

## AMENDMENTS

2008—Subsec. (b). Pub. L. 110-237 substituted “2009, \$5,062,500 for fiscal year 2010, \$5,125,000 for fiscal year 2011, \$5,187,500 for fiscal year 2012, and \$5,250,000 for fiscal year 2013.” for “2008, \$5,062,500 for fiscal year 2009, \$5,125,000 for fiscal year 2010, \$5,187,500 for fiscal year 2011, and \$5,250,000 for fiscal year 2012.”

**§ 300b-13. Interagency Coordinating Committee on Newborn and Child Screening**

**(a) Purpose**

It is the purpose of this section to—

(1) assess existing activities and infrastructure, including activities on birth defects and developmental disabilities authorized under section 247b-4 of this title, in order to make recommendations for programs to collect, analyze, and make available data on the heritable disorders recommended by the Advisory Committee on Heritable Disorders in Newborns and Children under section 300b-10 of this title, including data on the incidence and prevalence of, as well as poor health outcomes resulting from, such disorders; and

(2) make recommendations for the establishment of regional centers for the conduct of applied epidemiological research on effective interventions to promote the prevention of poor health outcomes resulting from such disorders as well as providing information and education to the public on such effective interventions.

**(b) Establishment**

The Secretary shall establish an Interagency Coordinating Committee on Newborn and Child Screening (referred to in this section as the “Interagency Coordinating Committee”) to carry out the purpose of this section.

**(c) Composition**

The Interagency Coordinating Committee shall be composed of the Director of the Centers for Disease Control and Prevention, the Administrator, the Director of the Agency for Healthcare Research and Quality, and the Director of the National Institutes of Health, or their designees.

**(d) Activities**

The Interagency Coordinating Committee shall—

(1) report to the Secretary and the appropriate committees of Congress on its recommendations related to the purpose described in subsection (a); and

(2) carry out other activities determined appropriate by the Secretary.

**(e) Authorization of appropriations**

For the purpose of carrying out this section, there are authorized to be appropriated \$1,000,000 for fiscal year 2009, \$1,012,500 for fiscal year 2010, \$1,025,000 for fiscal year 2011, \$1,037,500 for fiscal year 2012, and \$1,050,000 for fiscal year 2013.

(July 1, 1944, ch. 373, title XI, §1114, as added Pub. L. 110-204, §6, Apr. 24, 2008, 122 Stat. 710; amended Pub. L. 110-237, §1(a)(6), May 27, 2008, 122 Stat. 1557.)

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2008—Subsec. (e). Pub. L. 110-237 substituted “2009, \$1,012,500 for fiscal year 2010, \$1,025,000 for fiscal year

2011, \$1,037,500 for fiscal year 2012, and \$1,050,000 for fiscal year 2013.” for “2008, \$1,012,500 for fiscal year 2009, \$1,025,000 for fiscal year 2010, \$1,037,500 for fiscal year 2011, and \$1,050,000 for fiscal year 2012.”

**§ 300b-14. National contingency plan for newborn screening**

**(a) In general**

Not later than 180 days after April 24, 2008, the Secretary, acting through the Director of the Centers for Disease Control and Prevention and in consultation with the Administrator and State departments of health (or related agencies), shall develop a national contingency plan for newborn screening for use by a State, region, or consortia<sup>1</sup> of States in the event of a public health emergency.

**(b) Contents**

The contingency plan developed under subsection (a) shall include a plan for—

(1) the collection and transport of specimens;

(2) the shipment of specimens to State newborn screening laboratories;

(3) the processing of specimens;

(4) the reporting of screening results to physicians and families;

(5) the diagnostic confirmation of positive screening results;

(6) ensuring the availability of treatment and management resources;

(7) educating families about newborn screening; and

(8) carrying out other activities determined appropriate by the Secretary.

(July 1, 1944, ch. 373, title XI, §1115, as added Pub. L. 110-204, §7, Apr. 24, 2008, 122 Stat. 711.)

**§ 300b-15. Hunter Kelly Research Program**

**(a) Newborn screening activities**

**(1) In general**

The Secretary, in conjunction with the Director of the National Institutes of Health and taking into consideration the recommendations of the Advisory Committee, may continue carrying out, coordinating, and expanding research in newborn screening (to be known as “Hunter Kelly Newborn Screening Research Program”) including—

(A) identifying, developing, and testing the most promising new screening technologies, in order to improve already existing screening tests, increase the specificity of newborn screening, and expand the number of conditions for which screening tests are available;

(B) experimental treatments and disease management strategies for additional newborn conditions, and other genetic, metabolic, hormonal, or functional conditions that can be detected through newborn screening for which treatment is not yet available; and

(C) other activities that would improve newborn screening, as identified by the Director.

**(2) Additional newborn condition**

For purposes of this subsection, the term “additional newborn condition” means any

<sup>1</sup> So in original. Probably should be “consortium”.