

## S 1115

### Making Pharmaceutical Markets More Competitive Act

**Congress:** 115 (2017–2019, Ended)

**Chamber:** Senate

**Policy Area:** Health

**Introduced:** May 11, 2017

**Current Status:** Read twice and referred to the Committee on Health, Education, Labor, and Pensions.

**Latest Action:** Read twice and referred to the Committee on Health, Education, Labor, and Pensions. (May 11, 2017)

**Official Text:** <https://www.congress.gov/bill/115th-congress/senate-bill/1115>

### Sponsor

**Name:** Sen. Collins, Susan M. [R-ME]

**Party:** Republican • **State:** ME • **Chamber:** Senate

### Cosponsors (3 total)

| Cosponsor                     | Party / State | Role | Date Joined  |
|-------------------------------|---------------|------|--------------|
| Sen. Cotton, Tom [R-AR]       | R · AR        |      | May 11, 2017 |
| Sen. Franken, Al [D-MN]       | D · MN        |      | May 11, 2017 |
| Sen. McCaskill, Claire [D-MO] | D · MO        |      | May 11, 2017 |

### Committee Activity

| Committee  | Chamber | Activity    | Date         |
|--|---------|-------------|--------------|
| Health, Education, Labor, and Pensions Committee | Senate  | Referred To | May 11, 2017 |

### Subjects & Policy Tags

#### Policy Area:

Health

### Related Bills

| Bill        | Relationship   | Last Action   |
|-------------|----------------|---|
| 115 HR 2562 | Identical bill | <b>May 26, 2017:</b> Referred to the Subcommittee on Health.                                      |
| 115 S 934   | Related bill   | <b>May 11, 2017:</b> Placed on Senate Legislative Calendar under General Orders. Calendar No. 76. |

## **Making Pharmaceutical Markets More Competitive Act**

This bill amends the Federal Food, Drug, and Cosmetic Act to require the Food and Drug Administration (FDA) to prioritize the review of generic drug applications and supplements with respect to drugs that are in a shortage or for which there are not more than three approved drugs.

The holder of an approved drug application must notify the FDA within 180 days of withdrawing or transferring the application or withdrawing the drug from sale.

The FDA must maintain a list of generic drugs with three or fewer holders of approved applications.

## **Actions Timeline**

---

- **May 11, 2017:** Introduced in Senate
- **May 11, 2017:** Read twice and referred to the Committee on Health, Education, Labor, and Pensions.