

Definitions of Medical Intervention and Diagnosis, by Core RUSP Conditions

The purpose of this document is to serve as a guide for defining notification of the newborn screening results, medical intervention, and the date of diagnosis. Each definition correlates to [Quality Indicator](#) metrics and aims to provide standardization for Newborn Screening programs.

| Amino Acid Disorders | Notification of out-of-range NBS result ⁱ | Medical Intervention ⁱⁱ | Date of Diagnosis |
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| Argininosuccinic Aciduria (ASA) | <p>The notification of an out-of-range result, such as Citrulline or Argininosuccinate acid, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of ASA.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Diet change, drug, or supplement therapy. • Collection or testing for plasma ammonia and amino acids, urine amino acids, and argininosuccinate lyase enzyme. • Investigating poor feeding, vomiting, lethargy, tachypnea, or seizures. | <p>The date the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Amino acid and enzyme profile results +/- genotyping/sequencing of the ASL gene. |
| Citrullinemia, Type I (CIT) | <p>The notification of an out-of-range result, such as Citrulline or Argininosuccinate acid, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of CIT.</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Amino acid and enzyme profile results +/- genotyping/sequencing of |

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| | <p>Time-critical disorder.</p> | <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Diet change, drug, or supplement therapy. • Collection or testing for plasma ammonia and amino acids, urine amino acids, and arginosuccinate lyase enzyme. • Investigating poor feeding, vomiting, lethargy, tachypnea, or seizures. | <p>the ASS1 gene and SLC25A13 gene.</p> |
| <p>Classic Phenylketonuria (PKU)</p> | <p>The notification of an out-of-range result, such as Phenylalanine and/or Tyrosine, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of PKU.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Diet change, drug, or supplement therapy. • Collection or testing for plasma amino acids, urine pterins, or DHPR activity. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Amino acid and enzyme profile results +/- genotyping/sequencing of the PAH gene. |

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| | | <i>Note-</i> Newborns with PKU are asymptomatic. | |
| Homocystinuria (HCY) | The notification of an out-of-range result, such as Methionine or Homocysteine, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of HCY.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Diet change, drug, or supplement therapy. • Collection or testing plasma and urine amino acids. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Amino acids and total homocysteine profile results + genotyping/sequencing of the CBS gene. |
| Maple Syrup Urine Disease (MSUD) | <p>The notification of an out-of-range result, such as Leucine, Isoleucine, and Valine, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MSUD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Diet change, drug, or supplement therapy. • Collection or testing for plasma amino acids and urine organic acid analysis. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Plasma amino acids and urine organic acid profile results +/- genotyping/sequencing of the BCKDHA, BCKDHB, or DBT genes. |

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| | | <ul style="list-style-type: none"> Investigating poor feeding, lethargy, tachypnea seizures, or other symptoms. | |
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| Tyrosinemia, Type I (TYR I) | The notification of an out-of-range result, such as SUAC and Tyrosine, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of TYR I.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> Diet change or drug therapy. Collection or testing for liver function tests, Alpha-fetoprotein, electrolytes, plasma amino acids, or urine organic acids. Investigating failure to thrive, jaundice, diarrhea, or vomiting. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> Plasma amino acids and urine organic acid profile results +/- genotyping/sequencing of the FAH gene. |

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| Endocrine Disorders | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| Congenital Adrenal Hyperplasia (CAH) | <p>The notifications of an out-of-range result, such as 17-OHP or steroid profile, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or endocrine consultation to evaluate the potential diagnosis of CAH.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of oral glucocorticoid replacement therapy, or IV fluids, IM/IV hydrocortisone. • Collection or testing of serum 17-OHP, ACTH, urinary steroid profile, electrolyte panel, and plasma renin activity (PRA). • Investigating poor feeding, vomiting, lethargy, ambiguous genitalia, or non-palpable testes. | <p>The date of the clinical report reflects a confirmed diagnosis, with or without other abnormal adrenal hormone abnormalities, and evaluation by an endocrinologist.</p> <ul style="list-style-type: none"> • Serum and urine results + genotyping/sequencing of the CYP21A2 gene |

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| <p style="text-align: center;">Primary Congenital Hypothyroidism (CH)</p> | <p>The notifications of an out-of-range result, such as TSH, Total, and Free T4, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or endocrine consultation to evaluate the potential diagnosis of CH.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Serology evaluation of TSH, Free T4, and Total T4. • Initiates therapy like levothyroxine. • Investigating poor growth, prolonged jaundice, puffy facies, large fontanelles, macroglossia, or umbilical hernia. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Serology testing +/- imaging, |
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| Fatty Acid Oxidation Disorder | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| Carnitine Uptake Defect/Carnitine Transport Defect (CUD) | The notification of an out-of-range result, such as C0 and other acylcarnitines, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of CUD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Carnitine therapy • Collection or testing of plasma and urine carnitine. • Investigating poor feeding, lethargy, tachypnea, hepatomegaly, or hypotonia. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Acylcarnitine profile results +/- genotyping/sequencing of the SLC22A5 gene. |
| Long-chain L-3 Hydroxyacyl- CoA Dehydrogenase Deficiency (LCHAD) | <p>The notification of an out-of-range result, such as C16-OH and other long-chain acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of LCHAD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Acylcarnitine and enzyme profile results +/- genotyping/sequencing of the HADHA gene. |

