1 STATE OF OKLAHOMA 2 1st Session of the 58th Legislature (2021) 3 SENATE BILL 344 By: McCortney 4 5 6 AS INTRODUCED 7 An Act relating to cancer; amending 63 O.S. 2011, Section 1-551.1, as last amended by Section 1, 8 Chapter 99, O.S.L. 2018 (63 O.S. Supp. 2020, Section 1-551.1), which relates to tumor registry; modifying 9 applicability of section; updating language; and providing an effective date. 10 11 12 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 13 SECTION 1. 63 O.S. 2011, Section 1-551.1, as AMENDATORY 14 last amended by Section 1, Chapter 99, O.S.L. 2018 (63 O.S. Supp. 15 2020, Section 1-551.1), is amended to read as follows: 16 Section 1-551.1. A. The State Commissioner of Health shall 17 establish and maintain an up-to-date tumor registry to ensure an 18 accurate and continuing source of data concerning such cancerous, 19 precancerous and tumorous diseases as the State Board of Health may 20 by rule specify. Such registry may include data necessary for 21 epidemiological surveys and scientific research, and other data 22 which is necessary and proper to further the recognition, 23 prevention, control, treatment and cure of cancer, precancerous and

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tumorous diseases.

- B. The Commissioner, pursuant to rules of the State Board of Health, shall require any hospital, clinic, laboratory, pathologist, physician or dentist, or any facility which provides diagnostic or treatment services for cancerous diseases and precancerous conditions, to report any or all data and information necessary for the purposes of this act section which may include the following:
- 1. Patient name, address, age, race, sex, Social Security number and hospital identifier or other identifier;
- 2. Patient's residential, family, environmental, occupational and medical histories; and
- 3. Physician's name, diagnosis, stage of the disease, method of treatment and the name and address of any facility providing treatment.
- C. The provisions of subsection B of this section shall not apply to ambulatory surgical centers, as defined by Section 2657 of this title, upon:
- 1. Upon submission of a signed affidavit that the ambulatory surgical center utilizes a sole source pathology laboratory to report any or all data and information necessary for the purposes of this act section; or
- 2. That are not certified by the Centers for Medicare and Medicaid Services.
- D. The Commissioner shall protect the identity of the patient and physician involved in any report required by this act section,

and may not release their identity without written consent, except that:

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- 1. The Commissioner may grant any person involved in a legitimate research activity access to confidential information obtained by the Department concerning individual patients if:
 - a. the research activity is determined to be in the interest of the public health and welfare,
 - b. the person conducting the research provides written information about the purpose of the research project, the nature of the data to be collected and how the researcher intends to analyze it, the records the researcher wishes to review, and the safeguards the researcher will take to protect the identity of the patients whose records the researcher will be reviewing,
 - c. the proposed safeguards are adequate to protect the identity of each patient whose records will be reviewed, and
 - d. an agreement is executed between the Commissioner of

 Health and the researcher that specifies the

 researcher's use of the records and that prohibits the

 publication or release of the names of individual

 cancer patients or any facts tending to lead to the

 identification of individual cancer patients;

2. Researchers may, with the approval of the Commissioner, use the names of individual patients when requesting additional information for research purposes or soliciting an individual patient's participation in a research project. However, if a researcher requests additional information or an individual patient's participation in a research project, the researcher must first obtain the written consent of the patient's attending physician. If the consent of the patient's attending physician is obtained, the researcher must then obtain the individual cancer patient's written consent by having the patient complete a release of confidential medical information form;

3. Data on patients may be shared with other registries, private or governmental, within or without the state, provided that a reciprocal data-sharing agreement, approved by the Commissioner, is implemented with that registry. Such agreements must include patient identification confidentiality requirements; and

- 4. Provided further, that any confidential information released by the Commissioner under this act section shall be deemed to be a confidential communication within the meaning of the physician-patient and the psychotherapist-patient privilege.
- E. Nothing in this act section shall be construed to compel any individual to submit to any medical examination, treatment or supervision of any kind; nor shall anyone providing information in accordance with this act section be deemed to be, or held liable

1 for, divulging confidential information. An individual shall have 2 the right to deny registration on religious grounds. 3 F. The State Board of Health is empowered to adopt reasonable 4 regulations Commissioner may promulgate rules to carry out the 5 provisions of this act section. 6 G. Any person who, in violation of a written agreement to 7 maintain confidentiality, willfully discloses any information 8 provided pursuant to this section shall be denied further access to 9 any confidential information maintained by the Department. 10 person shall also be deemed guilty of a misdemeanor, and upon 11 conviction thereof shall be punished by a fine of Two Hundred 12 Dollars (\$200.00) or imprisonment in the county jail for not more 13 than thirty (30) days, or by both such fine and imprisonment. 14 SECTION 2. This act shall become effective November 1, 2021. 15 16 58-1-1444 DC 1/11/2021 3:26:58 PM 17 18 19 20 21 22 23 24