1		Louisiana Administrative Code
2 3		Title 46 - Professional and Occupational Standards
4 5		Part LIII: Pharmacists
6		
7 8	Chapte	er 25. Prescriptions, Drugs, and Devices
9	§2511.	Prescriptions and Chart Orders
10	A. De	finitions. As used in this Chapter, the following terms shall have the meaning ascribed to them in this
11	Se	ction:
12		* * *
13		tic Prescription – a prescription generated, signed and transmitted in electronic form which complies with
14		isions of 21 CFR 1311 or its successor, excluding electronically transmitted facsimile documents.
15	<i>Practice</i>	Affiliation—a practice relationship, collaboration, or practice under the supervision of a physician licensed
16	to practi	<del>ce medicine.</del>
17	<u>Practitio</u>	oner – an individual currently licensed, registered, or otherwise authorized by the appropriate licensing
18	board to	prescribe and administer drugs in the course of professional practice.
19		* * *
20	B. Pat	tient Authority to Acquire Prescription Drug
21	<u>1.</u>	A prescription or chart order represents the lawful authority for a person to acquire a prescription drug
22		from a pharmacy licensed to dispense prescription drugs.
23	<u>2.</u>	In the absence of refill instructions on the original prescription, the prescription shall not be refilled.
24		A pharmacy may dispense the total quantity authorized in one transaction, or in the alternative, may
25		dispense partial quantities in multiple transactions, provided however that the sum of the partial quantities
26		shall not exceed the total quantity authorized.
27	<u>3.</u>	In the event a prescription contains refill instructions, the prescription may be refilled when requested by
28		the person or caregiver. A pharmacy may dispense the quantity authorized for each refill in a single
29		transaction, or in the alternative, may dispense partial quantities in multiple transactions, provided
30		however that the sum of the partial quantities shall not exceed the total quantity authorized.
31	<u>4.</u>	While the documentation of a prescription or chart order shall be retained by the dispensing pharmacy as
32		evidence of its lawful dispensing of the prescription drug, the person's lawful authority to obtain the drug
33		conveyed by the prescription or chart order shall continue to exist until the earliest of the expiration date
34		of the prescription or chart order, or in the alternative, when the total quantity authorized has been
35		dispensed.
36	<u>5.</u>	In the event a person requests a pharmacy to transfer an unfilled prescription to another pharmacy, the
37		pharmacy shall comply with that request as soon as possible, but no later than the end of the next business
38		day. In the event a person requests a pharmacy to transfer the remainder of an unexpired prescription to

39			anoth	ner pharmacy, the pharmacy shall transfer that prescription information in compliance with the
40			provi	isions of this Chapter as soon as possible but no later than the end of the next business day. A
41			phari	macy shall not cancel the remainder of an unexpired prescription unless such action is required by
42			law c	or rule or is requested by the prescriber.
43	<u>C.</u>	Pra	ctitione	ers Authorized to Issue Prescriptions and Chart Orders
44		1.	A ph	armacist may dispense a prescription drug or device when the prescription or chart order is issued
45			by ar	ny of the following practitioners.
46			a.	advanced practice registered nurse
47			b.	<u>dentist</u>
48			<u>c.</u>	medical psychologist
49			<u>d</u> .	<u>optometrist</u>
50			<u>e.</u>	physician
51			<u>f.</u> 1	physician assistant
52			g.	<u>podiatrist</u>
53			h.	<u>veterinarian</u>
54		2.	A pre	escription may be prepared by the secretary or agent of the prescriber for the signature of the
55			presc	criber, but the prescriber retains accountability for the proper issuance of a valid prescription. A
56			presc	criber's secretary or agent may communicate a valid prescription to a pharmacy.
57		3.	In the	e event a practitioner's authority to issue prescriptions is restricted by his licensing authority, the
58			pharr	macist shall dispense any prescriptions issued by that practitioner according to such restriction(s).
59		<u>4.</u>	In the	e event a pharmacist receives a prescription issued by a practitioner in another jurisdiction, the
60			phari	macist may dispense a prescription issued in conformance with the requirements of that jurisdiction
61			How	ever, a prescription issued by a practitioner in another jurisdiction not in conformance with the
62			<u>requi</u>	rements of that jurisdiction shall not be considered a valid prescription in this state, and the
63			phari	macist shall not dispense medication pursuant to an invalid prescription.
64	<u>₿</u> <u>D</u>	. Re	quirem	ents Required Information.
65		<u>1</u> .	A pro	escription shall contain the following data elements:
66			<u>1 a</u> .	Prescriber's name, licensure designation, address, telephone number, and if for a controlled
67				substance, the Drug Enforcement Administration (DEA) registration number;
68			<u>2 b</u> .	Patient's name, and if for a controlled substance, address;
69			<u>3 c</u> .	Date prescription issued by the prescriber;
70			4 <u>d</u> .	Name of drug or device, and if applicable, strength, and quantity to be dispensed;
71			<u>5 e</u> .	Directions for use;
72			<u>6 f</u> .	Signature of the prescriber; and
73			7 <u>g</u> .	Refill instructions, if any. In the absence of refill instructions on the original prescription, the
74				prescription shall not be refilled

