1 ENGROSSED SENATE BILL NO. 207 By: Woods and Hicks of the 2 Senate 3 and West (Josh) and Waldron of 4 the House 5 6 7 [ public health - Council - report - educational and newborn screening programs - requirement - report codification - effective date -8 emergency ] 9 10 11 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 12 SECTION 1. NEW LAW A new section of law to be codified in the Oklahoma Statutes as Section 1-451 of Title 63, unless there 13 is created a duplication in numbering, reads as follows: 14 The Oklahoma Rare Disease Advisory Council shall be 15 Α. established within the State Department of Health in accordance with 16 this section. 17 The purpose of the Council shall be to provide guidance and 18 Β. recommendations to educate the public, the Legislature, and other 19 state agencies, as appropriate, on the needs of individuals with 20 rare diseases living in this state. 21 С. The Council shall conduct the following activities to 22 benefit those impacted by rare diseases in this state: 23 24

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Convene public hearings, make inquiries, and solicit
 comments from the general public in Oklahoma to assist the Council
 with a first-year landscape or survey of the needs of rare disease
 patients, caregivers, and providers in the state;

2. Provide testimony and comments on pending legislation and
regulations before the Legislature, if called, and other state
agencies that impact Oklahoma's rare disease community;

3. Consult with experts on rare diseases to develop policy
9 recommendations to improve patient access to, and quality of, rare
10 disease specialists, affordable and comprehensive health care
11 coverage, relevant diagnostics, timely treatment, and other needed
12 services;

4. Establish best practices and protocols to include in state
planning related to natural disasters, public health emergencies, or
other emergency declarations to enable continuity of care for rare
disease patients and ensure safeguards against discrimination for
rare disease patients are in place;

18 5. Identify areas of unmet need for research and opportunities 19 for collaboration with stakeholders and other states' rare disease 20 advisory councils that can inform future studies and work done by 21 the Council; and

6. Research and identify best practices to reduce health
disparities and achieve health equity in the research, diagnosis,
and treatment of rare disease in this state.

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D. 1. The Council's appointment process shall be conducted in a transparent manner to provide interested individuals an opportunity to apply for membership on the Council. All members of the Council shall be full-time residents of this state as practicable. Membership shall include a diverse set of stakeholders representative of the geographic and population diversity of the state.

8 2. The Governor shall appoint the chair of the Council not 9 later than December 1, 2025. The chair shall not hold any position 10 within the government of this state.

3. The chair shall appoint no fewer than thirteen (13) members,
 including the following:

- a. one representative of an academic research institution
  in the state that receives any grant funding for rare
  disease research,
- b. one representative of the Oklahoma Health CareAuthority,
- 18 c. one representative of the Insurance Department,
- d. one Registered Nurse or Advanced Practice Registered
   Nurse licensed and practicing in this state with
   experience treating rare diseases,
- e. one physician practicing in this state with experiencetreating rare diseases,
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1	f. one hospital administrator, or his or her designee,
2	from a hospital in this state that provides care to
3	persons diagnosed with a rare disease,
4	g. at least two patients who have a rare disease,
5	h. at least one caregiver of a person with a rare
6	disease,
7	i. one representative of a rare disease patient
8	organization that operates in this state,
9	j. one representative of the biopharmaceutical industry,
10	k. one representative of a health plan company, and
11	1. one member of the scientific community who is engaged
12	in rare disease research including, but not limited
13	to, a medical researcher with experience conducting
14	research on rare diseases.
15	4. The chair may appoint additional members on an ad hoc basis.
16	E. 1. The Council shall electronically submit a report to the
17	President Pro Tempore of the Senate, the Speaker of the House of
18	Representatives, and the Governor within one (1) year of
19	establishment of the Oklahoma Rare Disease Advisory Council and
20	annually thereafter. Prior to submission, a draft of the annual
21	report shall be made available for public comment and discussed at
22	an open public meeting.
23	2. Annual reports shall:
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1 describe the activities and progress of the Council a. under this section, and 2 provide recommendations to the Legislature and 3 b. Governor on ways to address the needs of people living 4 5 with rare diseases in this state. The initial meeting of the Council shall occur no later 6 F. 1. than February 1, 2026. Thereafter, the Council shall meet no less 7 than quarterly. 8 2. Meetings of the Council shall be conducted in accordance 9 with the Oklahoma Open Meeting Act. 10 The Council shall: G. 11 12 1. Provide opportunities for the public to hear updates and provide input into its work; and 13 2. Create and maintain a public website where meeting minutes, 14 notices of upcoming meetings, and public comments may be submitted. 15 Η. 1. The Governor shall appoint the chair for an initial term 16 of three (3) years. At the end of the chair's initial three-year 17 term, and every two (2) years thereafter, members of the Council 18 shall elect, by a majority vote, a new chair. 19 2. Council members shall serve no longer than three (3) years, 20 except that, to facilitate a staggered rotation of members to retain 21 continuity and knowledge transfer, during the initial five (5) years 22 after the establishment of the Council, members may serve up to a 23 four-year term. 24