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| | | <ul style="list-style-type: none"> • Advisement to avoid fasting or initiation of IV glucose. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, lethargy, vomiting, hypotonia, hepatomegaly, hypoglycemia, arrhythmias, or cardiac insufficiency. | |
| <p>Medium-chain Acyl-CoA Dehydrogenase Deficiency (MCAD)</p> | <p>The notification of an out-of-range result, such as C8, with or without C6 and C10, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MCAD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Advisement to avoid fasting or initiation of IV glucose. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, vomiting, lethargy, hypotonia, or hypoglycemia. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Acylcarnitine and enzyme profile results +/- genotyping/sequencing of the ACADM gene. |

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| <p style="text-align: center;">Trifunctional Protein Deficiency (TFP)</p> | <p>The notification of an out-of-range result, such as C16-OH and other long-chain acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of TFP.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Advisement to avoid fasting or initiation of IV glucose. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, lethargy, vomiting, hypotonia, hepatomegaly, hypoglycemia, arrhythmias, or cardiac insufficiency. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Acylcarnitine and enzyme profile results +/- genotyping/sequencing of the HADHA and HADHB genes. |
| <p style="text-align: center;">Very Long-chain Acyl-CoA Dehydrogenase Deficiency (VLCAD)</p> | <p>The notification of an out-of-range result, such as C14, C14:1, and other long-chain acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of VLCAD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Acylcarnitine enzyme profile results +/- genotyping/sequencing of the ACADVL gene. |

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| | | <ul style="list-style-type: none">• Advisement to avoid fasting or initiation of IV glucose.• Collection or testing of plasma acylcarnitines,• Investigating poor feeding, vomiting, lethargy, hypotonia, hepatomegaly, arrhythmias, or cardiac complications. | |
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| Hemoglobinopathies | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| <p>Presence of Hb S [S, Beta-Thalassemia, S,C Disease, S,S Disease (Sickle Cell Anemia)]</p> | <p>The notification of an out-of-range result by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or hematologist consultation to evaluate the possible diagnosis of Sickle Cell Anemia (FS), SC disease (FSC), Sickle Beta + Thalassemia (FSA). Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • The collection or testing of a complete blood count (CBC) with or without a manual smear and a hemoglobin electrophoresis analysis. • Investigating splenomegaly or jaundice. • Decision to initiate antibiotics, or not to initiate antibiotics. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Confirmation testing by an alternative method (e.g., secondary electrophoresis) +/- DNA studies. <ul style="list-style-type: none"> ○ Hemoglobin S |

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| Lysosomal Storage Disorders | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| <p>Glycogen Storage Disease Type II (Pompe)</p> <p><i>New disorder</i></p> | <p>The notification of an out-of-range result, such as GAA, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the clinic or hospital consultation to evaluate the potential diagnosis of Pompe.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Chest X-ray, EKG, ECHO, or other examination of cardiac status. • Collection or testing for urine glucose tetrasaccharide (HEX4), muscle enzymes CK, LDH, AST, ALT). | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Alpha-glucosidase enzyme analysis results + genotyping/sequencing of the GAA gene. |
| <p>Mucopolysaccharidosis Type I (MPS I)</p> <p><i>New disorder</i></p> | <p>The notification of an out-of-range result, such as IDUA enzyme or GAGs, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic or hospital consultation to evaluate the potential diagnosis of MPS I.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Urine testing for glycosaminoglycans analysis (GAGs) like dermatan sulfate, heparan sulfate, keratan sulfate, and chondroitin 6-sulfate. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Alpha-L-iduronidase enzyme analysis results, genotyping/sequencing of the IDUA gene, and GAG results. |