75	<u>2</u>	. <u>In tl</u>	ne event a pharmacist receives a prescription order or chart order lacking certain required information,
76		the	pharmacist may consult with the prescriber to clarify the prescriber's intent. Following a consultation
77		with	the prescriber and the appropriate documentation thereof on the order:
78		<u>1 a</u> .	A pharmacist may add the following data elements on the order:
79			a <u>i</u> . <u>Patient's address; or</u>
80			b ii. Drug dosage form.
81		<u> 2 b</u> .	A pharmacist may record changes in the following data elements on the order:
82			a <u>i.</u> <u>Patient's address;</u>
83			b ii. Drug strength;
84			e iii. Quantity prescribed; or
85			d iv. <u>Directions for use.</u>
86		<u>3 c</u> .	A pharmacist shall never add or make changes to the following data elements on the order:
87			a <u>i</u> . <u>Patient's name;</u>
88			b ii. Date of issue;
89			e iii. Drug name (except for generic interchange as permitted by law); or
90			d <u>iv</u> . <u>Prescriber signature.</u>
91	<u>E. N</u>	lanner o	of Issuance
92	<del>D.</del> <u>1.</u>	Oral	Prescriptions.
93		<u> 1 a</u> .	Upon the receipt of an oral prescription from an authorized prescriber, the pharmacist or pharmacy
94			intern or pharmacy technician shall reduce the order to a written form prior to dispensing the
95			medication. As an alternative to recording such prescriptions on paper forms, a pharmacist may enter
96			the prescription information directly into the pharmacy's dispensing information system.
97		<u>b.</u>	In the event a pharmacy intern or pharmacy technician transcribes such a prescription, the supervising
98			pharmacist shall initial or countersign the prescription form prior to processing the prescription.
99	<del>C.</del> <u>2.</u>	Writ	ten Prescriptions. A written prescription shall conform to the following format:
100		1 <u>a.</u>	The prescription form shall be of a size not less than 4 inches by 5 inches, and shall bear a single
101			printed signature line.
102		<u>2 b</u> .	The prescription form shall clearly indicate the authorized prescriber's name, licensure designation,
103			address, telephone number, and if for a controlled substance, the Drug Enforcement Administration
104			(DEA) registration number. In the event that multiple practitioners are identified on the prescription
105			form, the authorizing prescriber's specific identity shall be clear and unambiguous. This
106			identification may be indicated by any means, including but not limited to, a marked check box next
107			to, or circling, the authorized prescriber's printed name.
108		<u>3 c</u> . ]	No prescription form shall contain more than four <u>active</u> prescription drug orders. Each active
109		j	prescription drug order on the form shall provide the following:
110		ŧ	<u>a</u> <u>i</u> . check box labeled "Dispense as Written", or "DAW", or both; and

