

Sen. Christine Radogno

## Filed: 3/19/2015

|        | 09900SB0987sam001 LRB099 05579 JLK 32990 a   |
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| 1      | AMENDMENT TO SENATE BILL 987   |
| 2      | AMENDMENT NO Amend Senate Bill 987 by replacing  |
| 3      | everything after the enacting clause with the following:   |
| 4<br>5 | "Section 1. Short title. This Act may be known as the Down Syndrome Information and Awareness Act. |
| 6      | Section 5. Definitions. As used in this Act:   |
| 7      | "Clearinghouse" means a central institution or agency for  |
| 8      | the collection, maintenance, and distribution of materials   |
| 9      | related to Down syndrome.  |
| 10     | "Down syndrome" means a chromosomal condition caused by an   |
| 11     | error in cell division that results in the presence of an extra                                    |
| 12     | whole or partial copy of chromosome 21.  |
| 13     | "First call program" means a volunteer group of individuals  |
| 14     | who make themselves available to parents of children newly   |
| 15     | diagnosed with Down syndrome.  |
| 16     | "Health care provider" means any physician, hospital   |

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1 facility, or other person that is licensed or otherwise
2 authorized to deliver health care services.

3 Section 10. Down syndrome information. The Department 4 available up-to-date, evidence-based written shall make 5 information about Down syndrome that has been reviewed by State and national Down 6 medical experts and syndrome organizations, including physical, developmental, educational 7 8 and psychosocial outcomes, life expectancy, clinical course, 9 intellectual and functional development, and treatment 10 The written information shall include contact options. information regarding first call programs and clearinghouses, 11 12 national, State and local Down syndrome organizations, and 13 other educational and support programs. The Department shall 14 make this information available to persons who render prenatal 15 care, postnatal care, or genetic counseling to parents who receive a prenatal or postnatal diagnosis of Down syndrome. The 16 17 Department shall also make this information available to any person who has received a positive test result from a test for 18 19 Down syndrome. The information provided under this Section 20 shall be culturally and linguistically appropriate for a woman 21 receiving a positive prenatal diagnosis of Down syndrome and 22 for the family of a child receiving a postnatal diagnosis of 23 Down syndrome.

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Section 15. Distribution of information to parents. A

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health care provider who renders prenatal or postnatal care, or genetic counselor who renders genetic counseling shall, upon receipt of a positive test result from a test for Down syndrome, provide the expectant or new parent with the information provided by the Department under Section 10 of this Act.".