1	Louisiana Administrative Code
2 3	Title 46 – Professional and Occupational Standards
4 5	Part LIII: Pharmacists
6 7	Chapter 24. Limited Service Providers
8 9	Subchapter E. Marijuana Pharmacy
10 11	§2440. Preamble; Warning; Consultation Suggested
12	A. Pursuant to Act 261 of the Regular Session of the 2015 Louisiana Legislature as well as the subsequent
13	amendments found in Act 96 of the Regular Session of the 2016 Louisiana Legislature, the Board of Pharmac
14	was directed to:
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16	2. Adopt rules relating to the dispensing of recommended marijuana for therapeutic use, with such rules to
17	include, at a minimum, the following:
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19	e. Standards, procedures, and protocols to ensure all recommended therapeutic marijuana dispensed,
20	with the exception of raw or crude marijuana product, is consistently pharmaceutical grade;
21	A.2.f. – C
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23	AUTHORITY NOTE: Promulgated in accordance with R.S. 40:1046.
24	HISTORICAL NOTE: Promulgated by the Department of Health, Board of Pharmacy, LR 43:1538 (August 2017),
25	amended LR
26	* * *
27	§2443. Marijuana Products
28	A. Exclusive Source.
29	1. The exclusive source of marijuana products shall be the producers licensed for that activity by the
30	Department of Agriculture and Forestry (LDAF).
31	2. That Licensed producers shall prepare pharmaceutical grade marijuana products as well as raw marijuana
32	products for distribution to the marijuana pharmacies licensed by the board.
33	3. Marijuana products from any other source shall be deemed misbranded and/or adulterated and shall not be
34	distributed to any marijuana pharmacy, nor may such misbranded and/or adulterated products be
35	dispensed by any marijuana pharmacy.
36	B. – B.8 {Laboratory testing standards from LDAF?}
37	C. Product Dosage Forms.

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38 1. The producer shall limit their production of pharmaceutical grade marijuana products to the following 39 dosage forms: 40 Oils, extracts, tinctures, or sprays; 41 Solid oral dosage forms, e.g., capsules or pills; Liquid oral dosage forms, e.g., solutions or suspensions; 42 43 Gelatin-based or pectin-based chewables; 44 Topical applications, oils or lotions; 45 Transdermal patches; 46 Suppositories; 47 Metered-dose inhalers. The producer may produce other products from raw or crude marijuana, including dried flower, buds and 48 49 other plant material, intended for the following methods of administration: 50 Combustible forms for inhalation, including but not limited to pre-rolls; and 51 Edible forms for ingestion. 52 2 3. No marijuana product shall: 53 include alcoholic liquor, dietary supplements, or any drug, except for pharmaceutical grade 54 marijuana. For purposes of this provision, alcoholic liquor does not include any liquid or solid 55 containing less than 0.5 percent of alcohol by volume, or ethanol-based tinctures. 56 b. be manufactured or sold as a beverage. 57 e b. be manufactured or sold in a form or with a design that: 58 is obscene or indecent; 59 may encourage the use of marijuana for recreational purposes; 60 iii. may encourage the use of marijuana for a condition other than a debilitating medical condition; 61 62 iv. is customarily associated with persons under the age of 18 years; or 63 d c. have had pesticide chemicals or organic solvents used during the production or manufacturing 64 process other than those which may be approved by the commissioner of LDAF. 65 3.4. Any marijuana product not in compliance with the provisions of this Paragraph Section shall be deemed 66 adulterated. 67 Packaging and Labeling Requirements D. 68 1. Packaging 69 1.a - 1.b.v. ... 70 c. Products containing raw or crude marijuana shall be packaged in such a manner as to facilitate 71 compliance with the dispensing limitations imposed on marijuana pharmacies. 72 D.2. – E.4.f. ...

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74 AUTHORITY NOTE: Promulgated in accordance with R.S. 40:1046. 75 HISTORICAL NOTE: Promulgated by the Department of Health, Board of Pharmacy, LR 43:1540 (August 2017), 76 amended LR 45:1473 (October 2019), LR 46:568 (April 2020), amended LR 46:1227 (September 2020), LR 47:590 77 (May 2021), LR 78 79 §2457. Standards of Practice 80 A. – D.5. ... 81 **Professional Practice Standards** 82 Recommendation / opinion / referral (hereinafter "request") for Therapeutic Marijuana 83 1.a. – 1.b.vii. ... 84 c. Requests for raw or crude marijuana products intended for persons under 21 years of age shall 85 specifically indicate a recommendation for raw or crude forms of marijuana for such persons. 86 e d. Requests for marijuana products shall expire one year after the date of issue, unless a shorter period 87 of time is indicated by the physician. A pharmacist shall not dispense marijuana product pursuant to 88 an expired request. 89 e. Dispensing Limitations 90 i. A pharmacist shall not dispense more than two and one-half ounces, or 71 grams, of raw or 91 crude marijuana every 14 days to any person. 92 ii. Subject to the above limitation on dispensing raw or crude marijuana products, A a pharmacist 93 may dispense marijuana products on multiple occasions as indicated by the physician and 94 needed by the patient until the request expires; however, the pharmacist shall not dispense more 95 than a 90-day supply of marijuana product at one time nor more than a one-year supply 96 pursuant to a single request. 97 E.2. – E.6.e.iv. ... 98 99 AUTHORITY NOTE: Promulgated in accordance with R.S. 40:1046. 100 HISTORICAL NOTE: Promulgated by the Department of Health, Board of Pharmacy, LR 43:1550 (August 2017),

amended LR 45:1473 (October 2019), LR 47:246 (February 2021), LR