

Senate Substitute for HOUSE BILL No. 2149

By Committee on Public Health and Welfare

3-19

1 AN ACT concerning the Kansas program of medical assistance; relating to
2 donor human breast milk and medications used under medicaid;
3 amending K.S.A. 2014 Supp. 39-7,119, 39-7,120 and 39-7,121b and
4 repealing the existing sections.

5
6 *Be it enacted by the Legislature of the State of Kansas:*

7 New Section 1. (a) The department of health and environment shall
8 reimburse a medical care facility for prescribed medically necessary donor
9 human breast milk provided to a recipient of medical assistance under the
10 Kansas program of medical assistance if:

11 (1) Such recipient is:

12 (A) An infant under the age of three months;

13 (B) critically ill; and

14 (C) in the neonatal intensive care unit of the hospital;

15 (2) a person licensed to practice medicine and surgery orders the
16 donor human breast milk for the recipient;

17 (3) the department determines that the donor human breast milk is
18 medically necessary for the recipient;

19 (4) the parent or legal guardian of the recipient signs and dates an
20 informed consent form indicating the risks and benefits of using banked
21 donor human breast milk; and

22 (5) the donor human breast milk is obtained from a donor human
23 breast milk bank that meets the quality requirements established by the
24 department of health and environment.

25 (b) An electronic prior authorization system that uses the best medical
26 evidence and care and treatment guidelines consistent with national
27 standards shall be used by the department to determine medical necessity.

28 (c) The department shall promulgate rules and regulations necessary
29 to implement the provisions of this section *{prior to July 1, 2016}*.

30 (d) The department shall implement and administer the provisions of
31 this section in a manner consistent with applicable federal laws and
32 regulations. The department shall seek any necessary approvals of the
33 federal government that are required for the implementation of this
34 section.

35 (e) As used in this section:

36 (1) "Department" means the department of health and environment.

1 (2) "Medical care facility" shall mean the same as in K.S.A. 65-425,
2 and amendments thereto.

3 Sec. 2. K.S.A. 2014 Supp. 39-7,119 is hereby amended to read as
4 follows: 39-7,119. (a) There is hereby created the medicaid drug utilization
5 review board which shall be responsible for the implementation of
6 retrospective and prospective drug utilization programs under the Kansas
7 medicaid program.

8 (b) Except as provided in subsection (i), the board shall consist of at
9 least seven members appointed as follows:

10 (1) Two licensed physicians actively engaged in the practice of
11 medicine, nominated by the Kansas medical society and appointed by the
12 secretary of health and environment from a list of four nominees;

13 (2) one licensed physician actively engaged in the practice of
14 osteopathic medicine, nominated by the Kansas association of osteopathic
15 medicine and appointed by the secretary of health and environment from a
16 list of four nominees;

17 (3) two licensed pharmacists actively engaged in the practice of
18 pharmacy, nominated by the Kansas pharmacy association and appointed
19 by the secretary of health and environment from a list of four nominees;

20 (4) one person licensed as a pharmacist and actively engaged in
21 academic pharmacy, appointed by the secretary of health and environment
22 from a list of four nominees provided by the university of Kansas; *and*

23 (5) one licensed professional nurse actively engaged in long-term
24 care nursing, nominated by the Kansas state nurses association and
25 appointed by the secretary of health and environment from a list of four
26 nominees.

27 (c) The secretary of health and environment may add two additional
28 members so long as no class of professional representatives exceeds 51%
29 of the membership.

30 (d) The physician and pharmacist members shall have expertise in the
31 clinically appropriate prescribing and dispensing of outpatient drugs.

32 (e) The appointments to the board shall be for terms of three years. In
33 making the appointments, the secretary of health and environment shall
34 provide for geographic balance in the representation on the board to the
35 extent possible. Subject to the provisions of subsection (i), members may
36 be reappointed.

37 (f) The board shall elect a chairperson from among board members
38 who shall serve a one-year term. The chairperson may serve consecutive
39 terms.

40 (g) The board, in accordance with K.S.A. 75-4319, and amendments
41 thereto, may recess for a closed or executive meeting when it is
42 considering matters relating to identifiable patients or providers.

43 (h) All actions of the medicaid drug utilization review board shall be

1 upon the affirmative vote of five members of the board and the vote of
2 each member present when action was taken shall be recorded by roll call
3 vote.

