

Newborn Screening Report Form

Infant's Name: _____ Infant's Birth Date ___ / ___ / ___

Newborn Screening Program Lab #: _____ Mother's Name: _____

Diagnosis pending, Follow-up Plan:

Final Diagnosis (please attach CONFIRMATION LAB RESULTS)

- Normal
- Trait Condition (specify carrier status) _____
- Classic Galactosemia (GG phenotype/genotype)
- Duarte/Galactosemia Compound Heterozygote (DG phenotype/genotype)
- Congenital Hypothyroidism
- Congenital Adrenal Hyperplasia due to 21-Hydroxylase Deficiency
- Cystic Fibrosis
- Amino Acid Disorder (specify type) _____
- Fatty Acid Oxidation Disorder (specify type) _____
- Organic Acid Disorder (specify type) _____
- Mucopolysaccharidosis Type I (specify type) _____
- Pompe (specify type) _____
- Sickle Cell Disease (specify type) _____
- Hemoglobin disease (specify type) _____
- Biotinidase Deficiency (specify type) _____
- Other (specify) _____

Treatment Indicated? yes no

Date treatment started ___ / ___ / ___

Referred to pediatric sub-specialist (specify name):

Endocrinologist: _____

Metabolic Specialist/Geneticist: _____

Neurologist: _____

Date of first clinic visit to specialist: ___ / ___ / ___

Hematologist: _____

Pulmonologist: _____

Family referred for (check all that apply):

Genetic counseling (check provider):

___ Geneticist ___ Other (specify) _____

Enrollment in Newborn Screening Long-term Follow-up Program

Early Intervention Services

Print Physician's Name _____ Telephone ___ / ___ / ___

Physician Signature _____ Date ___ / ___ / ___

Mail or Fax this follow-up form with complete diagnostic information and confirmation lab results to:
Oklahoma State Department of Health
ATTN: Newborn Screening Program Coordinator
1000 NE Tenth Street
Oklahoma City, OK 73117-1299
Fax: (405) 271-4892

Questions or Referral Information
Phone: (405) 271-6617
(800) 766-2223