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HOUSE BILL 201

**49TH LEGISLATURE - STATE OF NEW MEXICO - SECOND SESSION, 2010**

INTRODUCED BY

Rhonda S. King

AN ACT

RELATING TO PUBLIC HEALTH; AMENDING A SECTION OF THE PUBLIC HEALTH ACT TO ADD FIVE ADDITIONAL DISEASES FOR WHICH TESTING IS REQUIRED FOR NEWBORNS.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

Section 1. Section 24-1-6 NMSA 1978 (being Laws 1973, Chapter 359, Section 6, as amended) is amended to read:

"24-1-6. TESTS REQUIRED FOR NEWBORN INFANTS.--

A. The department shall adopt screening tests for the detection of congenital diseases that shall be given to every newborn infant, except that, after being informed of the reasons for the tests, the parents or guardians of the newborn child may waive the requirements for the tests in writing. The screening tests shall include at a minimum:

- (1) 3-methylcrotonyl-CoA deficiency;

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- 1 (2) 3-OH 3-CH<sub>3</sub> glutaric aciduria;
- 2 (3) argininosuccinic acidemia;
- 3 (4) mitochondrial acetoacetyl-CoA thiolase
- 4 deficiency;
- 5 (5) biotinidase deficiency;
- 6 (6) carnitine uptake defect;
- 7 (7) citrullinemia;
- 8 (8) congenital adrenal hyperplasia;
- 9 (9) congenital hypothyroidism;
- 10 (10) cystic fibrosis;
- 11 (11) galactosemia;
- 12 (12) glutaric acidemia type I;
- 13 (13) Hb S/beta-thalassemia;
- 14 (14) hearing deficiency;
- 15 (15) homocystinuria;
- 16 (16) isovaleric academia;
- 17 (17) long-chain L-3-OH acyl-CoA dehydrogenase
- 18 deficiency;
- 19 (18) maple syrup urine disease;
- 20 (19) medium chain acyl-CoA dehydrogenase
- 21 deficiency;
- 22 (20) methylmalonic acidemia;
- 23 (21) multiple carboxylase deficiency;
- 24 (22) phenylketonuria;
- 25 (23) propanic acidemia;

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- 1 (24) sickle cell anemia;
- 2 (25) trifunctional protein deficiency;
- 3 (26) tyrosinemia type I; [~~and~~]
- 4 (27) very long-chain acyl-CoA dehydrogenase
- 5 deficiency;
- 6 (28) acid maltase deficiency or glycogen
- 7 storage disease type II;
- 8 (29) globoid cell leukodystrophy;
- 9 (30) Gaucher's disease;
- 10 (31) Niemann-Pick disease; and
- 11 (32) Fabry disease.

12 B. In determining which other congenital diseases  
13 to screen for, the secretary shall consider the recommendations  
14 of the New Mexico pediatrics society of the American academy of  
15 pediatrics.

16 C. The department shall institute and carry on such  
17 laboratory services or may contract with another agency or  
18 entity to provide such services as are necessary to detect the  
19 presence of congenital diseases.

20 D. The department shall, as necessary, carry on an  
21 educational program among physicians, hospitals, public health  
22 nurses and the public concerning congenital diseases.

23 E. The department shall require that all hospitals  
24 or institutions having facilities for childbirth perform or  
25 have performed screening tests for congenital diseases on all

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1 newborn infants except if the parents or guardians of a child  
2 object to the tests in writing."

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