TO BE RESCINDED

3701-45-02 **Required screening: facility requirements.**

(A) All newborn children shall be screened in accordance with the procedures set forth in Chapter 3701-45 of the Administrative Code for the presence of the following genetic, endocrine, or metabolic disorders:

(1) Phenylketonuria;

(2) Homocystinuria;

(3) Galactosemia;

(4) Hypothyroidism;

(5) Sickle cell and other hemoglobinopathies;

(6) Medium chain acyl-coA dehydrogenase deficiency;

(7) Maple syrup urine disease;

(8) Propionic Aciduria;

(9) Isovaleric Aciduria;

(10) Methylmalonic Aciduria;

(11) Citrullinemia; and

(12) Argininosuccinic Aciduria.

(B) All hospitals and freestanding birthing centers that are required by this chapter to cause specimens to be collected for newborn screening for genetic, endocrine, or metabolic disorders shall:

(1) Designate a newborn screening coordinator and physician responsible for the coordination of the facility's newborn screening;

(2) Notify the chief of the Ohio department of health bureau of public laboratories of the name of the individual designated as the newborn screening

coordinator on a yearly basis and whenever the designated individual changes; and

(3) Develop a written protocol for tracking newborn screening. The protocol must include a requirement that the name of the physician attending the child after birth or a designee be placed on the specimen slip sent with the initial specimen to the Ohio department of health public health laboratory.

Effective:

R.C. 119.032 review dates: 01/17/2003

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Certification

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Date

Promulgated Under: 119.03
Statutory Authority: 3701.501
Rule Amplifies: 3701.501
Prior Effective Dates: 1/31/02, 1/13/01, 2/15/99