HR 741

To amend the Federal Food, Drug, and Cosmetic Act to prohibit the marketing of authorized generic drugs.

Congress: 112 (2011–2013, Ended)
Chamber: House
Policy Area: Health
Introduced: Feb 16, 2011
Current Status: Referred to the Subcommittee on Health.
Latest Action: Referred to the Subcommittee on Health. (Feb 28, 2011)
Official Text: https://www.congress.gov/bill/112th-congress/house-bill/741

Sponsor

Name: Rep. Emerson, Jo Ann [R-MO-8]
Party: Republican • **State:** MO • **Chamber:** House

Cosponsors

No cosponsors are listed for this bill.

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Feb 28, 2011

Subjects & Policy Tags

Policy Area:

Health

Related Bills

Bill	Relationship	Last Action
112 S 373	Related bill	**Feb 16, 2011:** Read twice and referred to the Committee on Health, Education, Labor, and Pensions.

Summary (as of Feb 16, 2011)

Amends the Federal Food, Drug, and Cosmetic Act to prohibit a holder of a new, approved drug application from commencing to manufacture, market, sell, or distribute a generic version of such drug from the time of the receipt of notice from the generic manufacturer that an abbreviated new drug application has been submitted for approval until the expiration or forfeiture of the exclusivity period granted to the generic manufacturer.

Actions Timeline

- **Feb 28, 2011:** Referred to the Subcommittee on Health.
- **Feb 16, 2011:** Introduced in House
- **Feb 16, 2011:** Referred to the House Committee on Energy and Commerce.