

## HR 4056

Science and Technology Regulatory Relief Act of 2012

**Congress:** 112 (2011–2013, Ended)

**Chamber:** House

**Policy Area:** Health

**Introduced:** Feb 16, 2012

**Current Status:** Referred to the House Committee on Energy and Commerce.

**Latest Action:** Referred to the House Committee on Energy and Commerce. (Feb 16, 2012)

**Official Text:** <https://www.congress.gov/bill/112th-congress/house-bill/4056>

### Sponsor

**Name:** Rep. Bilbray, Brian P. [R-CA-50]

**Party:** Republican • **State:** CA • **Chamber:** House

### Cosponsors (6 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Bono Mack, Mary [R-CA-45]	R · CA		Feb 16, 2012
Rep. Calvert, Ken [R-CA-44]	R · CA		Feb 16, 2012
Rep. Davis, Susan A. [D-CA-53]	D · CA		Feb 16, 2012
Rep. Hunter, Duncan D. [R-CA-52]	R · CA		Feb 16, 2012
Rep. Lewis, Jerry [R-CA-41]	R · CA		Feb 16, 2012
Rep. Royce, Edward R. [R-CA-40]	R · CA		Feb 16, 2012

### Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Referred To	Feb 16, 2012

### Subjects & Policy Tags

#### Policy Area:

Health

### Related Bills

*No related bills are listed.*

## Summary (as of Feb 16, 2012)

---

Science and Technology Regulatory Relief Act of 2012 - Amends the Federal Food, Drug, and Cosmetic Act to prohibit a state or political subdivision from conducting or requiring an inspection of a factory, warehouse, or establishment in which a drug or device is manufactured, processed, packed, or held by a manufacturer or wholesale distributor for introduction into interstate commerce, or after such introduction, for purposes of verifying compliance with such Act, Public Health Service Act requirements regarding the regulation of biological products, or any similar requirements established pursuant to state law.

Makes exceptions for such an inspection if: (1) the state or subdivision makes a determination that a drug or device presents a threat of serious adverse health consequences or death; (2) the Secretary of Health and Human Services (HHS) orders a recall of a drug, biological product, or device manufactured, processed, packed, or held at the factory, warehouse, or establishment; or (3) the Secretary requests or authorizes the state to conduct or require the inspection.

## Actions Timeline

---

- **Feb 16, 2012:** Introduced in House
- **Feb 16, 2012:** Referred to the House Committee on Energy and Commerce.

# LegiList

CONGRESS, MADE CLEAR.

## Search Every Federal Bill, Law, and Vote

LegiList is the fastest way to research Congress. Track any bill from introduction to enactment, see how every legislator voted, follow committee activity, and read the full text of every bill — all in one place, always up to date.

[legilist.com](https://legilist.com)

## Free Course: Learn How Congress Actually Works

LegiList Learn is a free, self-paced course that walks through the entire legislative process — from drafting a bill to a presidential signature. Seven modules, plain language, no politics. Earn a certificate when you finish.

[legilist.com/learn](https://legilist.com/learn)

## Developer API: Build Apps on Legislative Data

The LegiList API gives developers direct access to bills, votes, legislators, committees, and more. Start free with 1,000 requests per day — no credit card required. Upgrade to Pro when you need to scale.

[legilist.com/api](https://legilist.com/api)

Public data belongs to the public. — [legilist.com](https://legilist.com)