Time Critical Disorders:

The following table is from the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children's (ACHDNC) recommendations on timeliness in newborn screening and was created based on the Society for Inherited Metabolic Disorders (SIMD) position statement and expert opinion from metabolic geneticists, hematologists, endocrinologist and pulmonologists.

Organic Acid Conditions	Fatty Acid Oxidation Disorders	Amino Acid Disorders	Other Disorders
Propionic acidemia (PROP)	Medium chain acyl-CoA-dehydrogenase deficiency (MCAD)	Argininosuccinic aciduria (ASA)	Classic Galactosemia (GALT)
Methylmalonic acidemia (methylmalonyl-CoA mutase) (MUT)	Very Long chain acyl-CoA dehydrogenase deficiency (VLCAD)	Citrullinemia type-1 (CIT)	Congenital adrenal hyperplasia (CAH)
Isovaleric acidemia (IVA)	Long chain L-3-hydroxyacyl-CoA dehydrogenase deficiency (LCHAD)	Maple syrup urine disease (MSUD)	
3-Hydroxy-3-methyglutaric aciduria (HMG)	Trifunctional protein deficiency (TFP)		
Holocarboxylase synthase deficiency (MCD)			
β-Ketothiolase deficiency (BKT)			
Glutaric Aciduria, Type 1 (GA1)			