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| <p>Mucopolysaccharidosis Type II (MPS II)</p> <p><i>New disorder</i></p> <p>Draft- Public health surveillance algorithm in development.</p> | <p>The notifications of an out-of-range result, such as I2S enzyme or GAGs, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic or hospital consultation to evaluate the potential diagnosis of MPS II.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Urine testing for glycosaminoglycans analysis (GAGs) like dermatan sulfate, heparan sulfate, keratan sulfate, and chondroitin 6-sulfate. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Iduronate-2-Sulfatase enzyme analysis results, genotyping/sequencing of the IDS molecular, and GAG results. |
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| Organic Acid Disorders | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| <p>3-Hydroxy-3-Methylglutaric Aciduria (HMG)</p> <p><i>Draft- Public health surveillance algorithm in development.</i></p> | <p>The notification of an out-of-range result, such as C5-OH, C6-DC, and other acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of HMG.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, vomiting, lethargy, hypoglycemia, ketonuria, or metabolic acidosis. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Urine organic acid and plasma acylcarnitine profile results +/- genotyping/sequencing of the HMGCL gene. |
| <p>3-Methylcrotonyl-CoA Carboxylase Deficiency (3-MCC)</p> | <p>The notification of an out-of-range result, such as C5-OH and other acycarnitines, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MCC.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Urine organic acids, plasma amino acids, and plasma acylcarnitine profile results +/- genotyping/sequencing of the MCCC1 and MCCC2 gene. |

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| | | <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, vomiting, lethargy, hypoglycemia, ketonuria, or metabolic acidosis. | |
| <p>β-Ketothiolase Deficiency (BKT)</p> <p><i>Draft- Public health surveillance algorithm in development.</i></p> | <p>The notification of the out-of-range result, such as C5:1, C5-OH, and other acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of BKT:</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, vomiting, lethargy, hypoglycemia, ketonuria, or metabolic acidosis. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Urine organic acids, plasma amino acids, and plasma acylcarnitine profile results +/- genotyping/sequencing of the ACAT1 gene. |
| <p>Glutaric Acidemia Type I (GA1)</p> | <p>The notification of the out-of-range result, such as C5-DC and other acylcarnitines, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of GA 1.</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Urine and plasma organic acids and acylcarnitine profile results +/- |

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| | <p>Time-critical disorder.</p> | <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of urine and plasma organic acids and acylcarnitines. • Investigating poor feeding, macrocephaly, or hypotonia. | <p>genotyping/sequencing of the GCDH gene.</p> |
| <p style="text-align: center;">Holocarboxylase Synthetase Deficiency (MCD)</p> | <p>The notification of the out-of-range result, C3, C5-OH, and other acylcarnitines, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MCD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of plasma acylcarnitines and urine organic acids. • Investigating poor feeding, vomiting, lethargy, hypoglycemia, ketonuria, or metabolic acidosis. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Urine organic acids, plasma amino acids, and plasma acylcarnitine profile results +/- genotyping/sequencing of the HLCS gene. |

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| <p>Isovaleric Acidemia (IVA)</p> | <p>The notification of the out-of-range result, C5 and other acylcarnities, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of IVA</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change, drug, or supplement therapy. • Collection or testing of plasma and urine acylcarnitines, and urine organic acids. • Investigating poor feeding, vomiting, lethargy, tachypnea, or odor of sweaty feet. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Plasma acylcarnitines, and urine acylcarnitines and organic acids profile results +/- Genotyping/sequencing of the IVD gene. |
| <p>Methylmalonic Acidemia Cbl A, B (cobalamin disorders) (MMA)</p> | <p>The notification of the out-of-range result, such as C3, C3/C2, and other acylcarnities, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MMA.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet, drug, or supplement therapy. • Collection or testing of plasma and urine organic | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Plasma and urine organic acid profile results + genotyping/sequencing of the MMAA and MMAB gene. |

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| | | <p>acids, and vitamin B12 analysis.</p> <ul style="list-style-type: none"> Investigating poor feeding, vomiting, lethargy, tachypnea, ketonuria, hypoglycemia, or failure to thrive. | |
| <p>Methylmalonic Acidemia (methylmalonyl-CoA mutase) (MUT)</p> | <p>The notification of the out-of-range, C3, C3/C2, and other acylcarnities, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of MUT.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> Date of diet, drug, or supplement therapy. Collection or testing of plasma and urine organic acids, and vitamin B12 analysis. Investigating poor feeding, vomiting, lethargy, tachypnea, ketonuria, hypoglycemia, or failure to thrive. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> Plasma and urine organic acid profile results + genotyping/sequencing of the MMUT gene. |
| <p>Propionic Acidemia (PROP)</p> | <p>The notification of the out-of-range result, C3, C3/C2, and other acylcarnities, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of PROP.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> Plasma and urine organic acid profile results +/- genotyping/sequencing of the PCCA and PCCB. |

