

By: LaMantia

S.R. No. 79

R E S O L U T I O N

1           WHEREAS, In January 2020, the Food and Drug Administration  
2 issued industry guidance outlining its enforcement priorities  
3 regarding e-cigarettes, vapes, and other electronic nicotine  
4 delivery system products; and

5           WHEREAS, This guidance emphasized the agency's focus on  
6 enforcing regulations against all flavored, cartridge-based ENDS  
7 products, but it unintentionally created a loophole, exempting all  
8 disposable vaping products from enforcement for nearly three years;  
9 and

10           WHEREAS, Since January 2020, a significant influx of illicit  
11 disposable vaping products has entered the United States; these  
12 originate primarily in China and mostly feature flavors designed to  
13 appeal to children; and

14           WHEREAS, According to the Centers for Disease Control and  
15 Prevention's National Youth Tobacco Survey, youth consumption of  
16 disposable vaping products has surged by an alarming 2,188 percent  
17 since 2019; a substantial majority of disposable vaping products  
18 introduced to the market have either entered after the FDA's  
19 regulatory submission cutoff date of August 8, 2016, or have failed  
20 to comply with the FDA's regulatory pathways designed to ensure  
21 that products available in stores promote public health and safety;  
22 the urgency of this matter is reinforced by the fact that, in 2021,  
23 46 percent of high school e-cigarette users vaped at least 20 days a  
24 month, and over 30 percent reported vaping every day; and

1           WHEREAS, The continued rise in youth vaping is a serious  
2 public health concern that demands immediate attention and action;  
3 much-needed measures include the publication of a directory of  
4 disposable vapor products that can be sold subject to FDA  
5 enforcement discretion, allowing retailers to remove all illegal  
6 disposable vapor products from shelves, as well as the allocation  
7 of resources for federal enforcement across all jurisdictions and  
8 at all ports and border control points of entry; now, therefore, be  
9 it

10           RESOLVED, That the Senate of the 88th Texas Legislature, 3rd  
11 Called Session, hereby urge the Food and Drug Administration to  
12 provide clear enforcement guidance regarding the distribution and  
13 sales of disposable vapor products; and, be it further

14           RESOLVED, That the Secretary of the Senate forward official  
15 copies of this resolution to the President of the United States, to  
16 the Speaker of the House of Representatives and the President of the  
17 Senate of the United States Congress, to the Commissioner of the  
18 FDA, and to all the members of the Texas delegation to Congress with  
19 the request that this resolution be officially entered in the  
20 Congressional Record as a memorial to the Congress of the United  
21 States of America.