# HR 3203

Novel Device Regulatory Relief Act of 2011

**Congress:** 112 (2011–2013, Ended)
**Chamber:** House
**Policy Area:** Health
**Introduced:** Oct 14, 2011
**Current Status:** Referred to the Subcommittee on Health.
**Latest Action:** Referred to the Subcommittee on Health. (Oct 18, 2011)
**Official Text:** https://www.congress.gov/bill/112th-congress/house-bill/3203

## Sponsor

**Name:** Rep. Bilbray, Brian P. [R-CA-50]
**Party:** Republican • **State:** CA • **Chamber:** House

## Cosponsors (18 total)

| Cosponsor | Party / State | Role | Date Joined |
|---|---|---|---|
| Rep. Blackburn, Marsha [R-TN-7] | R · TN | | Oct 14, 2011 |
| Rep. Burgess, Michael C. [R-TX-26] | R · TX | | Oct 14, 2011 |
| Rep. Capps, Lois [D-CA-23] | D · CA | | Oct 14, 2011 |
| Rep. Dent, Charles W. [R-PA-15] | R · PA | | Oct 14, 2011 |
| Rep. Guthrie, Brett [R-KY-2] | R · KY | | Oct 14, 2011 |
| Rep. Hunter, Duncan D. [R-CA-52] | R · CA | | Oct 14, 2011 |
| Rep. Lance, Leonard [R-NJ-7] | R · NJ | | Oct 14, 2011 |
| Rep. Latta, Robert E. [R-OH-5] | R · OH | | Oct 14, 2011 |
| Rep. Paulsen, Erik [R-MN-3] | R · MN | | Oct 14, 2011 |
| Rep. Shimkus, John [R-IL-19] | R · IL | | Oct 14, 2011 |
| Rep. Stearns, Cliff [R-FL-6] | R · FL | | Oct 14, 2011 |
| Rep. Culberson, John Abney [R-TX-7] | R · TX | | Oct 24, 2011 |
| Rep. Calvert, Ken [R-CA-44] | R · CA | | Nov 1, 2011 |
| Rep. Walden, Greg [R-OR-2] | R · OR | | Nov 1, 2011 |
| Rep. Mulvaney, Mick [R-SC-5] | R · SC | | Jan 18, 2012 |
| Rep. Rokita, Todd [R-IN-4] | R · IN | | Jan 23, 2012 |
| Rep. Matheson, Jim [D-UT-2] | D · UT | | Feb 1, 2012 |
| Rep. Gosar, Paul A. [R-AZ-1] | R · AZ | | Apr 24, 2012 |

## Committee Activity

| Committee | Chamber | Activity | Date |
|---|---|---|---|
| Energy and Commerce Committee | House | Referred to | Oct 18, 2011 |

## Subjects & Policy Tags

**Policy Area:**

Health

## Related Bills

*No related bills are listed.*

## Summary (as of Oct 14, 2011)

Novel Device Regulatory Relief Act of 2011 - Amends the Federal Food, Drug, and Cosmetic Act to revise the process for requesting classification for a type of medical device that has not been previously classified by removing the requirement that a person can only file a request for classification of a new medical device after the Secretary of Health and Human Services (HHS) classifies the device as a class III device in response to the person filing a notice of intent to market the device.

## Actions Timeline

- **Oct 18, 2011:** Referred to the Subcommittee on Health.
- **Oct 14, 2011:** Introduced in House
- **Oct 14, 2011:** Referred to the House Committee on Energy and Commerce.