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| | | <p>before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none">• Date of diet, drug, or supplement therapy.• Collection or testing of plasma and urine organic acids, and vitamin B12 analysis.• Investigating poor feeding, vomiting, lethargy, tachypnea, ketonuria, hypoglycemia, or failure to thrive. | |
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| Other Disorders | Notification of out-of-range NBS result | Medical Intervention | Date of Diagnosis |
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| Biotinidase Deficiency (BIOT) | The notification of an out-of-range result, such as Biotinidase, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic, hospital, or metabolic specialist consultation to evaluate the potential diagnosis of BIOT.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Biotin supplements. • Collection or testing of serum Biotinidase assay. • Investigation of seizures, hypotonia, ataxia, skin rash, etc. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Serum biotinidase and urine organic acids, +/- genotyping/sequencing of the BTD gene. |
| Classic Galactosemia (GALT) | <p>The notification of an out-of-range result, such as Galactose-1-phosphate and Total Galactose, by the NBS Program to the appropriate medical professional.</p> <p>Time-critical disorder.</p> | <p>Date of the first clinic, hospital, or metabolic consultation with a metabolic specialist to evaluate the potential diagnosis of GALT.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Date of diet change. • Urine reducing substances. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Red blood cell GALT activity and/or galactose-1-phosphate (gal-1-P), +/- GALT mutation analysis. |
| Cystic Fibrosis (CF) | The notifications of an out-of-range result, such as immunoreactive trypsinogen | Date of the first clinic or hospital consultation to evaluate the | The date of the clinical report reflects a confirmed diagnosis: |

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| | (IRT), by the NBS Program to the appropriate medical professional. | <p>potential diagnosis of CF or initiate therapy.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening result is reported and may include any of the following:</p> <ul style="list-style-type: none"> • Pancreatic enzymes/Salt supplements. • Antibiotics or other medication to support or manage symptoms. • Appointment for sweat chloride testing | <ul style="list-style-type: none"> • Positive sweat chloride test. • Genotype/sequencing identifying variants in the CFTR gene. • Nasal Potential Difference (NPD) results. |
| <p>Guanidinoacetate Methyltransferase Deficiency (GAMT)</p> <p><i>New disorder</i></p> <p>Draft- Public health surveillance algorithm in development.</p> | The notification of an out-of-range result, such as GUAC and Creatine, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic or hospital consultation to evaluate the potential diagnosis of GAMT.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Neurological investigation for hypotonia, epilepsy, etc. • Urine/plasma testing for creatine. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Plasma and urine GUAC and creatine analysis, + genotyping/sequencing of the GAMT gene. • Rule out of arginase deficiency. |
| <p>Severe Combined Immunodeficiencies (SCID)</p> | The notification of an out-of-range result, such as TREC, by the NBS Program to the appropriate medical professional. | <p>Date of the first clinic or hospital consultation with an immunologist or infectious disease specialists to evaluate the potential diagnosis for SCID:</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Flow cytometry and CD 4 results + genotyping/sequencing for IL2RG or other gene |

Definitions of Medical Intervention and Diagnosis, by Core RUSP Conditions

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| | | <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • Includes testing (but not limited to) CBC with differential and Flow Cytometry. • Maternal T-cell engraftment • Proliferation of PHA • Antibiotic therapy or isolation protocol. | <p>mutations known to cause SCID.</p> |
| <p>Spinal Muscular Atrophy (SMA)</p> | <p>The notification of an out-of-range result, such as SMN1 and SMN2, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic or hospital consultation to evaluate the potential diagnosis for SMA.</p> <p>Medical intervention should be the first date that alters the child's care. This may include any of the following:</p> <ul style="list-style-type: none"> • The neuromuscular investigation includes but is not limited to, hypotonia, weakness, poor feeding, or respiratory difficulties. | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • SMN1 sequencing/ SMN1 and SMN2 dosage analysis. |
| <p>X-linked Adrenoleukodystrophy (X-ALD)</p> <p><i>New disorder</i></p> | <p>The notification of an out-of-range result, such as C24:0 and C26:0, by the NBS Program to the appropriate medical professional.</p> | <p>Date of the first clinic or hospital consultation to evaluate the potential diagnosis for X-ALD.</p> <p>Medical intervention should be the first date that alters the child's care. This date may be before the newborn screening</p> | <p>The date of the clinical report reflects a confirmed diagnosis:</p> <ul style="list-style-type: none"> • Serum fatty acid profile and imaging + genotyping/sequencing ABCD1 gene. |

Definitions of Medical Intervention and Diagnosis, by Core RUSP Conditions

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| | | result is reported and may include any of the following: <ul style="list-style-type: none">• Adrenal insufficiency investigation.• Adrenal steroid replacement.• Specimen collection for very long chain fatty acids (VLCFA)/Plasmalogens. | |
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ⁱ The notification of an out-of-range result(s) may be communicated by phone, fax, mailer, or other method deemed acceptable by the program's policy.

ⁱⁱ Medical intervention is any interaction between a medical professional and the newborn/infant's family that alters the care of the child based on out-of-range newborn screening results. The intervention will be the earliest date that clinical action was rendered. Medical intervention includes a medical professional advising against fasting or the date therapy was initiated. This may also include the date additional samples were collected for confirmation testing. See the [Quality Indicator Source Document](#) for further explanation.