111	$\frac{b}{ii}$ . the number of refills, if any.
112	4 d. The prescription shall be written with ink or indelible pencil, typewriter, or printed on a computer
113	printer and shall be manually signed by the practitioner on the date issued and in the same manner as
114	he would sign a check or legal document (e.g., J. H. Smith or John H. Smith). Examples of invalid
115	signatures include rubber stamps, signatures of anyone other than the prescriber, and computer-
116	generated signatures.
117	5 <u>e</u> . Receipt via Facsimile
118	a i. Pharmacies may elect to receive written prescriptions via a facsimile machine located within
119	the prescription department. The paper used to print such prescriptions shall produce a non-
120	fading image. The pharmacy may elect to scan such documents in compliance with Section
121	1123 of this Part.
122	b ii. Pharmacies may elect to receive written prescriptions via electronic facsimile directly within
123	their pharmacy information system. The pharmacy shall retain such records in compliance with
124	Section 1123 of this Part.
125	6 f. Chart orders and forms used by pharmacists to record telephoned or transferred prescriptions are
126	exempt from the format requirements listed above in this Section.
127	D. Oral Prescriptions.
128	1. Upon the receipt of an oral prescription from an authorized prescriber, the pharmacist or pharmacy intern
129	or pharmacy technician shall reduce the order to a written form prior to dispensing the medication. As an
130	alternative to recording such prescriptions on paper forms, a pharmacist may enter the prescription
131	information directly into the pharmacy's dispensing information system. In the event a pharmacy intern
132	or pharmacy technician transcribes such a prescription, the supervising pharmacist shall initial or
133	countersign the prescription form prior to processing the prescription.
134	E. 3. Electronic Prescriptions.
135	4 a. The prescription shall clearly indicate the authorized prescriber's name, licensure designation,
136	address, telephone number, and if for a controlled substance, the DEA registration number.
137	F. Completion of Prescription Orders and Chart Orders. In the event a pharmacist receives a prescription order or
138	chart order lacking certain required information, the pharmacist may consult with the prescriber to clarify the
139	prescriber's intent. Following a consultation with the prescriber and the appropriate documentation thereof on
140	the order:
141	1. A pharmacist may add the following data elements on the order:
142	a. Patient's address; or
143	b. Drug dosage form.
144	2. A pharmacist may record changes in the following data elements on the order:
145	a. Patient's address;
146	b. Drug strength;

147	c. Quantity prescribed; or
148	d. Directions for use.
149	3. A pharmacist shall never add or make changes to the following data elements on the order:
150	a. Patient's name;
151	b. Date of issue;
152	c. Drug name (except for generic interchange as permitted by law); or
153	d. Prescriber signature.
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155 156 157 158 159 160	AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182.  HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 14:708 (October 1988) effective January 1, 1989, amended LR 29:2102 (October 2003), effective January 1, 2004, LR 41:98 (January 2015), LR 41:2147 (October 2015), amended by the Department of Health, Board of Pharmacy, LR 43:2162 (November 2017), LR 46:585 (April 2020), LR.
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## §2519. Prescription Refills; Medication Synchronization and Refill Consolidation

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- B. <u>Dispensing of Refills Requests</u>. Prescription refills authorized by the prescriber shall not be dispensed in the absence of a patient or caregiver's request or approval. This prohibition shall not apply to refills authorized by the prescriber which are to be dispensed to a patient residing in a long-term care facility.
- C. A pharmacy submitting a request for refill authorization to a prescriber shall differentiate between those originating from a specific patient inquiry from those originating from the pharmacy staff.
- 170 C D. Controlled Dangerous Substances.

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- 1. The refilling of a prescription for a drug listed in Schedule II is prohibited.
- 2. A prescription for a drug listed in Schedule III or IV may be refilled up to five times, if so indicated at the time issued.
- 3. A prescription for a drug listed in Schedule V may be refilled if so indicated at the time issued subject to the one year expiration date of the prescription.
- <u>P</u> <u>E</u>. Medication Synchronization and Refill Consolidation. These terms refer to a service which a pharmacist may perform for his patient, at the request of the patient, wherein he may proactively adjust the medication dispensing quantity and/or the refill schedule of a prescription in order to manage the patient's medication therapy, with the goal of improved medication adherence by the patient.
  - For the performance of this service, the pharmacist may adjust the dispensing quantity and/or the refill
    schedule originally ordered by the prescriber; however, the pharmacist shall not exceed the total quantity
    prescribed [dispensing quantity multiplied by the total number of fills authorized (original plus refills)], or
    what is otherwise allowed by law.
  - 2. With respect to prescriptions for controlled substances where refills have been authorized, pharmacists

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185	may utilize partial fills, as described in Paragraph 2747.C.5 of this Part, but may not exceed the
186	dispensing quantity noted on the original prescription.
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188 189 190 191 192	AUTHORITY NOTE: Promulgated in accordance with R.S. 37:1182. HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Board of Pharmacy, LR 14:708 (October 1988), effective January 1, 1989, amended LR 29:2104 (October 2003), effective January 1, 2004, LR 33:1133 (June 2007), amended by the Department of Health, Board of Pharmacy, LR 42:1519 (September 2016), LR 46:575 (April 2020), LR

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