

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.

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. 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.

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

62
 63 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

64 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 65 (June 2000) effective July 1, 2000, amended LR 38:1235 (May 2012).
 66

67 **§1203. Registrations** *{A suggested replacement section}*

68 A. Requirement for Registration

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

85 B. Eligibility for Registration

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

97 C. Application for Initial Issuance of Registration

- 98 1. The board shall develop an application form suitable for the AMS registration. The board may
 99 revise that application form on its own initiative in order to collect the information it deems
 100 necessary to properly evaluate an applicant.
 101 2. The application shall be accompanied by payment of the registration fee authorized by R.S.
 102 37:1184.
 103 3. The board shall not process applications received by facsimile, or that are incomplete, or
 104 submitted with the incorrect fee.

- 105 4. The submission of a false or fraudulent application shall substantiate a violation of R.S.
106 37:1241(A)(2) and shall subject the applicant to disciplinary action by the board.
107 5. When determined appropriate by the board, the applicant may be required to meet with a
108 committee or agent of the board prior to the issuance of the registration.
109 D. Maintenance of Registration
110 1. A registration shall be valid only for the pharmacy to which it was issued and the physical location
111 of the AMS identified on the application. The registration shall not be subject to sale, assignment
112 or other transfer, voluntary or involuntary, nor shall the registration be valid for any premises
113 other than the physical location for which it was issued.
114 2. A duplicate or replacement registration shall be issued upon the written request of the owner of the
115 registration and payment of the fee authorized by R.S. 37:1184. A duplicate or replacement
116 registration shall be marked as such, and it shall not serve or be used as an additional or second
117 registration.
118 3. In the event a pharmacy intends to relocate an automated medication system to a different address,
119 the pharmacy shall notify the board of its intent to do so, providing both current and new
120 addresses. A change in business address may require an inspection by the board or its designee.
121 E. Application for Renewal of Registration
122 1. The pharmacy shall complete an application for the renewal of the registration and submit it to the
123 board prior to the expiration date of the registration. The application shall be accompanied by the
124 fee authorized by R.S. 37:1184.
125 2. The board shall not process applications received by facsimile, or that are incomplete, or
126 submitted with the incorrect fee.
127 3. An AMS registration not renewed by the expiration date shall be classified as expired. The
128 operation of an automated medication system with an expired registration shall substantiate a
129 violation of R.S. 37:1241(A)(12) and shall subject the pharmacy to disciplinary action by the
130 board.
131 F. Relinquishment of Registration
132 1. In the event a pharmacy intends to cease supplying medications or devices to an automated
133 medication system, it shall relinquish the registration to the board no later than 10 days following
134 the effective date of such decision.
135 2. A pharmacy may not transfer a registration to another pharmacy.
136 G. Application for Reinstatement of Suspended or Revoked Registration
137 1. An application for the reinstatement of an AMS registration previously suspended or revoked by
138 the board may only be approved in compliance with R.S. 37:1249.
139 2. The applicant shall complete an application form for this specific purpose supplied by the board
140 and shall attach any documentation and fees identified in R.S. 37:1184.
141

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

143 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
144 (June 2000) effective July 1, 2000, amended LR 38:1235 (May 2012), amended by the Department of Health, Board
145 of Pharmacy, LR
146

147 §1205. Pharmacist-in-Charge Responsibilities

- 148 A. The pharmacist-in-charge shall be a Louisiana licensed pharmacist ~~and has~~ with the following
149 responsibilities:
150 1. assuring that the system is in good working order and accurately provides the correct strength,
151 dosage form, and quantity of the drug prescribed while maintaining appropriate record-
152 keeping and security safeguards.
153 2. establishment of a quality assurance program prior to implementation of a system and the
154 supervision of an ongoing quality assurance program that monitors appropriate use and
155 performance of a system, which is evidenced by ~~written~~ policies and procedures developed by
156 the pharmacist-in-charge.
157 3. ~~provide 30 days written notice to the board of removal of the system.~~