4 (i) Upon the expiration of the term of office of any member of the
5 medicaid drug utilization review board on or after the effective date of this
6 act and in any case of a vacancy existing in the membership position of
7 any member of the medicaid drug utilization review board on or after the
8 effective date of this act, a successor shall be appointed by the secretary of
9 health and environment so that as the terms of members expire, or
10 vacancies occur, members are appointed and the composition of the board
11 is changed in accordance with the following and such appointment shall be
12 made by the secretary of health and environment in the following order of
13 priority:

14 (1) One member shall be a licensed pharmacist who is actively
15 performing or who has experience performing medicaid pharmacy services
16 for a hospital and who is nominated by the Kansas hospital association and
17 appointed by the secretary of health and environment from a list of two or
18 more nominees;

19 (2) one member shall be a licensed pharmacist who is actively
20 performing or who has experience performing medicaid pharmacy services
21 for a licensed adult care home and who is nominated by the state board of
22 pharmacy and appointed by the secretary of health and environment from a
23 list of two or more nominees;

24 (3) one member shall be a licensed physician who is actively engaged
25 in the general practice of allopathic medicine and who has practice
26 experience with the state medicaid plan and who is nominated by the
27 Kansas medical society and appointed by the secretary of health and
28 environment from a list of two or more nominees;

29 (4) one member shall be a licensed physician who is actively engaged
30 in mental health practice providing care and treatment to persons with
31 mental illness, who has practice experience with the state medicaid plan
32 and who is nominated by the Kansas psychiatric society and appointed by
33 the secretary of health and environment from a list of two or more
34 nominees;

35 (5) one member shall be a licensed physician who is the medical
36 director of a nursing facility, who has practice experience with the state
37 medicaid plan and who is nominated by the Kansas medical society and
38 appointed by the secretary of health and environment from a list of two or
39 more nominees;

40 (6) one member shall be a licensed physician who is actively engaged
41 in the general practice of osteopathic medicine, who has practice
42 experience with the state medicaid plan and who is nominated by the
43 Kansas association of osteopathic medicine and who is appointed by the

1 secretary of health and environment from a list of two or more nominees;

2 (7) one member shall be a licensed pharmacist who is actively
3 engaged in retail pharmacy, who has practice experience with the state
4 medicaid plan and who is nominated by the state board of pharmacy and
5 appointed by the secretary of health and environment from a list of two or
6 more nominees;

7 (8) one member shall be a licensed pharmacist who is actively
8 engaged in or who has experience in research pharmacy and who is
9 nominated jointly by the Kansas task force for the pharmaceutical research
10 and manufacturers association and the university of Kansas and appointed
11 by the secretary of health and environment from a list of two or more
12 jointly nominated persons; and

13 (9) one member shall be a licensed advanced practice registered nurse
14 or physician assistant actively engaged in the practice of providing the
15 health care and treatment services such person is licensed to perform, who
16 has practice experience with the state medicaid plan and who is nominated
17 jointly by the Kansas state nurses' association and the Kansas academy of
18 physician assistants and appointed by the secretary of health and
19 environment from a list of two or more jointly nominated persons.

20 (j) *The medicaid drug utilization review board shall meet at least*
21 *quarterly and such meetings shall be open to the public and shall provide*
22 *an opportunity for public comments. The board shall post notice of such*
23 *meetings at least 14 business days before the scheduled meetings.*

24 Sec. 3. K.S.A. 2014 Supp. 39-7,120 is hereby amended to read as
25 follows: 39-7,120. (a) ~~The secretary of health and environment shall not~~
26 ~~restrict patient access to prescription-only drugs pursuant to a program of~~
27 ~~prior authorization or a restrictive formulary except by rules and~~
28 ~~regulations adopted in accordance with K.S.A. 75-5625, and amendments~~
29 ~~thereto. Prior to the promulgation of any such rules and regulations, the~~
30 ~~secretary of health and environment shall submit such proposed rules and~~
31 ~~regulations to the medicaid drug utilization review board for written~~
32 ~~comment~~ *may implement prior authorization of any new prescription-only*
33 *drugs until such drugs are reviewed by the medicaid drug utilization*
34 *review board at the next scheduled meeting. New drugs shall be approved*
35 *for use when such drugs are used within package insert guidelines*
36 *approved by the federal food and drug administration and clinically*
37 *reputable compendia, such as the United States pharmacopeia, as*
38 *approved by the secretary of health and environment in the rules and*
39 *regulations, during the period before such drugs are reviewed by the*
40 *medicaid drug utilization review board. The secretary of health and*
41 *environment may not implement permanent prior authorization until 30*
42 *days after receipt of comments by the drug utilization review board.*

43 (b) When considering recommendations from the medicaid drug

1 utilization review board regarding the prior authorization of a drug, the
2 secretary of health and environment shall consider the net economic
3 impact of such prior authorization, including, but not limited to, the costs
4 of specific drugs, rebates or discounts pursuant to 42 U.S.C. § 1396r-8,
5 dispensing costs, dosing requirements and utilization of other drugs or
6 other medicaid health care services which may be related to the prior
7 authorization of such drug.

