

## HR 1499

### Protecting Consumer Access to Generic Drugs Act of 2019

**Congress:** 116 (2019–2021, Ended)

**Chamber:** House

**Policy Area:** Health

**Introduced:** Mar 5, 2019

**Current Status:** Placed on the Union Calendar, Calendar No. 30.

**Latest Action:** Placed on the Union Calendar, Calendar No. 30. (May 10, 2019)

**Official Text:** <https://www.congress.gov/bill/116th-congress/house-bill/1499>

## Sponsor

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**Name:** Rep. Rush, Bobby L. [D-IL-1]

**Party:** Democratic • **State:** IL • **Chamber:** House

**Cosponsors** (38 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Craig, Angie [D-MN-2]	D · MN		Mar 26, 2019
Rep. Dingell, Debbie [D-MI-12]	D · MI		Mar 26, 2019
Rep. Eshoo, Anna G. [D-CA-18]	D · CA		Mar 26, 2019
Rep. Kennedy, Joseph P., III [D-MA-4]	D · MA		Mar 26, 2019
Rep. Pallone, Frank, Jr. [D-NJ-6]	D · NJ		Mar 26, 2019
Rep. Ruiz, Raul [D-CA-36]	D · CA		Mar 26, 2019
Rep. Matsui, Doris O. [D-CA-6]	D · CA		Mar 27, 2019
Rep. Clarke, Yvette D. [D-NY-9]	D · NY		Apr 1, 2019
Rep. Van Drew, Jefferson [D-NJ-2]	D · NJ		Apr 1, 2019
Rep. Khanna, Ro [D-CA-17]	D · CA		Apr 3, 2019
Rep. Pappas, Chris [D-NH-1]	D · NH		Apr 4, 2019
Rep. Schakowsky, Janice D. [D-IL-9]	D · IL		Apr 4, 2019
Rep. Cohen, Steve [D-TN-9]	D · TN		Apr 8, 2019
Rep. Welch, Peter [D-VT-At Large]	D · VT		Apr 8, 2019
Rep. Hastings, Alcee L. [D-FL-20]	D · FL		Apr 9, 2019
Rep. Larson, John B. [D-CT-1]	D · CT		Apr 9, 2019
Rep. Case, Ed [D-HI-1]	D · HI		Apr 15, 2019
Rep. Finkenauer, Abby [D-IA-1]	D · IA		Apr 15, 2019
Rep. Neguse, Joe [D-CO-2]	D · CO		Apr 15, 2019
Rep. Cooper, Jim [D-TN-5]	D · TN		Apr 25, 2019
Rep. Langevin, James R. [D-RI-2]	D · RI		Apr 25, 2019
Rep. Quigley, Mike [D-IL-5]	D · IL		Apr 25, 2019
Rep. Wild, Susan [D-PA-7]	D · PA		Apr 25, 2019
Rep. Mucarsel-Powell, Debbie [D-FL-26]	D · FL		Apr 29, 2019
Rep. Raskin, Jamie [D-MD-8]	D · MD		Apr 29, 2019
Rep. Scott, David [D-GA-13]	D · GA		Apr 29, 2019
Rep. Cisneros, Gilbert Ray, Jr. [D-CA-39]	D · CA		May 1, 2019
Rep. Stevens, Haley M. [D-MI-11]	D · MI		May 1, 2019
Rep. Thompson, Bennie G. [D-MS-2]	D · MS		May 1, 2019
Rep. Carbajal, Salud O. [D-CA-24]	D · CA		May 7, 2019
Rep. Carson, Andre [D-IN-7]	D · IN		May 7, 2019
Rep. Casten, Sean [D-IL-6]	D · IL		May 7, 2019
Rep. Davids, Sharice [D-KS-3]	D · KS		May 7, 2019
Rep. Houlahan, Chrissy [D-PA-6]	D · PA		May 7, 2019
Rep. Kelly, Robin L. [D-IL-2]	D · IL		May 7, 2019
Rep. Connolly, Gerald E. [D-VA-11]	D · VA		May 9, 2019
Rep. Smith, Adam [D-WA-9]	D · WA		May 9, 2019
Rep. Underwood, Lauren [D-IL-14]	D · IL		May 9, 2019

## Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Reported by	Mar 27, 2019
Judiciary Committee	House	Referred to	Apr 8, 2019

## Subjects & Policy Tags

### Policy Area:

Health

## Related Bills

Bill	Relationship	Last Action
116 HR 19	Related bill	Dec 19, 2019: Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law.
116 HR 2700	Related bill	Jun 26, 2019: Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law.
116 HR 987	Related bill	May 20, 2019: Received in the Senate and Read twice and referred to the Committee on Health, Education, Labor, and Pensions.

## Summary (as of May 10, 2019)

### Protecting Consumer Access to Generic Drugs Act of 2019

This bill prohibits the manufacturer of a brand-name, generic, or biosimilar drug from entering into certain agreements to resolve or settle a patent infringement claim in connection with the sale of a drug or biological product.

Specifically, such an agreement shall, with some exceptions, be a violation of the bill if the filer of a subsequent application to market a drug or biological product receives anything of value and agrees to limit or forego research, development, manufacturing, marketing, or sales of the subsequent drug or biological product. (Typically, a subsequent application seeks to market a generic or biosimilar version of a patented drug or biological product.)

Penalties for violations of the bill include civil penalties and loss of the 180-day exclusivity period for a generic drug. The Federal Trade Commission (FTC) shall have exclusive authority to litigate to enforce the bill.

When a generic or biosimilar drug manufacturer enters into an agreement with another drug manufacturer related to the manufacturing, marketing or sale of a drug, the manufacturers must certify that the material they have given the FTC concerning the agreement contains (1) the complete agreement; and (2) any agreements related to the main agreement, including descriptions of any oral agreements or representations.

## **Actions Timeline**

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- **May 10, 2019:** Reported (Amended) by the Committee on Energy and Commerce. H. Rept. 116-52, Part I.
- **May 10, 2019:** Committee on the Judiciary discharged.
- **May 10, 2019:** Placed on the Union Calendar, Calendar No. 30.
- **Apr 8, 2019:** Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law.
- **Apr 3, 2019:** Committee Consideration and Mark-up Session Held.
- **Apr 3, 2019:** Ordered to be Reported (Amended) by Voice Vote.
- **Mar 27, 2019:** Subcommittee Consideration and Mark-up Session Held.
- **Mar 27, 2019:** Forwarded by Subcommittee to Full Committee (Amended) by Voice Vote .
- **Mar 6, 2019:** Referred to the Subcommittee on Health.
- **Mar 5, 2019:** Introduced in House
- **Mar 5, 2019:** Referred to the Committee on Energy and Commerce, and in addition to the Committee on the Judiciary, for a period to be subsequently determined by the Speaker, in each case for consideration of such provisions as fall within the jurisdiction of the committee concerned.