- 158 4. define access to the system in policy and procedures of the pharmacy, in compliance with
 159 state and federal regulations.
 160 5. assign, discontinue, or change access to the system.
 161 6. ensure that access to the medications complies with state and federal regulations as
 162 applicable.
 163 7. ensure that the system is stocked/restocked accurately and in accordance with established
 164 written pharmacy policies and procedures.
 165 8. maintain or have access to all records of documentation specified in this Section for two years
 166 or as otherwise required by law.
 167 9. ~~notify each licensed prescriber that his medication orders/prescriptions are not restricted to the~~
 168 ~~limited number of medications which are stocked within a facility's automated medication~~
 169 ~~system by placing a prominent notice to that effect on the cover of or near the beginning of~~
 170 ~~such patient's medical chart or medical record.~~
 171 10. continuous monitoring and documentation of temperature in the drug storage areas including
 172 a mechanism to alert the pharmacist when defined parameters are out of range as well as an
 173 action plan to address such excursions. A pharmacy's failure to document the integrity of the
 174 drug supply or remediate for excursions as appropriate shall substantiate a violation of R.S.
 175 37:1241(A)(18) and shall subject the pharmacy to disciplinary action by the board.
 176

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

178 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 179 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR
 180

181 §1207. Pharmacist Review

- 182 A. System shall be used in settings that ensure medication orders are reviewed by a pharmacist prior to
 183 administration and in accordance with established policies and procedures and good pharmacy
 184 practice. A policy and procedure ~~protocol~~ shall be adopted for non-profile driven systems to
 185 retrospectively review medications orders which cannot be reviewed prior to medication
 186 administration, ~~as provided in LAC 46:LH.1209.2.~~
 187

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

189 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 190 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR
 191

192 §1209. Policies and Procedures

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

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

- 265 vi. ~~activity.~~
 266 vii. ~~problem.~~
 267 p. ~~medication inventory management.~~
 268 q. ~~staff education and training.~~
 269 r. ~~system set up.~~
 270

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

272 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 273 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
 274

275 §1211. Documentation

- 276 A. Documentation as to type of equipment, serial number, content, policies and procedures and location
 277 shall be maintained ~~on-site~~ in the pharmacy for review by the board. Such documentation shall
 278 include, but is not limited to:
- 279 1. name, address, and permit number of the pharmacy ~~or licensed health care facility~~ and the
 280 location where the system is operational;
 - 281 2. manufacturer's name and model;
 - 282 3. quality assurance policies and procedures to determine continued appropriate use and
 283 performance of the system;
 - 284 4. policies and procedures for system operation, safety, security, accuracy, patient
 285 confidentiality, access, controlled substances, data retention, definitions, downtime
 286 procedures, emergency or first dose procedures, inspection, installation requirements,
 287 maintenance security, quality assurance, medication inventory, staff education and training,
 288 system set-up, and malfunction procedures; and
 - 289 5. security procedures sufficient to prevent unauthorized access or use, prevent the illegal use or
 290 disclosure of protected health information, and comply with any applicable federal or state
 291 regulations.
- 292 B. A current copy of all pharmacy policies and procedures related to the use of the system shall be
 293 maintained at all ~~off-site facility~~ locations where the system is being used, ~~as well as the pharmacy of~~
 294 ~~the pharmacist in charge.~~
 295

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

297 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 298 (June 2000) effective July 1, 2000, amended by the Department of Health, Board of Pharmacy, LR
 299

300 §1213. Records

- 301 A. Records and/or electronic data kept by the system shall meet the following requirements:
- 302 1. All events involving access to the contents of the system shall be recorded electronically.
 - 303 2. In the event controlled substances are stored in the system, the records shall include the
 304 positive identification (as defined in Section 1119 of ~~the Board's rules~~ this Part) of the
 305 personnel retrieving and administering the controlled substances to the patient.
 - 306 3. These internal records shall be maintained for one year by the pharmacist-in-charge and shall
 307 be readily available to the board. Such records shall include:
 - 308 a. identity of system accessed;
 - 309 b. identification of the individual accessing the system;
 - 310 c. type of transaction;
 - 311 d. name, strength, dosage form, and quantity of the drug accessed;
 - 312 e. name or identification number of the patient, ~~or identification numbers~~ for whom
 313 the drug was ordered;
 - 314 f. identification of the certified pharmacy technician or pharmacist stocking or
 315 restocking the medications in the system; and
 - 316 g. such additional information as the pharmacist-in-charge may deem necessary.
- 317

318 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.
 319 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 320 (June 2000) effective July 1, 2000, amended LR 40:2256 (November 2014), effective January 1, 2015, amended by
 321 the Department of Health, Board of Pharmacy, LR
 322

323 **§1215. Security System(s)**

- 324 A. ~~System shall have adequate security system and procedures, evidenced by written pharmacy policies~~
 325 ~~and procedures, to:~~
 326 1. ~~prevent unauthorized access or use;~~
 327 2. ~~comply with any applicable federal and state regulations; and~~
 328 3. ~~maintain patient confidentiality.~~
 329

