

Louisiana Administrative Code

Title 46 – Professional and Occupational Standards

Part LIII: Pharmacists

Chapter 12. Automated Medication Systems

§1201. Definitions

Automated Medication System—includes, but is not limited to, a mechanical system that performs operations or activities, other than compounding or administration, relative to the storage, packaging, or delivery of medications, and which collects, controls, and maintains all transaction information. An automated medication system may be profile driven, non profile driven, or a combination of both.

Final Checks of Work—the requirement that only a pharmacist supervises and releases the completed product prepared by a pharmacy technician.

Floor Stock/First Dose Cabinet—a medication storage device, which shall be used by personnel, authorized by a protocol established by the pharmacist in charge, to gain access to doses as needed and first doses in patient care areas. In addition, a floor stock/first dose cabinet may be used to store medications in such specialty areas including, but not limited to, emergency room, surgery suite, and endoscopy suites.

Healthcare Setting—a place where healthcare services are rendered on a routine basis by credentialed healthcare professionals.

Non Profile Driven—system does not require prior or concomitant pharmacist review of medication order/prescriptions in order to gain access to the system for medication administration. A non-profile driven system may include, but is not limited to, a night drug cabinet, emergency drug kits, or floor stock/first dose cabinet.

Off Site Facility—the location of a building that houses a licensee of the Department of Health and Hospitals, but which does not house a board permitted pharmacy.

On Site Facility—the location of a building that houses a board permitted pharmacy.

Profile Driven—system requires that medication orders/prescriptions be reviewed by the pharmacist for appropriateness, dosage, and contraindications prior to, or concomitantly with, being entered into the system, and before access is allowed into the system for medication administration.

Remote Dispensing System—a profile driven automated medication dispensing system employing bidirectional audio-visual technology to facilitate pharmacist communication with a patient or caregiver.

AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR

§1203. Automated Medication System(s) Registration

A. The entire system shall be registered with the board and facilities shall meet the following conditions:

1. Facility shall possess a:

a. license from the Health Standards Section of the Department of Health and Hospitals, or

b. Controlled Dangerous Substance License from the Health Standards Section of the Department of Health and Hospitals, or

c. permit from the board.

2. Registration fee for a facility not permitted by the board is as identified in [R.S. 37:1184.C.xii](#).

3. No registration fee will be assessed a board permitted pharmacy.

4. Registration expires annually on June 30.

5. Initial application shall be completed and signed by the registrant of the facility and the pharmacist in charge of the system(s). The completed, signed application and required fee shall be submitted to the board office no later than 30 days prior to installation of the system.

- 53 6. ~~Annual Renewal.~~ The board shall make available an application for renewal to each registrant
54 ~~on or before May 1 each year. Said application shall be completed, signed, and, with annual~~
55 ~~fee, returned to the board office to be received on or before June 1 each year.~~
56 7. ~~Expired Registration.~~ A registration that is not renewed shall be null and void. A renewal
57 ~~application for an expired registration shall be requested by the registrant and the completed,~~
58 ~~signed application may be referred to the board's reinstatement committee for disposition in~~
59 ~~accordance with [R.S. 37:1230](#).~~
60 8. ~~Reinstatement.~~ The holder of a registration that has expired may be reinstated only upon
61 ~~written application to the board and upon payment of all lapsed fees and a penalty to be fixed~~
62 ~~by the board. Other conditions of reinstatement may be required at the discretion of the~~
63 ~~board.~~

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65 A. Requirement for Registration

- 66 1. A pharmacy intending to supply medications for use within an automated medication system, as
67 defined at R.S. 37:1164, shall obtain an AMS registration prior to engaging in such activity.
68 2. The placement of medications within an automated medication system in the absence of an AMS
69 registration shall substantiate a violation of R.S. 37:1241(A)(12) and shall subject the pharmacy
70 to disciplinary action by the board.
71 3. A pharmacy intending to supply controlled substances for use within an automated medication
72 system shall obtain a controlled dangerous substance (CDS) license in addition to the AMS
73 registration. The pharmacy shall also obtain a federal registration from the U.S. Drug
74 Enforcement Administration (DEA) prior to placing controlled substances within the AMS.
75 4. The placement of controlled substances within an automated medication system in the absence of
76 an AMS registration, CDS license, and DEA registration shall substantiate a violation of R.S.
77 37:1241(A)(12) and R.S. 40:973 and shall subject the pharmacy to disciplinary action by the
78 board.
79 5. The operation of a remote dispensing system without an AMS registration shall substantiate a
80 violation of R.S. 37:1241(A)(12) and shall subject the pharmacy to disciplinary action by the
81 board.

