

D. No person is authorized to distribute any industrial hemp-derived cannabidiol products in the state of Louisiana unless that person has first obtained a Certificate of IHDCP Registration from the department.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§519. Industrial Hemp-Derived Cannabidiol Products Labeling Requirements: Certificate of Analysis**

A. In addition to the requirements enumerated in R.S. 40:608, industrial hemp-derived cannabidiol products must bear labeling that includes a scannable bar code, QR code, or a web address linked to a document or website containing the certificate of analysis for that product.

B. The certificate of analysis must be from a laboratory that is accredited by LDH/OPH.

C. The certificate of analysis must include, at a minimum, the following information:

1. the batch number of the product;
2. the date the batch was received by the laboratory;
3. the date the testing was completed;
4. the laboratory methodology used for each analysis referenced in the report;
5. the amount of THC by dry weight in milligrams;
6. the amount of CBD by dry weight in milligrams;
7. the amount of any detected residual solvent in the product in parts per million;
8. the amount of any detected pesticide residues in the product in parts per million;
9. the amount of any microbiological contaminants in the product in appropriate units; and
10. the amount of any detected heavy metal traces in the product in parts per million.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§521. Industrial Hemp-Derived Cannabidiol Products Labeling Requirements: Disclaimer**

A. Each primary container of industrial hemp-derived cannabidiol product must bear the following statement: "This product has not been evaluated by the Food and Drug Administration and is not intended to diagnose, treat, cure, or prevent any disease."

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§523. Industrial Hemp-Derived Cannabidiol Products Labeling Requirements: Medical Claims Prohibited**

A. No product labeling or advertising material for any industrial hemp-derived cannabidiol product sold or otherwise distributed in the state of Louisiana may bear any implicit or explicit medical claims.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§525. Industrial Hemp-Derived Cannabidiol Products Labeling Requirements: Dietary Supplements Prohibited**

A. No industrial hemp-derived cannabidiol product may be marketed as a dietary supplement.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§527. Penalties for Violations of Requirements to Register Industrial Hemp-Derived Cannabidiol Products**

A. Any person who violates the provisions requiring registration of industrial hemp-derived cannabidiol products is subject to the penalties provided for by R.S. 3:1484 and other sanctions as provided for by the State Food, Drug, and Cosmetic Law.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**§529. Exemptions**

A. Industrial hemp-derived cannabidiol products that have been produced in accordance with R.S. 40: 1046 or that are Food and Drug Administration (FDA)-approved pharmaceuticals are not subject to the requirements of this regulation.

AUTHORITY NOTE: Promulgated in accordance with R.S. 3:1482(J) and R.S. 40:604.

HISTORICAL NOTE: Promulgated by the Department of Health, Office of Public Health, LR 46:

**Title 51**

**Public Health—Sanitary Code**

**Part VI. Manufacturing, Processing, Packing and Holding of Food, Drugs and Cosmetics**

**Chapter 3. Current Good Manufacturing Practices in Manufacturing, Processing, Packing or Holding Human Food**

**§301. General Provisions; Code of Federal Regulations [formerly paragraph 6:039]**

A. The Criteria in 21 CFR 117 Subpart A, Subpart B and Subpart F (Code of Federal Regulations) shall apply in determining whether the facilities, methods, practices, and controls used in the manufacturing, processing, packing or holding of food are in conformance with or are operated or administered in conformity with good manufacturing practices to assure that food for human consumption is safe and has been prepared, packed and held under sanitary conditions.

B. In accordance with R.S. 3:1468, facilities producing industrial hemp-derived cannabidiol products intended for human consumption will be inspected under the provisions of this Chapter.

AUTHORITY NOTE: Promulgated in accordance with the provisions of R.S. 40:4(A)(1)(a). Also see R.S. 40:601 et seq., and R.S. 3:1482(J).

HISTORICAL NOTE: Promulgated by the Department of Health and Hospitals, Office of Public Health, LR 28:1234 (June 2002), amended by the Department of Health, Office of Public Health, LR 46: