SF3125 REVISOR EM S3125-1 1st Engrossment

SENATE STATE OF MINNESOTA NINETY-FIRST SESSION

S.F. No. 3125

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DATE 02/13/2020 D-PG OFFICIAL STATUS 4754 Introduction and first reading Referred to Health and Human Services Finance and Policy 03/02/2020 5130 Authors added Abeler; Hoffman 03/04/2020 5241a Comm report: To pass as amended 5246 5629 Second reading 04/16/2020 Special Order Third reading Passed 5630 See SF13, Art. 13, Sec. 28

1.1 A bill for an act

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relating to medical assistance; providing coverage for routine patient costs that are incurred in the course of a clinical trial if the medical assistance program would provide coverage for the same routine patient costs not incurred in a clinical trial; amending Minnesota Statutes 2018, section 256B.0625, subdivision 64.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:

Section 1. Minnesota Statutes 2018, section 256B.0625, subdivision 64, is amended to read:

Subd. 64. **Investigational drugs, biological products, and devices.** (a) Medical assistance and the early periodic screening, diagnosis, and treatment (EPSDT) program do not cover the costs of any services that are incidental to, associated with, or resulting from the use of investigational drugs, biological products, or devices as defined in section 151.375 or any other treatment that is part of an approved clinical trial as defined in section 62Q.526. Participation of an enrollee in an approved clinical trial does not preclude coverage of medically necessary services covered under this chapter that are not related to the approved clinical trial.

- (b) Notwithstanding paragraph (a), stiripentol may be covered by the EPSDT program if all the following conditions are met:
- 1.19 (1) the use of stiripentol is determined to be medically necessary;
- (2) the enrollee has a documented diagnosis of Dravet syndrome, regardless of whether
 an SCN1A genetic mutation is found, or the enrollee is a child with malignant migrating
 partial epilepsy in infancy due to an SCN2A genetic mutation;

Section 1.

(3) all other available covered prescription medications that are medically necessary for
 the enrollee have been tried without successful outcomes; and

- (4) the United States Food and Drug Administration has approved the treating physician's individual patient investigational new drug application (IND) for the use of stiripental for treatment.
- 2.6 This paragraph does not apply to MinnesotaCare coverage under chapter 256L.

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