82 B. Eligibility for Registration

- 83 1. A pharmacy intending to supply medications for use within an automated medication system may
84 do so when the AMS is placed at any of the following locations:
85 a. within a facility in possession of a controlled dangerous substance license issued by the board.
86 b. within a hospital or other institutional facility in possession of an operating license issued by
87 the state department of health.
88 c. within a detention or correctional facility operated by or under contract with the state
89 department of public safety and corrections or other local governmental entity.
90 2. A pharmacy may operate a remote dispensing system when the system is placed within a
91 healthcare setting where the pharmacist-in-charge can ensure the security and environmental
92 integrity of the medications and devices placed within the system as well as the security and
93 confidentiality of the protected health information used therein.

94 C. Application for Initial Issuance of Registration

- 95 1. The board shall develop an application form suitable for the AMS registration. The board may
96 revise that application form on its own initiative in order to collect the information it deems
97 necessary to properly evaluate an applicant.
98 2. The application shall be accompanied by payment of the registration fee authorized by R.S.
99 37:1184.
100 3. The board shall not process applications received by facsimile, or that are incomplete, or
101 submitted with the incorrect fee.
102 4. The submission of a false or fraudulent application shall substantiate a violation of R.S.
103 37:1241(A)(2) and shall subject the applicant to disciplinary action by the board.
104 5. When determined appropriate by the board, the applicant may be required to meet with a
105 committee or agent of the board prior to the issuance of the registration.
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- 107 D. Maintenance of Registration
- 108 1. A registration shall be valid only for the pharmacy to which it was issued and the physical location
- 109 of the AMS identified on the application. The registration shall not be subject to sale, assignment
- 110 or other transfer, voluntary or involuntary, nor shall the registration be valid for any premises
- 111 other than the physical location for which it was issued.
- 112 2. A duplicate or replacement registration shall be issued upon the written request of the owner of the
- 113 registration and payment of the fee authorized by R.S. 37:1184. A duplicate or replacement
- 114 registration shall be marked as such, and it shall not serve or be used as an additional or second
- 115 registration.
- 116 3. In the event a pharmacy intends to relocate an automated medication system to a different address,
- 117 the pharmacy shall notify the board of its intent to do so, providing both current and new
- 118 addresses. A change in business address may require an inspection by the board or its designee.
- 119 E. Application for Renewal of Registration
- 120 1. The pharmacy shall complete an application for the renewal of the registration and submit it to the
- 121 board prior to the expiration date of the registration. The application shall be accompanied by the
- 122 fee authorized by R.S. 37:1184.
- 123 2. The board shall not process applications received by facsimile, or that are incomplete, or
- 124 submitted with the incorrect fee.
- 125 3. An AMS registration not renewed by the expiration date shall be classified as expired. The
- 126 operation of an automated medication system with an expired registration shall substantiate a
- 127 violation of R.S. 37:1241(A)(12) and shall subject the pharmacy to disciplinary action by the
- 128 board.
- 129 F. Relinquishment of Registration
- 130 1. In the event a pharmacy intends to cease supplying medications or devices to an automated
- 131 medication system, it shall relinquish the registration to the board no later than 10 days following
- 132 the effective date of such decision.
- 133 2. A pharmacy may not transfer a registration to another pharmacy.
- 134 G. Application for Reinstatement of Suspended or Revoked Registration
- 135 1. An application for the reinstatement of an AMS registration previously suspended or revoked by
- 136 the board may only be approved in compliance with R.S. 37:1249.
- 137 2. The applicant shall complete an application form for this specific purpose supplied by the board
- 138 and shall attach any documentation and fees identified in R.S. 37:1184.
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140 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

141 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271

142 (June 2000) effective July 1, 2000, amended LR 38:1235 (May 2012), amended by the Department of Health, Board

