FOR OFFICE USE ONLY	

HOUSE FLOOR AMENDMENTS

2023 Regular Session

Amendments proposed by Representative Hughes to Engrossed House Bill No. 179 by Representative Wheat

AMENDMENT NO. 1

Delete the set of House Floor Amendments by Representative Hughes (#3145)

AMENDMENT NO. 2

On page 1, line 2, after "R.S. 26:911(A)(7)" and before the comma "," insert "and 926"

AMENDMENT NO. 3

On page 1, line 6, after "R.S. 26:911(A)(7)" and before "hereby" delete "is" and insert "and 926 are"

AMENDMENT NO. 4

On page 1, line 13, after "products" insert a comma "," and insert "in each case, only if the e-liquid or vapor products contain nicotine from any source,"

AMENDMENT NO. 5

On page 1, at the end of line 19, insert "This Paragraph shall not apply to any e-liquid or vapor product that has received a marketing order from the United States Food and Drug Administration pursuant to 21 U.S.C. 387(j)."

AMENDMENT NO. 6

On page 1, after line 20, add the following:

"§926. Vapor product and alternative nicotine product directory

A. Beginning October 1, 2023, every vapor product manufacturer and alternative nicotine product manufacturer whose products are sold in this state, whether directly or through a wholesale dealer, retail dealer, or similar intermediary or intermediaries, shall execute and deliver on a form prescribed by the secretary, a certification to the secretary certifying, under penalty of perjury, either of the following:

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

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

(b) The FDA has issued a no marketing order for the vapor product or alternative nicotine product from the FDA; however, the agency or a federal court has issued a stay order or injunction during the pendency of the manufacturer's appeal of the no marketing order.

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

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

C. Any manufacturer submitting a certification pursuant to Subsection A shall notify the secretary within 30 days of any material change to the certification, including issuance by the FDA of any of the following:

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

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

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

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

D. The secretary shall develop and maintain a directory listing all vapor product manufacturers and alternative nicotine product manufacturers that have provided certifications that comply with Subsection A and all products that are listed in those certifications.

E. The secretary shall do all of the following:

(1) Make the directory available for public inspection on its website by November 1, 2023.

(2) Update the directory as necessary in order to correct mistakes and to add or remove vapor product manufacturers and alternative nicotine product manufacturers or products manufactured by those manufacturers consistent with the requirements of Paragraphs (1) and (2) of this Subsection on a monthly basis.

(3) Send monthly notifications to each wholesale dealer, retail dealer, or manufacturer of vapor products and manufacturer of alternative nicotine products that have qualified or registered with the department, by electronic communication, containing a list of all changes that have been made to the directory in the previous month. In lieu of sending monthly notifications, the secretary may make the information available in a prominent place on the department's public website.

F. Notwithstanding Subsection A of this Section, if a vapor product manufacturer or alternative nicotine product manufacturer can demonstrate to the secretary that the FDA has issued a rule, guidance, or any other formal statement that temporarily exempts a vapor product or alternative nicotine product from the federal premarket tobacco application requirements, the vapor product or alternative product may be added to the directory upon request by the manufacturer if the manufacturer

provides sufficient evidence that the vapor product or alternative nicotine product is compliant with the federal rule, guidance, or other formal statement, as applicable.

G. No wholesale dealer or retail dealer shall be permitted to remit tax with respect to a vapor product or alternative nicotine product unless such vapor product or alternative nicotine product is listed on the directory, and the sale, possession, or transportation of such vapor products or alternative nicotine products by any person, including a permitted wholesale dealer or retail dealer, shall be subject to provisions of R.S. 47:858. 859, and 860 as if such wholesale dealer or retail dealer did not possess a valid permit.

H. The secretary shall adopt rules for the implementation and enforcement of this Section."