

# HOUSE RESEARCH

## Bill Summary

**FILE NUMBER:** H.F. 236  
**Version:** As introduced

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**Subject:** Right to Try Act

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### Overview

This bill creates the Right to Try Act. The act allows certain eligible patients with a terminal disease to use investigation drugs, biological products, or devices that are not approved by the Food and Drug Administration but have completed phase 1 of a clinical trial. The act specifies eligibility requirements, and, among other things, states that health insurance, private or public, is not required to cover the cost of these products.

#### Section

**1** **Investigational drug use.** Adds § 151.375.

**Subd. 1. Title; citation.** States this section may be cited as the “Right to Try Act.”

**Subd. 2. Definitions.** Defines terms.

**Subd. 3. Eligibility.** Provides requirements a patient must meet in order to access an investigational drug, biological product, or device under this section. Requires a physician to document in writing that a patient, among other things, has a terminal disease and has given written informed consent.

**Subd. 4. Availability.** Allows, but specifically does not require, a manufacturer of an investigational drug, biological product, or device to make those products available to eligible patients.

**Subd. 5. Costs.** Allows a manufacturer to provide an investigation drug, biological product or device without receiving compensation and allows a manufacturer to require a patient to pay associated costs.

**Section**

**Subd. 6. Insurance coverage.** States that this section is not requiring private health insurance or a state health care program to cover costs of an investigational drug, biological product, or device.

**Subd. 7. Professional licensing.** Prohibits a health-related licensing board from taking disciplinary action against a licensee solely based on the licensee providing a prescription or recommendation under this section.

**Subd. 8. Penalty.** Provides that any official, employee, or agent of the state of Minnesota shall be guilty of a misdemeanor for attempting to block or blocking access of an eligible patient to an investigational drug, biological product, or device.

**Subd. 9. Severability.** States that if any section or its application is held to be invalid, it shall not affect any other provision of the section.