143 of Pharmacy, LR

144

145 §1205. Pharmacist-in-Charge Responsibilities

- 146 A. The pharmacist-in-charge shall be a Louisiana licensed pharmacist ~~and has~~ with the following
- 147 responsibilities:
- 148 1. assuring that the system is in good working order and accurately provides the correct strength,
- 149 dosage form, and quantity of the drug prescribed while maintaining appropriate record-
- 150 keeping and security safeguards.
- 151 2. establishment of a quality assurance program prior to implementation of a system and the
- 152 supervision of an ongoing quality assurance program that monitors appropriate use and
- 153 performance of a system, which is evidenced by ~~written~~ policies and procedures developed by
- 154 the pharmacist-in-charge.
- 155 3. ~~provide 30 days written notice to the board of removal of the system.~~
- 156 4. define access to the system in policy and procedures of the pharmacy, in compliance with
- 157 state and federal regulations.
- 158 5. assign, discontinue, or change access to the system.
- 159 6. ensure that access to the medications complies with state and federal regulations as
- 160 applicable.

- 161 7. ensure that the system is stocked/restocked accurately and in accordance with established
 162 written pharmacy policies and procedures.
 163 8. maintain or have access to all records of documentation specified in this Section for two years
 164 or as otherwise required by law.
 165 9. ~~notify each licensed prescriber that his medication orders/prescriptions are not restricted to the~~
 166 ~~limited number of medications which are stocked within a facility's automated medication~~
 167 ~~system by placing a prominent notice to that effect on the cover of or near the beginning of~~
 168 ~~such patient's medical chart or medical record.~~
 169 10. continuous monitoring and documentation of temperature in the drug storage areas including
 170 a mechanism to alert the pharmacist when defined parameters are out of range as well as an
 171 action plan to address such excursions. A pharmacy's failure to document the integrity of the
 172 drug supply or remediate for excursions as appropriate shall substantiate a violation of R.S.
 173 37:1241(A)(18) and shall subject the pharmacy to disciplinary action by the board.
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175 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

176 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 177 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR
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179 **§1207. Pharmacist Review**

- 180 A. System shall be used in settings that ensure medication orders are reviewed by a pharmacist prior to
 181 administration and in accordance with established policies and procedures and good pharmacy
 182 practice. A policy and procedure ~~protocol~~ shall be adopted for non-profile driven systems to
 183 retrospectively review medications orders which cannot be reviewed prior to medication
 184 administration, as provided in LAC 46:111.1209.2.
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186 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

187 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 188 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR
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190 **§1209. Policies and Procedures**

- 191 A. ~~The development of an automated medication system policy and procedures is the responsibility of the~~
 192 ~~pharmacist in charge, who shall submit the complete automated medication system policy and~~
 193 ~~procedures to the board for approval, on request. These policies and procedures shall be reviewed by~~
 194 ~~the pharmacist in charge, at least annually and modified if needed, and such review documented.~~
 195 ~~They shall include, but are not limited to, the following:~~
 196 1. ~~criteria for selection of medications to be stored in each system, provided that in facilities~~
 197 ~~licensed by the Department of Health and Hospitals, but not by the board, the selection~~
 198 ~~criteria shall not include the substitution by the pharmacist of a product that is not an~~
 199 ~~equivalent drug product to the product originally prescribed by the physician or practitioner~~
 200 ~~without the explicit consent of the physician or practitioner;~~
 201 2. ~~criteria for medications qualifying for use with a non-profile driven system and the locations~~
 202 ~~and situations that this type of system can be used in; and~~
 203 3. ~~information on the system as outlined below:~~
 204 a. ~~access-~~
 205 i. ~~system entry-~~
 206 ii. ~~access codes-~~
 207 iii. ~~system access privileges-~~
 208 iv. ~~changing access privileges-~~
 209 v. ~~termination of user-~~
 210 vi. ~~temporary access codes-~~
 211 vii. ~~password assignment-~~
 212 b. ~~controlled substances-~~
 213 i. ~~chain of custody-~~

