

Bill Fact Sheet - December 5, 2025 https://legilist.com

Bill page: https://legilist.com/bill/110/s/1479

S 1479

Safe Tissue Act

Congress: 110 (2007–2009, Ended)

Chamber: Senate Policy Area: Health Introduced: May 24, 2007

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 24, 2007)

Official Text: https://www.congress.gov/bill/110th-congress/senate-bill/1479

Sponsor

Name: Sen. Schumer, Charles E. [D-NY]

Party: Democratic • State: NY • Chamber: Senate

Cosponsors (1 total)

| Cosponsor | Party / State | Role | Date Joined |
|-------------------------------|---------------|------|--------------|
| Sen. Leahy, Patrick J. [D-VT] | $D \cdot VT$ | | May 24, 2007 |

Committee Activity

| Committee | Chamber | Activity | Date |
|--|---------|-------------|--------------|
| Health, Education, Labor, and Pensions Committee | Senate | Referred To | May 24, 2007 |

Subjects & Policy Tags

Policy Area:

Health

Related Bills

| Bill | Relationship | Last Action |
|-------------|----------------|---|
| 110 HR 2731 | Identical bill | Jun 14, 2007: Referred to the Subcommittee on Health. |

Safe Tissue Act - Requires the Food and Drug Administration (FDA) to inspect, at least once every two years, each establishment that engages in the manufacture of human cells, tissues, and cellular and tissue-based products. Allows the Secretary of Health and Human Services to establish a user fee to fund such inspections.

Requires the FDA to conduct periodic audits of all documentation submitted by each such establishment to determine compliance with all applicable requirements, including requirements related to ensuring: (1) that human cells, tissues, or cellular or tissue-based products are obtained legally; (2) that donor eligibility and donor medical history interviews are based on accurate information that was not provided or obtained in a fraudulent manner; and (3) current good tissue practice.

Requires the Secretary to publish a model form containing minimum requirements for establishments to use in obtaining consent from a potential donor of human cells, tissues, or cellular or tissue-based products. Sets forth penalties for failing to comply with model form requirements or for knowingly using fraudulent information.

Directs the Secretary to: (1) accredit establishments and the personnel of such establishments who participate in the recovery, processing, storage, labeling, packaging, or distribution of human cells, tissues, or cellular or tissue-based products; and (2) define "reasonable payments" that are associated with donation of human tissue and tissue-based products for purposes of the National Organ Transplant Act.

Actions Timeline

- May 24, 2007: Introduced in Senate
- May 24, 2007: Read twice and referred to the Committee on Health, Education, Labor, and Pensions.