

NEWBORN SCREENING PROGRAM
New York State Department of Health
Wadsworth Center, David Axelrod Institute
120 New Scotland Avenue
Albany, NY 12208
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INHERITED METABOLIC DISORDER DIAGNOSIS FORM

Dear Doctor:

Please complete this form in its entirety and return it to the Newborn Screening Program as soon as possible.

Attach Clinical Laboratory results including any available mutation analysis.

Your response is required, as specified in Title 10 New York Code of Rules and Regulations subpart 69-1.5e.

NEWBORN INFORMATION:

Name at Time of Birth: _____

Other Names (AKA): _____

Single Birth Twin A Twin B Other _____

Mother's Name: _____

Date of Birth: _____

Gender: Male Female

Hospital of Birth: _____

Medical Record #: _____

Galactosemia

- GALT01 Expired. If cause of death is known, choose the appropriate diagnosis below.
- GALT10 Disease, Galactosemia ó classical
- GALT11 Disease, Galactosemia ó variant
- GALT29 Disease, not on NBS panel. Specify: _____
- GALT30 Inconclusive, Galactosemia
- GALT40 No disease
- GALT41 No disease, transient abnormality due to prematurity/TPN
- GALT49 No disease, polymorphisms only
- GALT71 Other, maternal disease or medication

Biotinidase

- BIOT01 Expired. If cause of death is known, choose the appropriate diagnosis below.
- BIOT10 Disease Biotinidase ó classical
- BIOT11 Disease ó partial Biotinidase deficiency
- BIOT29 Disease, not on NBS panel. Specify: _____
- BIOT30 Inconclusive, Biotinidase
- BIOT40 No disease
- BIOT41 No disease, transient abnormality due to prematurity/TPN
- BIOT71 Other, maternal disease or medication

COMMENTS: _____

PHYSICIAN'S SIGNATURE: _____ **DATE:** _____

PRINT NAME: _____