§ 26:2-110. Policy declaration

It is hereby declared to be the public policy of this State that in the interests of public health every effort should be made to detect in newborn infants, hypothyroidism, galactosemia, phenylketonuria, and other preventable biochemical disorders which may cause mental retardation or other permanent disabilities and to treat affected individuals.


NOTES:

Cross References:

1. Coverage for treatment of inherited metabolic diseases by hospital service corporation, see 17:48-6s.
2. Coverage for treatment of inherited metabolic diseases by medical service corporation, see 17:48A-7q.
3. Coverage for treatment of inherited metabolic diseases by health service corporation, see 17:48E-35.16.
4. Coverage for treatment of inherited metabolic diseases by individual health insurance policy, see 17B:26-2.10.
5. Coverage for treatment of inherited metabolic diseases by group health insurance policy, see 17B:27-46.1r.
8. Option of additional screening for disorders in infant required; cost, see 26:2-111.1.
9. Coverage for treatment of inherited metabolic diseases by health maintenance organization, see 26:2J-4.17.
10. Coverage for treatment of inherited metabolic diseases by State Health Benefits Program, see 52:14-17.29c.

LexisNexis (R) Notes:
Related Statutes & Rules:

ADMINISTRATIVE CODE:

*N.J.A.C. 8:17-2.2* (2013), CHAPTER EARLY INTERVENTION SYSTEM, Referral process.

*N.J.A.C. 8:18* (2013), CHAPTER NEWBORN BIOCHEMICAL SCREENING PROGRAM, 8, Chapter 18 -- Chapter: Notes.

*N.J.A.C. 8:18-l.1* (2013), CHAPTER NEWBORN BIOCHEMICAL SCREENING PROGRAM, Purpose and scope.

*N.J.A.C. 8:18-l.2* (2013), CHAPTER NEWBORN BIOCHEMICAL SCREENING PROGRAM, Definitions.

*N.J.A.C. 8:18-l.4* (2013), CHAPTER NEWBORN BIOCHEMICAL SCREENING PROGRAM, Responsibilities of the chief executive officer.