1	SENATE FLOOR VERSION
2	February 23, 2015
3	SENATE BILL NO. 9 By: Barrington of the Senate
4	and
5	Denney of the House
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8	An Act relating to genetic counseling; providing definition; permitting health care entities
9	administering prenatal care, postnatal care, or genetic counseling to provide certain information to
10	expectant or new parents; establishing standards for information provided to expectant parents;
11	authorizing the State Department of Health to provide certain information under certain circumstances;
12	requiring information to be culturally and linguistically appropriate; prohibiting certain
13	limitation; providing for codification; and providing an effective date.
14	an effective date.
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16	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:
17	SECTION 1. NEW LAW A new section of law to be codified
18	in the Oklahoma Statutes as Section 1-571 of Title 63, unless there
19	is created a duplication in numbering, reads as follows:
20	A. For the purposes of this section, "Down syndrome" means a
21	chromosomal condition caused by an error in cell division that
22	results in the presence of an extra whole or partial copy of
23	chromosome twenty-one (21).
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B. Any hospital, physician, health care provider, nurse
midwife, genetic counselor or other entity that renders prenatal
care, postnatal care or genetic counseling, upon receipt of a
positive test result from a test for Down syndrome, may provide the
expectant or new parent with information provided by the State
Department of Health. The information provided shall be:

1. Up-to-date, evidenced-based written information about Down
syndrome that has been reviewed by medical experts and national Down
syndrome organizations. The written information provided shall
include physical, developmental, educational, and psychological
outcomes, life expectancy, clinical course, intellectual and
functional development, and treatment options; and

Contact information regarding programs and support services,
 including information hotlines specific to Down syndrome, resource
 centers or clearinghouses, national and local Down syndrome
 organizations, and other education and support programs.

17 C. The Department may also make such information available to 18 any other person who has received a positive test result from a test 19 for Down syndrome.

20 D. Information provided under this section shall be culturally 21 and linguistically appropriate for women receiving a positive 22 prenatal diagnosis or for the family of a child receiving a 23 postnatal diagnosis of Down syndrome.

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1	E. Nothing in this section shall limit the rights of any person
2	under the Genetic Counseling Licensure Act.
3	SECTION 2. This act shall become effective November 1, 2015.
4	COMMITTEE REPORT BY: COMMITTEE ON HEALTH AND HUMAN SERVICES February 23, 2015 - DO PASS
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