STATE OF NEW YORK

4497

2019-2020 Regular Sessions

IN SENATE

March 13, 2019

Introduced by Sen. RIVERA -- read twice and ordered printed, and when printed to be committed to the Committee on Health

AN ACT to amend the public health law, in relation to establishing the rare disease advisory council

The People of the State of New York, represented in Senate and Assembly, do enact as follows:

Section 1. The public health law is amended by adding a new article 27-L to read as follows:

ARTICLE 27-L

RARE DISEASE ADVISORY COUNCIL

5 Section 2799-u. Rare disease advisory council.

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- § 2799-u. Rare disease advisory council. 1. There shall be established within the department a rare disease advisory council to advise and consult on policy matters relating to rare disease. For the purposes of this section, rare disease shall mean any disease or condition that affects less than 200,000 people in the United States.
- 10 2. The council shall be composed of twenty-one members consisting of: seventeen members shall be appointed by the governor, two members shall 12 13 be appointed by the speaker of the assembly, and two members shall be 14 appointed by the temporary president of the senate. The governor shall 15 designate the commissioner, or his or her designee, as chair of the 16 council, and the superintendent of the department of financial services, or his or her designee, as co-chair of the council. The council shall at 17 all times include in its membership the following, which shall be 18 19 appointed by the governor: two physicians licensed and practicing in the 20 state with experience researching, diagnosing, or treating rare 21 <u>diseases; one registered nurse or advanced practice registered nurse</u> 22 licensed and practicing in the state with experience treating rare 23 diseases; two hospital administrators from hospitals in the state that 24 provide care to persons diagnosed with a rare disease, one of whom must 25 represent a hospital in which the scope of service focuses on rare

EXPLANATION--Matter in italics (underscored) is new; matter in brackets [-] is old law to be omitted.

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diseases of pediatric patients; three persons age eighteen or older who either have a rare disease or are a caregiver of a person with a rare disease; two representatives of a rare disease patient organization that operates in the state; one pharmacist with experience with drugs used to treat rare diseases; one representative of the biotechnology industry; one representative of health insurance companies; one member of the scientific community who is engaged in rare disease research; and one genetic counselor with experience providing services to persons diagnosed with a rare disease or caregivers of those persons.

- 3. Members of the advisory council shall serve at the pleasure of the appointing authority. Vacancies on the advisory council shall be filled in the same manner as the original appointment. Members of the advisory council shall receive no compensation for their services, but may be reimbursed for necessary and actual expenses incurred in the performance of their duties hereunder.
- 4. The advisory council shall meet at least three times in each calendar year. The council shall meet at the request of its chairs.
 - 5. The advisory council's duties may include, but are not limited to:
- a. Convening public hearings, and receiving and considering reports and testimony from individuals, the department, health care providers, and public or private organizations recognized as having expertise in rare diseases, as well as other interested parties, to assist the council in developing recommendations under this section;
- b. Studying the economic burden of the treatment of rare diseases with regard to quality of care, health care coverage, and access to treatment and services;
 - c. Developing recommendations pertaining to the impact of particular health care coverage and the impact of particular coverage on the provision of treatment and care services;
- 30 <u>d. Identifying best practices for rare diseases that could improve</u> 31 <u>treatment of rare diseases in New York state; and</u>
 - e. Developing a list of existing, publicly accessible resources on research, diagnosis, treatment, coverage options, and education relating to rare diseases, which shall be published on the department's website upon completion.
- 6. The council shall prepare and submit a written report no later than
 Cotober first, two thousand twenty, and biennially thereafter, to the
 governor, speaker of the assembly, and temporary president of the senate
 that sets forth the findings and recommendations of the council on
 matters within the scope of its duties as set forth in this section.
 - § 2. This act shall take effect immediately.