

HR 1463

To prohibit the use of Federal funds to implement, administer, or enforce a final rule of the Food and Drug Administration relating to "Medical Devices; Laboratory Developed Tests", and for other purposes.

Congress: 119 (2025–2027, Current)

Chamber: House

Policy Area: Health

Introduced: Feb 21, 2025

Current Status: Referred to the House Committee on Energy and Commerce.

Latest Action: Referred to the House Committee on Energy and Commerce. (Feb 21, 2025)

Official Text: <https://www.congress.gov/bill/119th-congress/house-bill/1463>

Sponsor

Name: Rep. Finstad, Brad [R-MN-1]

Party: Republican • **State:** MN • **Chamber:** House

Cosponsors (7 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Crenshaw, Dan [R-TX-2]	R · TX		Feb 21, 2025
Rep. Miller-Meeks, Mariannette [R-IA-1]	R · IA		Feb 24, 2025
Rep. Weber, Randy K. Sr. [R-TX-14]	R · TX		Feb 27, 2025
Rep. Pfluger, August [R-TX-11]	R · TX		Mar 4, 2025
Rep. Dunn, Neal P. [R-FL-2]	R · FL		Mar 21, 2025
Rep. Barr, Andy [R-KY-6]	R · KY		Apr 1, 2025
Rep. Goldman, Craig A. [R-TX-12]	R · TX		Aug 8, 2025

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred To	Feb 21, 2025

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Summary (as of Feb 21, 2025)

This bill prohibits the use of federal funds to implement, administer, or enforce the final rule issued by the Food and Drug Administration (FDA) titled *Medical Devices; Laboratory Developed Tests* and published on May 6, 2024, or any substantially similar rule.

The rule specifies that laboratory developed tests (medical screening tests developed by laboratories rather than medical device manufacturers) are medical devices subject to FDA regulation with limited exceptions. Prior to the rule, the FDA had exercised discretion as to whether to apply regulations for medical devices to laboratory developed tests.

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