- 214 ii. discrepancy resolution.
- 215 e. data:
- 216 i. archiving.
- 217 ii. stored/uploading to database.
- 218 iii. backup.
- 219 d. definitions.
- 220 e. downtime procedures (see malfunction).
- 221 f. emergency procedures.
- 222 g. information security/confidentiality:
- 223 i. patient information.
- 224 ii. medication information.
- 225 iii. transaction files.
- 226 iv. information update plan.
- 227 v. patient update plan.
- 228 vi. information access.
- 229 h. inspection.
- 230 i. installation requirements.
- 231 j. maintenance, e.g., service and repair protocols.
- 232 k. medication administration:
- 233 i. medication and patient validation.
- 234 ii. administration verification.
- 235 l. medication security:
- 236 i. acquisition and disposition records.
- 237 ii. proof of delivery.
- 238 iii. chain of custody of controlled substances (institutions).
- 239 iv. security management and control.
- 240 v. medication loading and storage.
- 241 vi. medication loading records.
- 242 vii. medication containers.
- 243 viii. cross contamination.
- 244 ix. lot number control.
- 245 x. inventory.
- 246 xi. utilization review.
- 247 xii. research.
- 248 m. malfunction:
- 249 i. troubleshooting.
- 250 ii. power failure.
- 251 n. quality assurance/quality improvement
- 252 i. documentation and verification of proper loading and refilling of
- 253 device.
- 254 ii. removal of drugs for administration, return, or waste.
- 255 iii. recording, resolving, and reporting of discrepancies.
- 256 iv. periodic audits to assure compliance with policies and procedures.
- 257 o. reports:
- 258 i. system maintenance.
- 259 ii. administrative functions.
- 260 iii. inventory.
- 261 iv. error.
- 262 v. discrepancies.
- 263 vi. activity.
- 264 vii. problem.
- 265 p. medication inventory management.
- 266 q. staff education and training.
- 267 r. system set up.

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HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR

§1211. Documentation

- A. Documentation as to type of equipment, serial number, content, policies and procedures and location shall be maintained ~~on-site~~ in the pharmacy for review by the board. Such documentation shall include, but is not limited to:
1. name, address, and permit number of the pharmacy ~~or licensed health care facility~~ and the location where the system is operational;
 2. manufacturer's name and model;
 3. quality assurance policies and procedures to determine continued appropriate use and performance of the system;
 4. policies and procedures for system operation, safety, security, accuracy, patient confidentiality, access, controlled substances, data retention, definitions, downtime procedures, emergency or first dose procedures, inspection, installation requirements, maintenance security, quality assurance, medication inventory, staff education and training, system set-up, and malfunction procedures; and
 5. security procedures sufficient to prevent unauthorized access or use, prevent the illegal use or disclosure of protected health information, and comply with any applicable federal or state regulations.
- B. A current copy of all pharmacy policies and procedures related to the use of the system shall be maintained at all ~~off-site facility~~ locations where the system is being used, ~~as well as the pharmacy of the pharmacist-in-charge.~~

AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR

§1213. Records

- A. Records and/or electronic data kept by the system shall meet the following requirements:
1. All events involving access to the contents of the system shall be recorded electronically.
 2. In the event controlled substances are stored in the system, the records shall include the positive identification (as defined in Section 1119 of ~~the Board's rules this Part~~) of the personnel retrieving and administering the controlled substances to the patient.
 3. These internal records shall be maintained for one year by the pharmacist-in-charge and shall be readily available to the board. Such records shall include:
 - a. identity of system accessed;
 - b. identification of the individual accessing the system;
 - c. type of transaction;
 - d. name, strength, dosage form, and quantity of the drug accessed;
 - e. name or identification number of the patient, ~~or identification numbers~~ for whom the drug was ordered;
 - f. identification of the certified pharmacy technician or pharmacist stocking or restocking the medications in the system; and
 - g. such additional information as the pharmacist-in-charge may deem necessary.

AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, amended LR 40:2256 (November 2014), effective January 1, 2015, amended by the Department of Health, Board of Pharmacy, LR

§1215. Security System(s)

- A. ~~System shall have adequate security system and procedures, evidenced by written pharmacy policies and procedures, to:~~
- ~~1. prevent unauthorized access or use;~~
 - ~~2. comply with any applicable federal and state regulations; and~~
 - ~~3. maintain patient confidentiality.~~

AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR

§1217. Stocking and Restocking

- A. ~~*On Site Facility System(s)*. The stocking and restocking of all medications in the on site system shall be accomplished by Louisiana licensed pharmacists, and/or Louisiana certified pharmacy technicians under the supervision of Louisiana licensed pharmacists. A pharmacist must conduct final checks of work performed by a pharmacy technician. The pharmacy shall have a mechanism in place to identify the certified pharmacy technician stocking or restocking and the pharmacist checking the accuracy of the medications to be stocked or restocked in the automated medication systems.~~
- B. ~~*Off Site Facility System(s)*. The stocking and restocking of all medications in the off site system shall be accomplished by Louisiana licensed pharmacists; however, the certified pharmacy technician may stock or restock an off site facility system provided a pharmacist is physically present at the off site facility and supervises and verifies the stocking and/or restocking prior to use. The pharmacy shall have a mechanism in place to identify the certified pharmacy technician stocking or restocking and the pharmacist checking the accuracy of the medications to be stocked or restocked in the system.~~
- A. The stocking and restocking of medications and devices within an automated medication system shall be performed by a pharmacist, or in the alternative, a pharmacy intern, pharmacy technician, or pharmacy technician candidate under the supervision of a pharmacist.
- B. When the pharmacy employs electronic product verification procedures as described within this Subsection, the stocking and restocking of medications and devices within an automated medication system may be performed by other personnel approved by the pharmacist-in-charge.
- C. Electronic Product Verification
1. A bar code verification, electronic verification, or similar verification process which prohibits any human intervention following pharmacist verification of the product may be utilized to assure the correct selection of drugs to be placed into an automated medication system.
 2. The use of a bar code, electronic, or similar verification process shall require an initial quality assurance validation followed by ongoing quality assurance reviews at intervals no greater than 90 days since the previous review, all conducted by a pharmacist.
 3. When a bar code verification, electronic verification, or similar verification process is utilized as specified in the Paragraph, and in the absence of any human intervention in the product selection process, the stocking and restocking functions in systems located either on site or off site may be performed by a pharmacy technician without the necessity of direct pharmacist supervision.

AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271 (June 2000) effective July 1, 2000, amended LR 41:1488 (August 2015), amended by the Department of Health, Board of Pharmacy, LR

§1219. Packaging and Labeling

- A. ~~All containers of medications stored in the system shall be packaged and labeled in accordance with federal and state laws and regulations and contain an established satisfactory beyond use date based on U.S.P. standards.~~

374 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

375 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
376 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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378 **§1221. Proof of Use**

379 A. ~~For medication removed from the system for patient administration, the system shall document, at a
380 minimum, the following:~~

- 381 1. ~~name of the patient or resident;~~
- 382 2. ~~patient's or resident's medical record number or identification number, or room and bed
383 number;~~
- 384 3. ~~date and time medication was removed from the system;~~
- 385 4. ~~name, initials, or other unique identifier of the person removing the drug; and~~
- 386 5. ~~name, strength, and dosage form of the medication or description of the medical device
387 removed.~~

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389 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

390 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
391 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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393 **§1223. Wasted, Discarded, or Unused Medications**

394 A. ~~The system shall provide a mechanism for securing and accounting for wasted, discarded, or unused
395 medications removed from the system according to policies and procedures, and existing state and
396 federal law.~~

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398 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

399 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
400 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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402 **§1225. Inspection**

403 A. ~~System records shall be available and readily retrievable for board inspection and review during
404 regular working hours of operation. The system itself is also subject to inspection at that time.~~

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406 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

407 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
408 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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410 **§1227. Out of State Pharmacies**

411 A. ~~Out of state pharmacies must have applied for and been issued an out of state pharmacy permit by the
412 board as identified in regulations. Out of state pharmacies must have the proper pharmacy permit
413 issued by the state in which they reside in order to utilize a system in Louisiana.~~

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415 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

416 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
417 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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419 **§1229. Violations; Penalties**

420 A. ~~The board may refuse to issue or renew, or may revoke, summarily suspend, suspend, place on
421 probation, censure, reprimand, issue a warning against, or issue a cease and desist order against, the
422 licenses or the registration of, or assess a fine/civil penalty or costs/administrative costs against any
423 person pursuant to the procedures set forth in [R.S. 37:1245](#), for any violation of the provisions of this
424 Section.~~

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426 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

427 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
428 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
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430 **~~§1231. Revised Statutes and Louisiana Administrative Code~~**

431 A. ~~These regulations shall be read and interpreted jointly with [Chapter 14 of Title 37 of the Revised](#)~~
432 ~~[Statutes](#) and Part LIII of Title 46 of the Louisiana Administrative Code.~~

433
434 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

435 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
436 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR

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