HR 4747

Dietary Supplement Regulatory Implementation Act of 2004

Congress: 108 (2003–2005, Ended)
Chamber: House
Policy Area: Agriculture and Food
Introduced: Jun 25, 2004
Current Status: Referred to the Subcommittee on Health.
Latest Action: Referred to the Subcommittee on Health. (Jul 7, 2004)
Official Text: https://www.congress.gov/bill/108th-congress/house-bill/4747

Sponsor

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

Cosponsors (2 total)

Cosponsor	Party / State	Role	Date Joined
Del. Christensen, Donna M. [D-VI-At Large]	D · VI		Jul 15, 2004
Rep. Hinchey, Maurice D. [D-NY-22]	D · NY		Oct 5, 2004

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Jul 7, 2004

Subjects & Policy Tags

Policy Area:

Agriculture and Food

Related Bills

Bill	Relationship	Last Action
108 HR 4760	Related bill	**Jul 19, 2004:** Referred to the Subcommittee on Health.

Summary (as of Jun 25, 2004)

Dietary Supplement Regulatory Implementation Act of 2004 - Makes appropriations for FY 2005, and authorizes appropriations for FY 2006 through 2009: (1) to carry out the Dietary Supplement Health and Education Act of 1994 (DSHEA), the amendments made by DSHEA, and all applicable regulatory requirements for dietary supplements under the Federal Food, Drug, and Cosmetic Act; and (2) for expanded research and development of consumer information, including information on safety and beneficial effects, of dietary supplements by the Office of Dietary Supplements at the National Institutes of Health.

Directs the Secretary of Health and Human Services, acting through the Commissioner of Food and Drugs, to: (1) fully and appropriately use such funds to regulate dietary supplements; (2) report annually on DSHEA implementation and enforcement; (3) carry out programs to educate health professionals and consumers on the safety and health benefits of the dietary supplements, including the potential for interactions of dietary supplements and drugs (using specified funds authorized by this Act); and (4) establish a system for the requirements for the reporting of serious adverse experiences associated with the use of a dietary supplement received by the manufacturer, packer, or distributor whose name appears on the label of the product.

Expresses the sense of the Congress regarding dietary supplements containing ephedrine alkaloids.

Actions Timeline

- **Jul 7, 2004:** Referred to the Subcommittee on Health.
- **Jun 25, 2004:** Introduced in House
- **Jun 25, 2004:** Referred to the House Committee on Energy and Commerce.

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