

118TH CONGRESS
1ST SESSION

H. R. 901

To require the Food and Drug Administration to prioritize enforcement of disposable electronic nicotine delivery system products.

IN THE HOUSE OF REPRESENTATIVES

FEBRUARY 9, 2023

Mrs. CHERFILUS-McCORMICK introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

To require the Food and Drug Administration to prioritize enforcement of disposable electronic nicotine delivery system products.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Disposable ENDS
5 Product Enforcement Act of 2023”.

6 **SEC. 2. FINDINGS.**

7 Congress finds the following:

8 (1) In April 2020, the Food and Drug Adminis-
9 tration issued guidance entitled “Enforcement Prior-
10 ities for Electronic Nicotine Delivery System

1 (ENDS) and Other Deemed Products on the Market
2 Without Premarket Authorization”.

3 (2) In such guidance, Food and Drug Adminis-
4 tration describes how the agency intends to prioritize
5 enforcement against any flavored, cartridge-based
6 electronic nicotine delivery system product (other
7 than a tobacco- or menthol-flavored electronic nico-
8 tine delivery system product) marketed without au-
9 thorization from the Food and Drug Administration.

10 (3) In defining the term “cartridge-based
11 ENDS product,” the guidance cites “self-contained,
12 disposable products” as not being within the scope of
13 this prioritized category.

14 (4) The guidance explains: “FDA is continu-
15 ously evaluating new information and adjusting its
16 enforcement priorities in light of the best available
17 data, and it will continue to do so with respect to
18 these products. FDA will take appropriate action re-
19 garding tobacco products that are marketed without
20 premarket authorization, including as warranted
21 based on changed circumstances, new information,
22 or to better address minors’ use of those products”.

23 (5) In November 2022, the Food and Drug Ad-
24 ministration and the Centers for Disease Control
25 and Prevention released the findings from the 2022

1 National Youth Tobacco Survey. The data shows
2 that self-contained disposable electronic nicotine de-
3 livery system products were the most common device
4 type used by minors.

5 **SEC. 3. UPDATED ENFORCEMENT PRIORITIZATION.**

6 (a) GUIDANCE.—The Secretary of Health and
7 Human Services, acting through the Commissioner of
8 Food and Drugs (referred to in this Act as the “Sec-
9 retary”), shall not later than 90 days after the date of
10 the enactment of this Act, update the final guidance enti-
11 tled “Enforcement Priorities for Electronic Nicotine De-
12 livery System (ENDS) and Other Deemed Products on the
13 Market Without Premarket Authorization” issued in April
14 2020, to include a description of how the Secretary will
15 also prioritize enforcement against disposable ENDS
16 products, including such nicotine products not derived
17 from tobacco.

18 (b) ENFORCEMENT.—Nothing in this section shall be
19 construed as preventing the Secretary from prioritizing
20 enforcement against disposable ENDS products, including
21 nicotine products not derived from tobacco, in advance of
22 updating the guidance referred to in paragraph (1), par-
23 ticularly with respect to such products that are targeted
24 to minors or whose marketing is likely to promote use of
25 such products by minors.

1 (c) INCLUSION IN TOBACCO REGULATION ACTIVITIES
2 ANNUAL REPORT.—The Secretary shall include in each
3 annual report required to be submitted pursuant to section
4 112 of subtitle B of title I of division P of the Consolidated
5 Appropriations Act, 2022 (Public Law 117–103), the total
6 number of compliance and enforcement actions taken with
7 respect to disposable ENDS products during the year cov-
8 ered by the report.

9 (d) DISPOSABLE ENDS PRODUCT DEFINED.—In
10 this Act, the term “disposable ENDS product” means a
11 tobacco product (as defined by section 201(rr) of the Fed-
12 eral Food, Drug, and Cosmetic Act (21 U.S.C. 321(rr)))
13 that consists of a single unit including the battery and
14 the liquid, and the entire device is intended to be disposed
15 of after the liquid has been depleted.

○