330 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.
 331 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 332 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
 333

334 **§1217. Stocking and Restocking**

- 335 A. ~~On Site Facility System(s). The stocking and restocking of all medications in the on-site system shall~~
 336 ~~be accomplished by Louisiana licensed pharmacists, and/or Louisiana certified pharmacy technicians~~
 337 ~~under the supervision of Louisiana licensed pharmacists. A pharmacist must conduct final checks of~~
 338 ~~work performed by a pharmacy technician. The pharmacy shall have a mechanism in place to identify~~
 339 ~~the certified pharmacy technician stocking or restocking and the pharmacist checking the accuracy of~~
 340 ~~the medications to be stocked or restocked in the automated medication systems.~~
 341 B. ~~Off Site Facility System(s). The stocking and restocking of all medications in the off-site system shall~~
 342 ~~be accomplished by Louisiana licensed pharmacists; however, the certified pharmacy technician may~~
 343 ~~stock or restock an off-site facility system provided a pharmacist is physically present at the off-site~~
 344 ~~facility and supervises and verifies the stocking and/or restocking prior to use. The pharmacy shall~~
 345 ~~have a mechanism in place to identify the certified pharmacy technician stocking or restocking and the~~
 346 ~~pharmacist checking the accuracy of the medications to be stocked or restocked in the system.~~
 347 A. The stocking and restocking of medications and devices within an automated medication system shall
 348 be performed by a pharmacist, or in the alternative, a pharmacy intern, pharmacy technician, or
 349 pharmacy technician candidate under the supervision of a pharmacist.
 350 B. When the pharmacy employs electronic product verification procedures as described within this
 351 Subsection, the stocking and restocking of medications and devices within an automated medication
 352 system may be performed by other personnel approved by the pharmacist-in-charge.
 353 C. Electronic Product Verification
 354 1. A bar code verification, electronic verification, or similar verification process which prohibits
 355 any human intervention following pharmacist verification of the product may be utilized to
 356 assure the correct selection of drugs to be placed into an automated medication system.
 357 2. The use of a bar code, electronic, or similar verification process shall require an initial quality
 358 assurance validation followed by ongoing quality assurance reviews at intervals no greater
 359 than 90 days since the previous review, all conducted by a pharmacist.
 360 3. ~~When a bar code verification, electronic verification, or similar verification process is utilized~~
 361 ~~as specified in the Paragraph, and in the absence of any human intervention in the product~~
 362 ~~selection process, the stocking and restocking functions in systems located either on-site or~~
 363 ~~off-site may be performed by a pharmacy technician without the necessity of direct~~
 364 ~~pharmacist supervision.~~
 365

366 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.
 367 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
 368 (June 2000) effective July 1, 2000, amended LR 41:1488 (August 2015), amended by the Department of Health,
 369 Board of Pharmacy, LR
 370

371 **§1219. Packaging and Labeling**

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

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

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

379
380 **§1221. Proof of Use**

381 A. ~~For medication removed from the system for patient administration, the system shall document, at a~~
382 ~~minimum, the following:~~
383 1. ~~name of the patient or resident;~~
384 2. ~~patient's or resident's medical record number or identification number, or room and bed~~
385 ~~number;~~
386 3. ~~date and time medication was removed from the system;~~
387 4. ~~name, initials, or other unique identifier of the person removing the drug; and~~
388 5. ~~name, strength, and dosage form of the medication or description of the medical device~~
389 ~~removed.~~

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

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

394
395 **§1223. Wasted, Discarded, or Unused Medications**

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

399
400 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

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

403
404 **§1225. Inspection**

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

407
408 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

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

411
412 **§1227. Out of State Pharmacies**

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

416
417 AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.A.

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

420
421 **§1229. Violations; Penalties**

422 A. ~~The board may refuse to issue or renew, or may revoke, summarily suspend, suspend, place on~~
423 ~~probation, censure, reprimand, issue a warning against, or issue a cease and desist order against, the~~

424 licenses or the registration of, or assess a fine/civil penalty or costs/administrative costs against any
425 person pursuant to the procedures set forth in [R.S. 37:1245](#), for any violation of the provisions of this
426 Section.
427

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

429 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
430 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
431

432 ~~§1231. Revised Statutes and Louisiana Administrative Code~~

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

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

436 HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 26:1271
437 (June 2000) effective July 1, 2000, repealed by the Department of Health, Board of Pharmacy, LR
438

