1 HOUSE OF REPRESENTATIVES - FLOOR VERSION 2 STATE OF OKLAHOMA 1st Session of the 60th Legislature (2025) 3 COMMITTEE SUBSTITUTE 4 FOR 5 HOUSE BILL NO. 1600 By: Gise, Williams, Pae, Roberts, Caldwell (Chad), Adams, Steagall, Harris, 6 and **Kelley** of the House 7 and 8 Hines of the Senate 9 10 11 COMMITTEE SUBSTITUTE 12 An Act relating to health care; enacting the Lori Brand Patient Bill of Rights Act of 2025; creating a list of rights for a patient seeking treatment; 1.3 specifying certain responsibilities of patients 14 seeking treatment; creating certain rights for minor patients seeking treatment; specifying certain 15 responsibilities of parents of minor patients seeking treatment in this state; providing for codification; 16 and providing an effective date. 17 18 19 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 20 SECTION 1. NEW LAW A new section of law to be codified 21 in the Oklahoma Statutes as Section 3401 of Title 63, unless there 22 is created a duplication in numbering, reads as follows: 23 This act shall be known and may be cited as the "Lori Brand 24 Patient Bill of Rights Act of 2025".

- SECTION 2. NEW LAW A new section of law to be codified
 in the Oklahoma Statutes as Section 3401.1 of Title 63, unless there
 is created a duplication in numbering, reads as follows:
 - A. Each patient treated in this state shall have the following rights when being treated:
 - 1. The right to considerate and respectful care, provided in a safe environment, free from all forms of abuse, neglect, harassment, and exploitation;
 - 2. To receive information in a manner that he or she understands. Communications with the patient shall be effective and provided in a manner that facilitates understanding by the patient. Written information provided will be appropriate to the age, understanding, and, as appropriate, the language of the patient. As appropriate, communications specific to the vision-, speech-, hearing-, cognitive-, and language-impaired patient will be provided. The hospital shall meet the requirements of federal regulations that require program and facility accessibility;
 - 3. To receive as much information about any proposed treatment or procedure as he or she may need in order to give informed consent or to refuse the course of treatment. Except in emergencies, this shall include a description of the procedure or treatment, the medically significant risks involved in the procedure or treatment, alternate courses of treatment or nontreatment and the risks

involved in each, and the name of the person who will carry out the procedure or treatment;

- 4. To have an advance directive attorney for health care concerning treatment or to designate a surrogate decision-maker with the expectation that the hospital will honor the intent of that directive to the extent allowed by law and hospital policy. The health care provider must advise a patient of his or her rights under state law and hospital policy to make informed medical decisions, ask if the patient has an advance directive, and include that information in patient records. The patient has the right to timely information about hospital policy that may limit its ability to implement a legally valid advance directive;
- 5. To participate in the development and implementation of his or her plan of care and actively participate in decisions regarding his or her medical care;
- 6. To accept medical care or to refuse treatment, to the extent permitted by law, and to be informed of the consequences of such refusal;
- 7. To become informed of his or her rights as a patient in advance of, or when discontinuing, the provision of care. The patient may appoint a representative to receive this information should he or she so desire;
- 8. To have a family member or representative of his or her choice notified promptly of his or her admission to the hospital;

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- 9. To request that no information regarding his or her admittance, diagnosis, or treatment be released;
- 10. To review the records and obtain a copy of the medical records pertaining to his or her medical care and to have the information explained or interpreted as necessary, except when restricted by law;
- 11. To reasonable continuity of care, when appropriate, and to be informed by the doctor and other caregivers of available and realistic patient care options when hospital care is no longer appropriate;
- 12. To confidential treatment of all communications and records pertaining to his or her care and stay at the hospital;
- 13. To expect that, within its capability, capacity, and policies, the hospital will make a reasonable response to the request of a patient for appropriate and medically directed care and services. The hospital must provide evaluation, service, and a referral as indicated by the urgency of the case. When medically appropriate and legally permissible, or when a patient has requested a transfer, that patient may be transferred to another facility. That facility must have first accepted the patient for transfer. The patient must also have the benefit of the complete information and explanation concerning the need for, risks and benefits of, and alternatives to such a transfer;

