HR 4262

Cosmetics Safety Enhancement Act of 2012

Congress: 112 (2011–2013, Ended)
Chamber: House
Policy Area: Health
Introduced: Mar 26, 2012
Current Status: Referred to the Subcommittee on Health.
Latest Action: Referred to the Subcommittee on Health. (Mar 30, 2012)
Official Text: https://www.congress.gov/bill/112th-congress/house-bill/4262

Sponsor

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

Cosponsors (1 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Dingell, John D. [D-MI-15]	D · MI		Mar 26, 2012

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Mar 30, 2012

Subjects & Policy Tags

Policy Area:

Health

Related Bills

Bill	Relationship	Last Action
112 HR 14	Related bill	**Mar 21, 2012:** Referred to the Committee on Transportation and Infrastructure, and in addition to the Committees on Ways and Means, Natural Resources, Energy and Commerce, Agriculture, Science, Space, and Technology, the Budget, Oversight and Government Reform, Financial Services, Education and the Workforce, and Foreign Affairs, 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.

Summary (as of Mar 26, 2012)

Cosmetics Safety Enhancement Act of 2012 - Amends the the Federal Food, Drug, and Cosmetic Act to require the registration of cosmetic products and cosmetic manufacturing facilities, including, for each product: (1) a unique facility identifier, (2) every product's brand name, and (3) product ingredients. Sets registration fees.

Requires a cosmetic manufacturer: (1) before the introduction into interstate commerce of a cosmetic product, to establish a file containing scientific evidence substantiating the product's safety; (2) to report any serious adverse event information received that is associated with the use of a cosmetic product; and (3) maintain records concerning any cosmetic product that may be adulterated, misbranded, or otherwise in violation of the Act, including all records relating to cosmetic product safety substantiation or relating to serious adverse event reports.

Deems a cosmetic product adulterated if its manufacture, packing, or holding do not conform to current good manufacturing practice regulations.

Provides each manufacturer with an opportunity to cease distribution and recall of a cosmetic product if it is determined that there is a reasonable probability that the product is adulterated and the use of, or exposure to, such product will cause serious adverse health consequences or death to humans. Permits the issuance of an order requiring a recall if there is not a voluntary recall of an adulterated product.

Actions Timeline

- **Mar 30, 2012:** Referred to the Subcommittee on Health.
- **Mar 26, 2012:** Introduced in House
- **Mar 26, 2012:** Referred to the House Committee on Energy and Commerce.