8 Sec. 4. K.S.A. 2014 Supp. 39-7,121b is hereby amended to read as
9 follows: 39-7,121b. *(a) No requirements for prior authorization or other*
10 *restrictions on medications used to treat mental illnesses—such as—*
11 *schizophrenia, depression or bipolar disorder may be imposed on medicaid*
12 *recipients. Medications that will be available under the state medicaid plan*
13 *without restriction for persons with mental illnesses shall include atypical*
14 *antipsychotic medications, conventional antipsychotic medications and*
15 *other medications used for the treatment of mental illnesses., except on*
16 *medications subject to guidelines developed by the drug utilization review*
17 *board according to subsection (c). None of the following shall be*
18 *construed as restrictions under this subsection:*

19 *(1) Any alert to a pharmacist that does not deny the claim and can be*
20 *overridden by the pharmacist;*

21 *(2) prescriber education activities; or*

22 *(3) the consolidation of dosing regimens to equivalent doses—~~and~~*
23 *~~other such dose optimization policies.~~*

24 *(b) The mental health medication advisory committee shall provide*
25 *recommendations to the drug utilization review board for the*
26 *purpose of developing guidelines. The drug utilization review board may*
27 *accept the recommendations of the mental health medication advisory*
28 *committee in whole and such recommendations shall take effect*
29 *immediately upon such approval. The drug utilization review board may*
30 *reject the recommendations of the mental health medication advisory*
31 *committee in whole and such recommendations shall be referred back*
32 *to the mental health medication advisory committee for further*
33 *consideration. No medication guidelines related to mental health*
34 *medications shall be adopted by the drug utilization review board without*
35 *recommendations made by the mental health medication advisory*
36 *committee.*

37 *(c) For the medications used to treat mental illness that are available*
38 *for use on July 1, 2015, the drug utilization review board shall review all*
39 *such medications prior to July 1, 2016. For medications used to*
40 *treat mental illness that do not exist on July 1, 2015, but are later*
41 *developed or believed to be effective in the treatment of mental illness, the*
42 *drug utilization board shall review all such medications within six months*
43 *of presentation to the drug utilization review board.*

1 (d) *The mental health medication advisory committee is hereby*
2 *established.*

3 (1) *The mental health medication advisory committee shall be*
4 *appointed by the secretary of health and environment and consist of nine*
5 *members; including the secretary of health and environment, or the*
6 *secretary's designee, who shall be the chair of the committee; two*
7 *persons licensed to practice medicine and surgery with board certification*
8 *in psychiatry nominated by the Kansas psychiatric society, one of whom*
9 *specializes in geriatric mental health; two persons licensed to practice*
10 *medicine and surgery with board certification in psychiatry nominated by*
11 *the association of community mental health centers of Kansas, one of*
12 *whom specializes in pediatric mental health; two pharmacists nominated*
13 *by the Kansas pharmacy association; one person licensed to practice*
14 *medicine and surgery nominated by the Kansas medical society; and one*
15 *advanced practice registered nurse engaged in a role of mental health*
16 *nominated by the Kansas state nurses association. All nominating bodies*
17 *shall provide two nominees for each position for which they provide*
18 *nominations, with the secretary selecting the appointee from the provided*
19 *nominees.*

20 (2) *The mental health medication advisory committee shall meet*
21 *upon the request of the chair of the mental health medication advisory*
22 *committee, but shall meet at least one time each quarter.*

23 (3) *Members of the mental health medication advisory committee are*
24 *entitled to compensation and expenses as provided in K.S.A. 75-3223, and*
25 *amendments thereto. Members of the committee attending committee*
26 *meetings shall be paid mileage and all other applicable expenses,*
27 *provided such expenses are consistent with policies established by the*
28 *secretary of health and environment.*

29 Sec. 5. K.S.A. 2014 Supp. 39-7,119, 39-7,120 and 39-7,121b are
30 hereby repealed.

31 Sec. 6. This act shall take effect and be in force from and after its
32 publication in the statute book.