**South Carolina General Assembly**

123rd Session, 2019-2020

**S. 225**

**STATUS INFORMATION**

General Bill

Sponsors: Senators Gambrell and Nicholson

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Companion/Similar bill(s): 3036

Introduced in the Senate on January 8, 2019

Currently residing in the Senate Committee on **Medical Affairs**

Summary: Neonatal testing

**HISTORY OF LEGISLATIVE ACTIONS**

Date Body Action Description with journal page number

12/12/2018 Senate Prefiled

12/12/2018 Senate Referred to Committee on **Medical Affairs**

1/8/2019 Senate Introduced and read first time ([Senate Journal‑page 141](file:///h:\sj\20190108.docx))

1/8/2019 Senate Referred to Committee on **Medical Affairs** ([Senate Journal‑page 141](file:///h:\sj\20190108.docx))

View the latest [legislative information](http://www.scstatehouse.gov/billsearch.php?billnumbers=225&session=123&summary=B) at the website

**VERSIONS OF THIS BILL**

[12/12/2018](file:///p:\pprever\2019-20\225_20181212.docx)

**A** **BILL**

TO AMEND THE CODE OF LAWS OF SOUTH CAROLINA, 1976, BY ADDING SECTION 44‑37‑35 SO AS TO REQUIRE NEONATAL TESTING FOR CERTAIN GENETIC DISORDERS AND DISEASES AND FOR OTHER PURPOSES.

Be it enacted by the General Assembly of the State of South Carolina:

SECTION 1. Chapter 37, Title 44 of the 1976 Code is amended by adding:

“Section 44‑37‑35. (A) Neonatal testing conducted pursuant to Section 44‑37‑30 must include testing for the following:

(1) Krabbe disease;

(2) Fabry disease;

(3) Gaucher disease;

(4) Pompe disease;

(5) Hurler syndrome; and

(6) Niemann‑Pick disease.

(B) The department shall require additional lysosomal storage disorders to be tested in accordance with Section 44‑37‑30 pursuant to a duly promulgated regulation as testing for such disorders becomes available.”

SECTION 2. This act takes effect upon approval by the Governor.

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