SECOND REGULAR SESSION

HOUSE COMMITTEE SUBSTITUTE FOR

HOUSE BILL NO. 2407

99TH GENERAL ASSEMBLY

6334H.02C D. ADAM CRUMBLISS, Chief Clerk

AN ACT

To amend chapter 208, RSMo, by adding thereto one new section relating to an advisory council on rare diseases within the MO HealthNet division.

Be it enacted by the General Assembly of the state of Missouri, as follows:

Section A. Chapter 208, RSMo, is amended by adding thereto one new section, to be known as section 208.183, to read as follows:

208.183. 1. The "Advisory Council on Rare Diseases and Personalized Medicine" is hereby established within the MO HealthNet division. The advisory council on rare diseases and personalized medicine shall serve as an expert advisory committee to the drug utilization review board, providing necessary consultation to the board when the board makes recommendations or determinations regarding beneficiary access to drugs or biological products for rare diseases, or when the board itself determines that it lacks the specific scientific, medical, or technical expertise necessary for the proper performance of its responsibilities and the necessary expertise can be provided by external in-state experts.

2. The advisory council on rare diseases and personalized medicine shall be composed of the following health care professionals, who shall be appointed by the director of the department of social services:

(1) Two physicians affiliated with public schools of medicine who are licensed and practicing in this state with experience researching, diagnosing, or treating rare diseases;

(2) Two physicians affiliated with private schools of medicine headquartered in this state who are licensed and practicing in this state with experience researching, diagnosing, or treating rare diseases;

EXPLANATION — Matter enclosed in bold-faced brackets **[**thus**]** in the above bill is not enacted and is intended to be omitted from the law. Matter in **bold-face** type in the above bill is proposed language.

(3) A physician who holds a doctor of osteopathy degree and is active in medical practice and affiliated with a school of medicine in this state with experience researching, diagnosing, or treating rare diseases;

(4) Two medical researchers from either academic research institutions or medical research organizations in this state who have received federal or foundation grant funding for rare disease research;

(5) A registered nurse or advanced practice registered nurse licensed and practicing in this state with experience treating rare diseases;

(6) A pharmacist practicing in a hospital in this state that has a designated orphan disease center;

(7) A professor employed by a pharmacy program in this state that is fully accredited by the Accreditation Council for Pharmacy Education who has advanced scientific or medical training in orphan and rare disease treatments;

(8) One individual representing the rare disease community or who is living with a rare disease;

(9) One member who represents a rare disease foundation;

(10) A representative from a rare disease center located within one of the state's comprehensive pediatric hospitals;

(11) The chair of the joint committee on the life sciences or the chair's designee; and

(12) The chairperson of the drug utilization review board, or the chairperson's designee, who shall serve as an ex officio, nonvoting member of the advisory council.

3. The director shall convene the first meeting of the advisory council on rare diseases and personalized medicine no later than February 28, 2019. Following the first meeting, the advisory council shall meet upon the call of the chairperson of the drug utilization review board or upon the request of a majority of the council members.

4. The drug utilization review board, when making recommendations or determinations regarding beneficiary access to drugs and biological products for rare diseases, as defined in the federal Orphan Drug Act of 1983, Pub. L. 97-414, and drugs and biological products that are approved by the United States Food and Drug Administration and within the emerging fields of personalized medicine and noninheritable gene editing therapeutics, shall request and consider information from the advisory council on rare diseases and personalized medicine. "Beneficiary access", as used in this subsection, means developing prior authorization and reauthorization criteria for a rare disease drug, including placement on a preferred drug list or a formulary, payment, cost-sharing, drug utilization review, or medication therapy management.

5. The board shall seek the input of the advisory council on rare diseases and personalized medicine to address topics for consultation under this section including, but not limited to:

(1) Rare diseases;

(2) The severity of rare diseases;

(3) The unmet medical need associated with rare diseases;

(4) The impact of particular coverage, cost-sharing, tiering, utilization management, prior authorization, medication therapy management, or other Medicaid policies on access to rare disease therapies;

(5) An assessment of the benefits and risks of therapies to treat rare diseases;

(6) The impact of particular coverage, cost-sharing, tiering, utilization management, prior authorization, medication therapy management, or other policies on patients' adherence to the treatment regimen prescribed or otherwise recommended by their physicians;

(7) Whether beneficiaries who need treatment from or a consultation with a rare disease specialist have adequate access and, if not, what factors are causing the limited access; and

(8) The demographics and the clinical description of patient populations.

6. Nothing in this section shall be construed to create a legal right for a consultation on any matter or require the drug utilization review board to meet with any particular expert or stakeholder.

7. Recommendations of the advisory council on rare diseases and personalized medicine on an applicable treatment of a rare disease shall be explained in writing to members of the board during public hearings.

8. For purposes of this section, a "rare disease drug" is a drug used to treat a rare medical condition, defined as any disease or condition that affects fewer than two hundred thousand persons in the United States, such as cystic fibrosis, hemophilia, and multiple myeloma.

9. All members of the advisory council on rare diseases and personalized medicine shall annually sign a conflict of interest statement revealing economic or other relationships with entities that could influence a member's decisions, and at least twenty percent of the advisory council members shall not have a conflict of interest with respect to any insurer, pharmaceutical benefits manager, or pharmaceutical manufacturer.

✓