

HR 5839

Safeguarding America's Pharmaceuticals Act of 2008

Congress: 110 (2007–2009, Ended)

Chamber: House
Policy Area: Health
Introduced: Apr 17, 2008

Current Status: Referred to the Subcommittee on Health.

Latest Action: Referred to the Subcommittee on Health. (Apr 17, 2008) **Official Text:** https://www.congress.gov/bill/110th-congress/house-bill/5839

Sponsor

Name: Rep. Buyer, Steve [R-IN-4]

Party: Republican • State: IN • Chamber: House

Cosponsors (13 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Green, Gene [D-TX-29]	$D\cdotTX$		Apr 17, 2008
Rep. Matheson, Jim [D-UT-2]	$D \cdot UT$		Apr 17, 2008
Rep. Rogers, Mike J. [R-MI-8]	$R \cdot MI$		Apr 17, 2008
Rep. Bilbray, Brian P. [R-CA-50]	$R \cdot CA$		Apr 24, 2008
Rep. Ferguson, Mike [R-NJ-7]	$R \cdot NJ$		Apr 24, 2008
Rep. Bartlett, Roscoe G. [R-MD-6]	$R \cdot MD$		May 15, 2008
Rep. Butterfield, G. K. [D-NC-1]	$D \cdot NC$		May 15, 2008
Rep. Hunter, Duncan [R-CA-52]	$R \cdot CA$		May 15, 2008
Rep. Jackson-Lee, Sheila [D-TX-18]	$D\cdotTX$		May 15, 2008
Rep. Wilson, Joe [R-SC-2]	$R \cdot SC$		May 15, 2008
Rep. Jefferson, William J. [D-LA-2]	D·LA		Jun 4, 2008
Rep. Meek, Kendrick B. [D-FL-17]	D·FL		Jun 4, 2008
Rep. Souder, Mark E. [R-IN-3]	$R \cdot IN$		Sep 24, 2008

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred to	Apr 17, 2008

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Safeguarding America's Pharmaceuticals Act of 2008 - Amends the Federal Food, Drug, and Cosmetic Act to require the Secretary of the Treasury to destroy imported counterfeit drugs and certain adulterated or misbranded imported drugs.

Requires the manufacturer of a prescription drug to provide to each wholesale distributor or dispenser a packing list or comparable document that includes information identifying the drug's proprietary and established names and National Drug Code number.

Requires each person engaged in wholesale distribution of a prescription drug to provide information on each prior transaction involving the drug.

Requires the Secretary of Health and Human Services (the Secretary) to report to Congress on the feasibility and operational efficiencies of adopting security technologies, including track and trace technology throughout the prescription drug supply chain. Phases in requirements for standardized numerical identifiers that are unique to each unit of a prescription drug.

Requires the Secretary to issue regulations to establish an effective drug identification and tracking system to authenticate the wholesale distribution history of any prescription drug that is subject to a requirement for such an identifier.

Requires the Comptroller General to study the availability and cost of technologies to dispensers to comply with this Act.

Requires the Secretary to: (1) award matching grants for drug identification and tracking systems to ensure the security and integrity of the drug supply chain; and (2) issue guidelines for wholesale distributors that prescribe requirements that include mandatory background checks and physical inspection of facilities.

Sets forth civil penalties for violations of this Act.

Requires the Secretary to study threats to the domestic prescription drug supply chain and make recommendations for improvements.

Actions Timeline

- Apr 17, 2008: Introduced in House
- Apr 17, 2008: Referred to the House Committee on Energy and Commerce.
- Apr 17, 2008: Referred to the Subcommittee on Health.