S 3653

Resources To Prevent Youth Vaping Act

Congress: 118 (2023–2025, Ended)
Chamber: Senate
Policy Area: Health
Introduced: Jan 24, 2024
Current Status: Read twice and referred to the Committee on Health, Education, Labor, and Pensions.
Latest Action: Read twice and referred to the Committee on Health, Education, Labor, and Pensions. (Jan 24, 2024)
Official Text: https://www.congress.gov/bill/118th-congress/senate-bill/3653

Sponsor

Name: Sen. Shaheen, Jeanne [D-NH]
Party: Democratic • **State:** NH • **Chamber:** Senate

Cosponsors (6 total)

Cosponsor	Party / State	Role	Date Joined
Sen. Baldwin, Tammy [D-WI]	D · WI		Jan 24, 2024
Sen. Collins, Susan M. [R-ME]	R · ME		Jan 24, 2024
Sen. Durbin, Richard J. [D-IL]	D · IL		Jan 24, 2024
Sen. Murkowski, Lisa [R-AK]	R · AK		Jan 24, 2024
Sen. Romney, Mitt [R-UT]	R · UT		Jan 24, 2024
Sen. Blumenthal, Richard [D-CT]	D · CT		Jul 8, 2024

Committee Activity

Committee	Chamber	Activity	Date
Health, Education, Labor, and Pensions Committee	Senate	Referred To	Jan 24, 2024

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Summary (as of Jan 24, 2024)

Resources to Prevent Youth Vaping Act

This bill directs the Food and Drug Administration (FDA) to collect user fees on products that it deems by regulation to be tobacco products, including electronic nicotine delivery systems, and addresses related issues. Currently, the FDA is authorized to collect user fees only on specified classes of tobacco products.

The bill increases the total amount of such fees to be collected for FY2025. For each fiscal year after, the total amount of such fees shall be adjusted according to changes in a price index.

Starting in FY2027, the FDA must assess user fees on classes of products that it has deemed by regulation to be tobacco products, unless the FDA fails to finalize a formula for assessing such fees on time. Once it is finalized, the FDA may only revise this formula by regulation.

The bill also requires each tobacco manufacturer and importer to periodically submit certain information related to the tobacco products that it sells or distributes in the United States.

The FDA must annually report to Congress about its use of such tobacco product fees.

Actions Timeline

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