

S 2100

Personal Care Products Safety Act

Congress: 117 (2021–2023, Ended)

Chamber: Senate

Policy Area: Health

Introduced: Jun 17, 2021

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

Latest Action: Read twice and referred to the Committee on Health, Education, Labor, and Pensions. (Sponsor introductory remarks on measure: CR S4625) (Jun 17, 2021)

Official Text: <https://www.congress.gov/bill/117th-congress/senate-bill/2100>

Sponsor

Name: Sen. Feinstein, Dianne [D-CA]

Party: Democratic • **State:** CA • **Chamber:** Senate

Cosponsors (3 total)

| Cosponsor | Party / State | Role | Date Joined |
|---------------------------------|---------------|------|--------------|
| Sen. Collins, Susan M. [R-ME] | R · ME | | Jun 17, 2021 |
| Sen. Blumenthal, Richard [D-CT] | D · CT | | Oct 19, 2021 |
| Sen. Whitehouse, Sheldon [D-RI] | D · RI | | Feb 15, 2022 |

Committee Activity

| Committee | Chamber | Activity | Date |
|--|---------|-------------|--------------|
| Health, Education, Labor, and Pensions Committee | Senate | Referred To | Jun 17, 2021 |

Subjects & Policy Tags

Policy Area:

Health

Related Bills

No related bills are listed.

Personal Care Products Safety Act

This bill requires cosmetics brands and manufacturers to register with the Food and Drug Administration (FDA), provides the FDA with various regulatory authorities, and addresses related issues.

Cosmetics brand owners and contract manufacturers must annually register their manufacturing facilities with the FDA. Certain entities and facilities, such as manufacturers with gross sales below certain thresholds, are exempted. Registrants with gross annual sales above certain thresholds must pay a registration fee.

Each registration must contain certain information, including an ingredient list for all cosmetic products from a registered facility, with different requirements for registrants that qualify as small businesses.

The FDA may suspend a registration for various reasons, including if the registrant's product has a reasonable probability of causing serious adverse health consequences and the problem cannot be isolated to a single product.

The FDA must annually conduct a safety review of at least five cosmetics ingredients or nonfunctional constituents and, if appropriate, issue a final finding on the safety of that ingredient or constituent. The FDA must also implement regulations for good cosmetics manufacturing practices.

The bill also requires cosmetics brand owners and manufacturers to report to the FDA any serious adverse event associated with their products.

The FDA shall have various authorities to regulate cosmetics, including to (1) order a mandatory recall of a product, (2) inspect the records of manufacturers and processors, and (3) require warning labels for certain products.

The FDA must issue a proposed rule to ban using intentionally added perfluoroalkyl or polyfluoroalkyl substances in cosmetics.

Actions Timeline

- **Jun 17, 2021:** Introduced in Senate
- **Jun 17, 2021:** Read twice and referred to the Committee on Health, Education, Labor, and Pensions. (Sponsor introductory remarks on measure: CR S4625)

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