

ACT No. 274

2025 Regular Session

HOUSE BILL NO. 412

BY REPRESENTATIVE ROMERO

1 AN ACT

2 To amend and reenact R.S. 26:926.1, relative to alternative nicotine products; to provide for
3 a directory; and to provide for related matters.

4 Be it enacted by the Legislature of Louisiana:

5 Section 1. R.S. 26:926.1 is hereby amended and reenacted to read as follows:

6 §926.1. Vapor product and alternative nicotine product directory

7 A. Every vapor product manufacturer ~~and alternative nicotine product~~
8 ~~manufacturer~~ whose products are sold in this state, whether directly or through a
9 wholesale dealer, retail dealer, or similar intermediary or intermediaries, shall
10 execute and deliver on a form prescribed by the commissioner a certification to the
11 commissioner affirming, under penalty of perjury, either of the following:

12 (1) The product was on the market in the United States as of August 8, 2016,
13 and the manufacturer has applied for a marketing order pursuant to 21 U.S.C. 387j
14 for the vapor product or alternative nicotine product by submitting a premarket
15 tobacco product application on or before September 9, 2020, to the United States
16 Food and Drug Administration, hereinafter referred to in this Section as "FDA", and
17 either of the following is true:

18 (a) The premarket tobacco product application for the vapor product ~~or~~
19 ~~alternative nicotine product~~ remains under review by the FDA.

20 (b) The FDA has issued a no marketing order for the vapor product ~~or~~
21 ~~alternative nicotine product~~, but the agency or a federal court has issued a stay order
22 or injunction during the pendency of the manufacturer's appeal of the no marketing

1 order, or the order has been appealed either to the FDA or a challenge to the order
2 filed with a federal court and the appeal or challenge is still pending.

3 (2) The manufacturer has received a marketing order or other authorization
4 under 21 U.S.C. 387j for the vapor product ~~or alternative nicotine product~~ from the
5 FDA.

6 B. Every alternative nicotine product manufacturer whose products are sold
7 in this state, whether directly or through a wholesale dealer, retail dealer, or similar
8 intermediary or intermediaries, shall execute and deliver on a form prescribed by the
9 commissioner a certification to the commissioner affirming, under penalty of
10 perjury, either of the following:

11 (1) The product was on the market in the United States as of April 14, 2022,
12 and the manufacturer has applied for a marketing order pursuant to 21 U.S.C. 387j
13 for the vapor product or alternative nicotine product by submitting a premarket
14 tobacco product application on or before May 14, 2022, to the FDA, and either of
15 the following is true:

16 (a) The premarket tobacco product application for the alternative nicotine
17 product remains under review by the FDA.

18 (b) The FDA has issued a no marketing order for the alternative nicotine
19 product, but the agency or a federal court has issued a stay order or injunction during
20 the pendency of the manufacturer's appeal of the no marketing order, or the order has
21 been appealed either to the FDA or a challenge to the order filed with a federal court
22 and the appeal or challenge is still pending.

23 (2) The manufacturer has received a marketing order or other authorization
24 under 21 U.S.C. 387j for the alternative nicotine product from the FDA.

25 ~~B. C.~~ In addition to the requirements of ~~Subsection~~ Subsections A and B of
26 this Section, each manufacturer shall provide a copy of the cover page of the
27 premarket tobacco application with evidence of receipt of the application by the FDA
28 or a copy of the cover page of the marketing order or other authorization issued
29 pursuant to 21 U.S.C. 387j, whichever is applicable.

1 ~~C. D.~~ Any manufacturer submitting a certification pursuant to ~~Subsection A~~
 2 Subsection A or B of this Section shall notify the commissioner within thirty days
 3 of any material change to the certification, including issuance by the FDA of any of
 4 the following:

5 (1) A market order or other authorization pursuant to 21 U.S.C. 387j.

6 (2) An order requiring a manufacturer to remove a product from the market
 7 either temporarily or permanently.

8 (3) Any notice of action taken by the FDA affecting the ability of the new
 9 product to be introduced or delivered into interstate commerce for commercial
 10 distribution.