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3. If a vacancy occurs, the Council, by a majority vote, shall
 fill such vacancy in a timely manner and in compliance with
 requirements set forth in subsection D of this section.

I. As used in this section, "rare disease", sometimes called an
orphan disease, means a disease that affects fewer than two hundred
thousand (200,000) people in the United States.

SECTION 2. AMENDATORY 63 O.S. 2021, Section 1-533, as
amended by Section 1, Chapter 161, O.S.L. 2022 (63 O.S. Supp. 2024,
Section 1-533), is amended to read as follows:

10 Section 1-533. A. The State Commissioner of Health shall 11 provide, pursuant to the provisions of Section 1-534 of this title, 12 as technologies and funds become available, an intensive educational 13 and newborn screening program among physicians, hospitals, public 14 health nurses, and the public concerning phenylketonuria, related 15 inborn metabolic disorders, and other genetic or biochemical 16 disorders for which:

Newborn screening will provide early treatment and
 management opportunities that might not be available without
 screening; and

20 2. Treatment and management will prevent intellectual
 21 disabilities and/or reduce infant morbidity and mortality.

B. This educational and newborn screening program shall includeinformation about:

24 1. The nature of the diseases;

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2. Examinations for the detection of the diseases in infancy;
 and

3 3. Follow-up measures to prevent the morbidity and mortality4 resulting from these diseases.

5 C. For purposes of this section, "phenylketonuria" means an inborn error of metabolism attributable to a deficiency of or a 6 defect in phenylalanine hydroxylase, the enzyme that catalyzes the 7 conversion of phenylalanine to tyrosine. The deficiency permits the 8 9 accumulation of phenylalanine and its metabolic products in the body fluids. The deficiency can result in intellectual disabilities 10 (phenylpyruvic oligophrenia), neurologic manifestations (including 11 hyperkinesia, epilepsy, and microcephaly), light pigmentation, and 12 eczema. The disorder is transmitted as an autosomal recessive trait 13 and can be treated by administration of a diet low in phenylalanine. 14 To the extent practicable, the <del>list of</del> disorders screened 15 D. for under this section shall be identical to at a minimum include 16 those listed in the Recommended Uniform Screening Panel of the 17 United States Department of Health and Human Services. 18

E. On November 1, 2026, and each November 1 thereafter, the
State Department of Health shall compile an annual report listing
the disorders screened for under this section, specifying any
disorders added since the prior annual report, and detailing efforts
being undertaken to add additional disorders. The report shall be
published on the Department's website and shall be submitted

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1	electronically to the President Pro Tempore of the Senate, the
2	Speaker of the House of Representatives, the Governor, and the
3	Oklahoma Rare Disease Advisory Council established under Section 1
4	of this act.
5	<u>F.</u> The Commissioner shall promulgate any rules necessary to
6	effectuate the provision of this section.
7	SECTION 3. This act shall become effective July 1, 2025.
8	SECTION 4. It being immediately necessary for the preservation
9	of the public peace, health or safety, an emergency is hereby
10	declared to exist, by reason whereof this act shall take effect and
11	be in full force from and after its passage and approval.
12	Passed the Senate the 27th day of March, 2025.
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14	Presiding Officer of the Senate
14	Presiding Officer of the Senate
14 15	Presiding Officer of the Senate Passed the House of Representatives the day of,
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14 15 16 17 18 19	Passed the House of Representatives the day of, 2025. Presiding Officer of the House
14 15 16 17 18 19 20	Passed the House of Representatives the day of, 2025. Presiding Officer of the House
14 15 16 17 18 19 20 21	Passed the House of Representatives the day of, 2025. Presiding Officer of the House