HR 6433

Animal Generic Drug User Fee Act of 2008

Congress: 110 (2007–2009, Ended)
Chamber: House
Policy Area: Health
Introduced: Jul 8, 2008
Current Status: Placed on the Union Calendar, Calendar No. 521.
Latest Action: Placed on the Union Calendar, Calendar No. 521. (Jul 30, 2008)
Official Text: https://www.congress.gov/bill/110th-congress/house-bill/6433

Sponsor

Name: Rep. Pallone, Frank, Jr. [D-NJ-6]
Party: Democratic • **State:** NJ • **Chamber:** House

Cosponsors (5 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Barton, Joe [R-TX-6]	R · TX		Jul 8, 2008
Rep. Deal, Nathan [R-GA-9]	R · GA		Jul 8, 2008
Rep. Dingell, John D. [D-MI-15]	D · MI		Jul 8, 2008
Rep. Towns, Edolphus [D-NY-10]	D · NY		Jul 8, 2008
Rep. DeGette, Diana [D-CO-1]	D · CO		Jul 10, 2008

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Reported by	Jul 9, 2008

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Summary (as of Jul 30, 2008)

Animal Generic Drug User Fee Act of 2008 - Amends the Federal Food, Drug, and Cosmetic Act to require the Secretary of Health and Human Services to assess and collect fees for an abbreviated application for a generic new animal drug, including application fees, product fees, and sponsor fees.

Sets forth total revenue to be collected for each type of fee for FY2009-FY2013.

Provides for fee adjustments. Requires the Secretary to establish such fees each fiscal year.

Provides for fee waivers where the Secretary finds that the generic new animal drug is intended solely to provide for a minor use or minor species indication.

Authorizes appropriations for FY2009-FY2013.

Requires the Secretary to report to Congress and make publicly available information on: (1) progress toward the goal of expediting the generic new animal drug development process and the review of abbreviated applications for such drugs; and (2) the implementation of the authority for and use of generic new animal drug fees.

Sets forth the process for developing recommendations to present to Congress for the review of abbreviated applications for generic new animal drugs after FY2013.

Terminates the authority to assess and use generic new animal drug fees on October 1, 2013. Terminates reporting requirements on January 31, 2014.

Actions Timeline

- **Jul 30, 2008:** Reported (Amended) by the Committee on Energy and Commerce. H. Rept. 110-805.
- **Jul 30, 2008:** Placed on the Union Calendar, Calendar No. 521.
- **Jul 16, 2008:** Committee Consideration and Mark-up Session Held.
- **Jul 16, 2008:** Ordered to be Reported (Amended) by Voice Vote.
- **Jul 9, 2008:** Subcommittee Consideration and Mark-up Session Held.
- **Jul 9, 2008:** Forwarded by Subcommittee to Full Committee by Voice Vote .
- **Jul 8, 2008:** Introduced in House
- **Jul 8, 2008:** Referred to the House Committee on Energy and Commerce.
- **Jul 8, 2008:** Referred to the Subcommittee on Health.

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