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- 14. The patient or patient's representative has the right to participate in the consideration of ethical issues that might arise in the care of the patient. The hospital shall have a mechanism for the consideration of ethical issues arising in the care of patients and to provide education to caregivers and patients on ethical issues in health care;
- 15. To be advised of the hospital's complaint or grievance process should the patient wish to communicate a concern regarding the quality of care he or she receives. This includes whom to contact to file a complaint. The patient will be provided with a written notice of the complaint determination that contains the name of the hospital's contact person, the steps taken on his or her behalf to investigate the complaint, the results of the complaint and, when possible, the resolution of the complaint concerning the quality of care;
- 16. To examine and receive an explanation of his or her bill regardless of source of payment;
- 17. To remain free from restraints or seclusion in any forms that are not medically necessary or are used as a means of coercion, discipline, convenience, or retaliation by staff;
- 18. To receive the visitors whom he or she designates, including, but not limited to, a spouse, a domestic partner, including a same-sex domestic partner, another family member, or a friend. The patient has the right to withdraw or deny consent at

- any time. Visitation will not be restricted, limited, or otherwise denied on the basis of race, color, national origin, religion, sex, or disability; and
 - 19. Through use of the hospital-issued notice of noncoverage, Medicare beneficiaries have the right to be informed in advance of procedures or treatment for which Medicare may deny payment, and that the beneficiary may be personally responsible for full payment if Medicare denies payment.
- B. A patient, guardian of a patient, or legally authorized representative of a patient shall have the following responsibilities:
 - 1. To provide accurate and complete information concerning the patient's present complaints, past illnesses, hospitalizations, medications, and other matters relating to his or her health;
 - 2. To report perceived risks in the patient's care and unexpected changes in his or her condition to the responsible health care provider;
 - 3. For the patient's actions should he or she refuse treatment or not follow his or her doctor's orders;
- 4. To ask questions when the patient does not understand what
 he or she has been told about the patient's care or what he or she
 is expected to do;
- 5. To be considerate of the rights of other patients and hospital personnel;

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- 6. To participate in educational and discharge planning activities necessary to ensure that he or she has adequate knowledge and support services to provide him or her with a safe environment upon discharge from the hospital;
- 7. To ask the doctor or nurse what to expect regarding pain management, to discuss pain relief options with doctors and nurses and to help develop a pain management plan, to ask for pain relief when pain first begins, to help doctors and nurses assess the patient's pain, to tell the doctors and nurses if his or her pain is not relieved, and to tell doctors and nurses about any concerns about taking pain medication;
- 8. For keeping appointments and for notifying the hospital or doctor when he or she is unable to do so;
- 9. Being respectful of his or her personal property and that of other patients in the hospital;
 - 10. Following hospital procedures; and
 - 11. Assuring that the financial obligations of his or her care is fulfilled as promptly as possible.
- C. Any minor patient has the following rights when being treated in this state:
 - 1. To be treated with respect in regards to:
 - a. each child and adolescent as a unique individual, and
 - b. the caretaking role and individual response of the parent and legal guardian;

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1	2. To provisions for normal physical and physiological needs of
2	a growing child including nutrition, rest, sleep, warmth, activity,
3	and freedom to move and explore. Minors shall have the right to:
4	a. appropriate treatment in the least restrictive
5	setting,
6	b. not receive unnecessary or excessive medication,
7	c. an individualized treatment plan and the right to
8	participate in the plan,
9	d. a humane treatment environment that provides
10	reasonable protection from harm and appropriate
11	privacy for personal needs,
12	e. separation from adult patients when possible, and
13	f. regular communication between the minor patient and
14	the patient's family or legal guardian;
15	3. To consistent, supportive, and nurturing care which:
16	a. meets the emotional and psychosocial needs of the
17	minor, and
18	b. fosters open communication;
19	4. To provisions for self-esteem needs which will be met by
20	attempts to give the minor:
21	a. the reassuring presence of a caring person, especially
22	a parent,
23	b. freedom to express feelings or fears with appropriate
24	reactions,

- c. as much control as possible over both self and situation,
 - d. opportunities to work through experiences before and after they occur, verbally, in play, or in other appropriate ways, and
 - e. recognition for coping well during difficult situations;
 - 5. To provisions for varied and normal stimuli of life which contributes to cognitive, social, emotional, and physical developmental needs such as play, educational, and social activities essential to all children and adolescents;
 - 6. To information about what to expect prior to, during, and following a procedure or experience and support in coping with it;
 - 7. To participate in decisions affecting his or her own medical treatment; and
 - 8. To the minimization of stay duration by recognizing discharge planning needs.
 - D. All parents and legal guardians of minor patients in this state shall have the following responsibilities:
 - 1. To continue in his or her parenting role to the extent of his or her ability; and
- 22 2. To be available to participate in decision-making and provide staff with knowledge of other parent or family whereabouts.

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1	SECTION 3. This act shall become effective November 1, 2025.
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3	COMMITTEE REPORT BY: COMMITTEE ON HEALTH AND HUMAN SERVICES OVERSIGHT, dated 03/05/2025 - DO PASS, As Amended and Coauthored.
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