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Bill Fact Sheet - December 10, 2025 https://legilist.com Bill page: https://legilist.com/bill/115/hr/872

HR 872

DEVICE Act of 2017

Congress: 115 (2017–2019, Ended)

Chamber: House Policy Area: Health Introduced: Feb 6, 2017

Current Status: Referred to the Subcommittee on Health.

Latest Action: Referred to the Subcommittee on Health. (Feb 10, 2017) Official Text: https://www.congress.gov/bill/115th-congress/house-bill/872

Sponsor

Name: Rep. Lieu, Ted [D-CA-33]

Party: Democratic • State: CA • Chamber: House

Cosponsors (6 total)

Cosponsor	Party / State	Role	Date Joined
Del. Norton, Eleanor Holmes [D-DC-At Large]	D · DC		Feb 6, 2017
Rep. Chu, Judy [D-CA-27]	D · CA		Feb 6, 2017
Rep. Cummings, Elijah E. [D-MD-7]	$D\cdotMD$		Feb 6, 2017
Rep. Moore, Gwen [D-WI-4]	D · WI		Feb 6, 2017
Rep. Slaughter, Louise McIntosh [D-NY-25]	$D \cdot NY$		Feb 6, 2017
Rep. Grijalva, Raúl M. [D-AZ-3]	D · AZ		Feb 14, 2017

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Feb 10, 2017

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Disclosure; and Encouragement of Verification, Innovation, Cleaning, and Efficiency Act of 2017 or the DEVICE Act of 2017

This bill amends the Federal Food, Drug, and Cosmetic Act to require medical device manufacturers to notify the Food and Drug Administration (FDA): (1) before making changes to the design or reprocessing instructions of a device, and (2) no more than five days after widely disseminating to health care providers in a foreign country communications regarding changes to the design or reprocessing instructions of a device or regarding a safety concern about a device. A device may not be sold if the manufacturer violates these notification requirements.

Rapid assessment tests intended to ensure the proper reprocessing of reusable medical devices are defined as medical devices. The FDA must publish a list of the types of rapid assessment tests for which premarket notification must include validated instructions for use and validation data.

Actions Timeline

- Feb 10, 2017: Referred to the Subcommittee on Health.
- Feb 6, 2017: Introduced in House
- Feb 6, 2017: Referred to the House Committee on Energy and Commerce.