

HR 1108

Diagnostics Testing Preparedness Plan Act of 2025

Congress: 119 (2025–2027, Current)

Chamber: House

Policy Area: Health

Introduced: Feb 7, 2025

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

Latest Action: Referred to the House Committee on Energy and Commerce. (Feb 7, 2025)

Official Text: <https://www.congress.gov/bill/119th-congress/house-bill/1108>

Sponsor

Name: Rep. Miller-Meeks, Mariannette [R-IA-1]

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

Cosponsors (3 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Carson, André [D-IN-7]	D · IN		Feb 7, 2025
Rep. Crenshaw, Dan [R-TX-2]	R · TX		Feb 7, 2025
Rep. Schrier, Kim [D-WA-8]	D · WA		Feb 7, 2025

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred To	Feb 7, 2025

Subjects & Policy Tags

Policy Area:

Health

Related Bills

Bill	Relationship	Last Action
119 S 891	Related bill	Mar 6, 2025: Read twice and referred to the Committee on Finance.
119 HR 1768	Related bill	Mar 3, 2025: Referred to the Committee on Energy and Commerce, and in addition to the Committees on Ways and Means, the Budget, the Judiciary, and Education and Workforce, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned.

Diagnostics Testing Preparedness Plan Act of 2025

This bill requires the Department of Health and Human Services (HHS) to develop and publish a plan for the rapid development and distribution of diagnostic tests in response to a chemical, biological, radiological, or nuclear threat, including an emerging infectious disease, that causes or has significant potential to cause a declared public health emergency.

Specifically, HHS must coordinate with any relevant public and private entities, such as government entities and device manufacturers, in creating the plan and facilitating its collaborative implementation. The plan must take into consideration certain factors specified in the bill, including domestic testing capacity, novel technologies, and medical supply needs. HHS must publish the plan within one year after enactment of the bill and then update the plan every three years.

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