

## HR 4398

Affordable Prescriptions for Patients Through Promoting Competition Act of 2019

**Congress:** 116 (2019–2021, Ended)

**Chamber:** House

**Policy Area:** Commerce

**Introduced:** Sep 19, 2019

**Current Status:** Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law.

**Latest Action:** Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law. (Oct 2, 2019)

**Official Text:** <https://www.congress.gov/bill/116th-congress/house-bill/4398>

### Sponsor

**Name:** Rep. Cicilline, David N. [D-RI-1]

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

### Cosponsors

*No cosponsors are listed for this bill.*

### Committee Activity

Committee	Chamber	Activity	Date
Judiciary Committee	House	Referred to	Oct 2, 2019

### Subjects & Policy Tags

**Policy Area:**

Commerce

### Related Bills

Bill	Relationship	Last Action
116 HR 5133	Related bill	<b>Dec 24, 2020:</b> Placed on the Union Calendar, Calendar No. 578.
116 HR 2486	Related bill	<b>Jul 23, 2020:</b> Message on House action received in Senate and at desk: House amendments to Senate amendment.
116 S 1416	Related bill	<b>Jun 28, 2019:</b> Placed on Senate Legislative Calendar under General Orders. Calendar No. 132.

## Affordable Prescriptions for Patients Through Promoting Competition Act of 2019

This bill prohibits product hopping by drug manufacturers. Product hopping is presumed when a drug manufacturer engages in a *hard switch* or a *soft switch*. A hard switch occurs when, after a manufacturer receives notice of an application for a generic drug, either, (1) the manufacturer obtains removal of a drug from the Food and Drug Administration's approved drug list or the drug is moved to the discontinued products list, and the manufacturer markets or sells a follow-on product; or (2) a manufacturer announces the withdrawal or discontinuance of a listed drug, or the manufacturer destroys the inventory of a listed drug in a manner that impedes generic drug competitors, and the manufacturer markets or sells a follow-on product. A *follow-on product* is a changed, modified, or reformulated version of a manufacturer's already-approved drug or biological product that still treats the same medical condition.

A soft switch occurs when a manufacturer receives notice of an application for a generic drug, takes other actions that impede generic drug competitors, and the manufacturer markets or sells a follow-on product.

A drug manufacturer may rebut a presumption of product hopping by demonstrating that its conduct was not intended to limit competition.

### Actions Timeline

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- **Oct 2, 2019:** Referred to the Subcommittee on Antitrust, Commercial, and Administrative Law.
- **Sep 19, 2019:** Introduced in House
- **Sep 19, 2019:** Referred to the House Committee on the Judiciary.