

HR 3870

Prescription Drug Abuse Elimination Act of 2004

Congress: 108 (2003–2005, Ended)

Chamber: House Policy Area: Health Introduced: Mar 2, 2004

Current Status: Referred to the Subcommittee on Health.

Latest Action: Referred to the Subcommittee on Health. (Mar 11, 2004) **Official Text:** https://www.congress.gov/bill/108th-congress/house-bill/3870

Sponsor

Name: Rep. Norwood, Charles W. [R-GA-9]

Party: Republican • State: GA • Chamber: House

Cosponsors (7 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Strickland, Ted [D-OH-6]	D · OH		Mar 2, 2004
Rep. Boehner, John A. [R-OH-8]	$R \cdot OH$		Apr 1, 2004
Rep. Isakson, Johnny [R-GA-6]	$R \cdot GA$		Apr 1, 2004
Rep. Shays, Christopher [R-CT-4]	$R \cdot CT$		Apr 1, 2004
Rep. Weldon, Curt [R-PA-7]	$R \cdot PA$		Apr 1, 2004
Rep. Weldon, Dave [R-FL-15]	$R \cdot FL$		Apr 1, 2004
Rep. Doolittle, John T. [R-CA-4]	$R \cdot CA$		Sep 14, 2004

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Mar 11, 2004

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Prescription Drug Abuse Elimination Act of 2004 - Amends the Public Health Service Act to require the Secretary of Health and Human Services to provide grants to States to develop or enhance prescription drug monitoring programs, subject to specified requirements. Requires a dispenser to report to the State database when a patient presents a prescription for a Schedule II, III, or IV controlled substance.

Amends the Federal Food, Drug, and Cosmetic Act to require Internet pharmacies to meet specified criteria, including obtaining verification of a valid prescription in some circumstances. Allows district courts to order Internet providers to remove access to sites that violate this Act.

Directs the Secretary to promulgate chain-of-custody requirements to track prescription drugs from the manufacturer through each importer, distributor, and retailer. Deems a drug misbranded if such requirements are not met.

Amends the Controlled Substances Import and Export Act to restrict the importation of a schedule II, III, or IV drug by a U.S. resident without a valid prescription to once during any 30-day period.

Requires the Secretary of Health and Human Services, acting through the Commissioner of Food and Drugs, to convene a working group to study and report on pharmaceutical counterfeiting.

Requires the Secretary to conduct research and report on issues related to prescription drug abuse.

Requires the Administrator of the Substance Abuse and Mental Health Services Administration to maintain a comprehensive, national database on deaths occurring as a result of substance abuse.

Actions Timeline

- Mar 11, 2004: Referred to the Subcommittee on Health.
- Mar 2, 2004: Introduced in House
- Mar 2, 2004: Introduced in House
- Mar 2, 2004: Referred to the House Committee on Energy and Commerce.