HR 1082

Shandra Eisenga Human Cell and Tissue Product Safety Act

Congress: 119 (2025–2027, Current)
Chamber: House
Policy Area: Health
Introduced: Feb 6, 2025
Current Status: Received in the Senate and Read twice and referred to the Committee on Health, Education, Labor, and
Latest Action: Received in the Senate and Read twice and referred to the Committee on Health, Education, Labor, and Pensions. (Jun 24, 2025)
Official Text: https://www.congress.gov/bill/119th-congress/house-bill/1082

Sponsor

Name: Rep. Moolenaar, John R. [R-MI-2]
Party: Republican • **State:** MI • **Chamber:** House

Cosponsors (1 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Dingell, Debbie [D-MI-6]	D · MI		Feb 6, 2025

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Reported By	Jun 12, 2025
Health, Education, Labor, and Pensions Committee	Senate	Referred To	Jun 24, 2025

Subjects & Policy Tags

Policy Area:

Health

Related Bills

Bill	Relationship	Last Action
119 HR 340	Related bill	**Jan 13, 2025:** Referred to the House Committee on Energy and Commerce.

Summary (as of Feb 6, 2025)

Shandra Eisenga Human Cell and Tissue Product Safety Act

This bill establishes civil penalties for violations of regulations governing the donation and handling of human cell and tissue products. It also requires the Food and Drug Administration (FDA) to report on the regulation of these products and to provide related information to stakeholders. (*Human cell and tissue products* are articles containing or consisting of human cells or tissues that are intended for use in a human recipient.)

Specifically, the bill establishes civil penalties for violations of the FDA's regulations on donor eligibility and current good tissue practice for manufacturing and distributing human cell and tissue products.

Also, the bill requires the FDA to conduct workshops to educate stakeholders and facilitate discussion on the science and regulation of human cell and tissue products. The FDA must establish a public docket to receive written comments on this topic, and report to Congress with recommendations. The bill also requires the FDA to support the development of educational materials for health care professionals regarding organ, tissue, and eye donations and related topics.

Additionally, the bill requires the FDA to publish on its website educational materials about the Tissue Reference Group (a working group within the FDA) and best practices for obtaining a recommendation from them about human cell and tissue products. Also, annually for three years, the FDA must publish on its website information on inquiries submitted to the Tissue Reference Group and FDA registrations and inspections regarding human cell and tissue manufacturers.

Actions Timeline

- **Jun 24, 2025:** Received in the Senate and Read twice and referred to the Committee on Health, Education, Labor, and Pensions.
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