retusal to issue ofuci

27.10(17A)	Contents of declaratory order—effective date
27.11(17A)	Copies of orders
27.12(17A)	Effect of a declaratory order
	CHAPTER 28
	AGENCY PROCEDURE FOR RULE MAKING
	(Uniform Rules)
28.1(17A)	Applicability
28.2(17A)	Advice on possible rules before notice of proposed rule adoption
28.3(17A)	Public rule-making docket
28.4(17A)	Notice of proposed rule making
28.5(17A)	Public participation
28.6(17A)	Regulatory analysis
28.7(17A,25B)	Fiscal impact statement
28.8(17A)	Time and manner of rule adoption
28.9(17A)	Variance between adopted rule and published notice of proposed rule adoption
28.10(17Å)	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
30.0(13311)	110gram rando
	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

Reserved

CHAPTER 34

RIII ES FOR WAIVERS AND VARIANCES

	RULES FOR WAIVERS AND VARIANCE
34.1(17A)	Definition
34.2(17A,124,126,1	47,155A,205,272C) Scope of chapter
34.3(17A,124,126,1	47,155A,205,272C) Applicability of chapter
34.4(17A)	Criteria for waiver or variance
34.5(17A,124,126,1	47,155A,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

CHAPTER 35

CONTESTED CASES

35.1(17A,124,124E	3,126,147,155A,205,272C)	Scope and applicability
35.2(17A,272C)	Definitions	
25 2(17 A)	Tima raquiramanta	

35.3(17A) Time requirements 35.4 Reserved

35.4 Reserved

34.16(17A)

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

Judicial review

Disqualification 35.9(17A)

35.10(17A,272C) Consolidation—severance

35.11(17A,272C) Service and filing of pleadings and other papers

35.12(17A,272C) Discovery 35.13(17A,272C) Subpoenas 35.14(17A,272C) Motions

35.15(17A,272C) Prehearing conference

35.16(17A,272C) Continuances 35.17(17A) Withdrawals 35.18 Reserved

35.19(17A,124B,126,147,155A,205,272C) Hearing procedures in contested cases

35.20(17A,272C) Evidence 35.21(17A,272C) Default

35.22(17A,272C) Ex parte communication

35.23(17A,272C)	Recording costs		
35.24(17A,272C)	Interlocutory appeals		
35.25(17A)	Final decision		
35.26(17A,124B,12	26,147,155A,205,272C) Appeals a	nd review	
	26,147,155A,205,272C) Application	ons for rehearing	
35.28(17A,272C)	Stays of board actions		
35.29(17A,272C) No factual dispute contested cases			
35.30(17A,124B,12	26,147,155A,205,272C) Emergence	y adjudicative proceedings	
	СНАРТЕ	R 36	
	DISCIPL		
	(C) Authority and grounds for di	scipline	
36.2(155A,272C)			
	Peer review committees		
	3,126,147,155A,272C) Disciplinar		
	3,126,147,155A,272C) Notice of 6		
	7,155A,272C) Informal settlement		
36.7(272C)	Appearance		
	7,155A,272C) Order of proceeding	gs	
36.9(272C)	Confidentiality		
	Notification of decision		
	Board decision		
	Publication of decisions		
	17,155A,272C) Reinstatement		
	17,155A,272C) Informal reinstater		
		er of a license, permit, or registration	
	17,155A,272C) License, permit, or		
	Order for mental or physical exar		
36.18(272C)	Disciplinary hearings—fees and o	costs	
	СНАРТЕ		
	IOWA PRESCRIPTION MO	NITORING PROGRAM	
37.1(124)	Purpose		
37.2(124)	Definitions		
37.3(124)	Requirements for the PMP		
37.4(124)			
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		