11 (4) Any change in policy that results in a product no longer being exempt
 12 from federal enforcement oversight.

13 ~~D. E.~~ The commissioner shall develop and maintain a directory listing all
 14 vapor product manufacturers and alternative nicotine product manufacturers that
 15 have provided certifications that comply with ~~Subsection~~ Subsections A and B of this
 16 Section and all products that are listed in those certifications.

17 ~~E. F.~~ The commissioner shall do all of the following:

18 (1) Make the directory available for public inspection on the public website
 19 of the office of alcohol and tobacco control.

20 (2) Update the directory as necessary in order to correct mistakes and to add
 21 or remove vapor product manufacturers and alternative nicotine product
 22 manufacturers or products manufactured by those manufacturers.

23 (3) Send monthly notifications to each wholesale dealer, retail dealer, and
 24 manufacturer of vapor products and manufacturer of alternative nicotine products
 25 that has qualified or registered with the commissioner, by electronic communication,
 26 containing a list of all changes that have been made to the directory in the previous
 27 month. In lieu of sending monthly notifications, the commissioner may make the
 28 information available in a prominent place on the public website of the office of
 29 alcohol and tobacco control.

1 ~~F.~~ G. Notwithstanding ~~Subsection A~~ Subsections A and B of this Section, if
 2 a vapor product manufacturer or alternative nicotine product manufacturer can
 3 demonstrate to the commissioner that the FDA has issued a rule, guidance, or any
 4 other formal statement that temporarily exempts a vapor product or alternative
 5 nicotine product from the federal premarket tobacco application requirements, the
 6 vapor product or alternative product may be added to the directory upon request by
 7 the manufacturer if the manufacturer provides sufficient evidence that the vapor
 8 product or alternative nicotine product is compliant with the federal rule, guidance,
 9 or other formal statement, as applicable.

10 ~~G.~~ H. Each certifying vapor product manufacturer or alternative nicotine
 11 product manufacturer shall pay an initial fee of one hundred dollars per product stock
 12 keeping unit or SKU to offset the costs incurred by the commissioner for processing
 13 the certifications and operating the directory. The commissioner shall collect an
 14 annual renewal fee of one hundred dollars per product stock keeping unit or SKU to
 15 offset the costs associated with maintaining the directory and satisfying the
 16 requirements of this Section. The fees received pursuant to this Section by the
 17 commissioner shall be used by the office of alcohol and tobacco control exclusively
 18 for processing the certifications and operating and maintaining the directory.

19 ~~H.~~ I. Beginning on the date that the commissioner makes the directory
 20 available for public inspection on the public website of the office of alcohol and
 21 tobacco control as provided in ~~Subsection E~~ F of this Section, a vapor product
 22 manufacturer or alternative nicotine product manufacturer who offers for sale a
 23 vapor product or alternative nicotine product not listed on the directory is subject to
 24 a one thousand dollar daily fine for each vapor product or alternative nicotine
 25 product offered for sale in violation of this Section until the offending product is
 26 removed from the market or until the offending product is properly listed on the
 27 directory.

28 ~~F.~~ J.(1) The sale, possession, or transportation of vapor products or
 29 alternative nicotine products not listed on the directory by any person, including a

1 permitted wholesale dealer or retail dealer, shall be subject to provisions of R.S.
2 47:858, 859, and 860 as if such wholesale dealer or retail dealer did not possess a
3 valid permit.

4 (2) Each unit of vapor product or alternative nicotine product sold or offered
5 for sale, possessed, or transported shall constitute a separate violation for purposes
6 of Paragraph (1) of this Subsection.

7 ~~¶~~ K. Any other violation of this Section shall result in a fine of five hundred
8 dollars per offense.

9 ~~¶~~ L. The commissioner shall adopt rules for the implementation and
10 enforcement of this Section.

SPEAKER OF THE HOUSE OF REPRESENTATIVES

PRESIDENT OF THE SENATE

GOVERNOR OF THE STATE OF LOUISIANA

APPROVED: _____