

116TH CONGRESS 2D SESSION

S. 3595

To require a longitudinal study on the impact of COVID-19.

IN THE SENATE OF THE UNITED STATES

May 4, 2020

Ms. Rosen (for herself and Mr. Rubio) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

A BILL

To require a longitudinal study on the impact of COVID— 19.

- 1 Be it enacted by the Senate and House of Representa-
- 2 tives of the United States of America in Congress assembled,
- 3 SECTION 1. SHORT TITLE.
- 4 This Act may be cited as the "Ensuring Under-
- 5 standing of COVID-19 to Protect Public Health Act".
- 6 SEC. 2. STUDY ON THE IMPACT OF COVID-19.
- 7 Part A of title IV of the Public Health Service Act
- 8 (42 U.S.C. 281 et seq.) is amended by adding at the end
- 9 the following:

1 "SEC. 4040, STUDY ON THE IMPACT OF COVID-19.

- 2 "(a) IN GENERAL.—The Director of NIH, in con-
- 3 sultation with the Director of the Centers for Disease Con-
- 4 trol and Prevention, shall conduct a longitudinal study,
- 5 over not less than 10 years, on the full impact of COVID-
- 6 19 on infected individuals, including both short-term and
- 7 long-term health impacts.
- 8 "(b) Timing.—The Director of NIH shall begin en-
- 9 rolling patients in the study under this section not later
- 10 than 6 months after the date of enactment of this section.
- 11 "(c) REQUIREMENTS.—The study under this section
- 12 shall—
- 13 "(1) be nationwide;
- "(2) include diversity of enrollees to account for
- gender, age, race, ethnicity, geography,
- 16 comorbidities, and underrepresented populations, in-
- 17 cluding pregnant and lactating women;
- 18 "(3) study individuals who were infected with
- 19 COVID-19 who experienced mild symptoms, such
- 20 individuals who experienced moderate symptoms,
- and such individuals who experienced severe symp-
- 22 toms;
- 23 "(4) monitor the health outcomes and symp-
- toms of individuals who were infected with COVID-
- 25 19, or had prenatal exposure to COVID-19, includ-
- ing lung capacity and function, and immune re-

- sponse, taking into account any pharmaceutical
- 2 interventions such individuals may have received;
- 3 "(5) monitor the mental health outcomes of in-
- 4 dividuals infected with COVID-19, taking into ac-
- 5 count any interventions that affected mental health;
- 6 and
- 7 "(6) monitor individuals enrolled in the study
- 8 not less frequently than twice per year after the first
- 9 year of the individual's infection with COVID-19.
- 10 "(d) Public-Private Research Network.—For
- 11 purposes of carrying out the study under this section, the
- 12 Director of NIH may develop a network of public-private
- 13 research partners, provided that all research, including the
- 14 research carried out through any such partner, is available
- 15 publicly.
- 16 "(e) Summaries of Findings.—The Director of
- 17 NIH shall make public a summary of findings under this
- 18 section not less frequently than once every 3 months for
- 19 the first 2 years of the study, and not less frequently than
- 20 every 6 months thereafter. Such summaries may include
- 21 information about how the findings of the study under this
- 22 section compare with findings from research conducted
- 23 abroad.

- 1 "(f) AUTHORIZATION OF APPROPRIATIONS.—There
- 2 are authorized to be appropriated such sums as may be

3 necessary to carry out this section.".

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