

New Jersey Department of Health and Senior Services
Division of Family Health Services
Newborn Screening and Genetic Services Program

This Table represents the outcome of Newborn Screening test results for the 102,666 initial screens received during Calendar Year 2010.

2010 Data as of 8/20/2012 [Date of Birth 1/1/2010 – 12/31/2010]		# of Babies with Confirmed Classic Disease	# of Babies with Variant Disease or Carrier Status	# of Babies with cleared results
Newborn Screening Disorders				
Biotinidase Deficiency	BIOT	2	9	52
Congenital Adrenal Hyperplasia	CAH	2	3	653
Congenital Hypothyroidism	CH	57	13	1821
Cystic Fibrosis	CF	10	28	161
Galactosemia	GALT	2	41	50
Maple Syrup Urine Disease	MSUD	0	0	0
Phenylketonuria	PKU	2	5	2
Sickle Cell Anemia and Other Hemoglobinopathies	S/S, S/C, Var Hb	43	38	7
Hemoglobin Traits	2931			
<i>Amino Acid Disorders</i>				234
Homocystinuria	HCY	0	0	
Hypermethioninemia	MET	0	1	
Tyrosinemia	TYR	0	6	
<i>Fatty Acid Disorders</i>				99
Carnitine Uptake Defect	CUD	1	1	
Short Chain Acyl-CoA Dehydrogenase Deficiency	SCAD	8	1	
Glutaric Aciduria, Type II	GA-II	0	0	
Medium Chain Acyl-CoA Dehydrogenase Deficiency	MCAD	5	0	
Long/Very Long Chain Acyl-CoA Dehydrogenase Deficiency	LCAD/VLCAD	2	0	
Long Chain 3-Hydroxyacyl-CoA Dehydrogenase Deficiency	LCHAD	0	0	
Trifunctional Protein Deficiency	TFP	1	0	
Carnitine Palmitoyltransferase Deficiency, Type II	CPT-II	0	0	
Carnitine/Acylcarnitine Translocase Deficiency	CACT	0	0	
Carnitine Palmitoyltransferase Deficiency, Type IA	CPT-1A	0	0	
Medium/Short Chain 3-OH Acyl-CoA Dehydrogenase Deficiency	M/SCHAD	0	0	
Medium Chain Ketoacyl-CoA Thiolase Deficiency	MCKAT	0	0	
Dienoyl-CoA Reductase Deficiency	DERED	0	0	
<i>Organic Acid Disorders</i>				178
Propionyl-CoA Carboxylase Deficiency	PROP	1	0	
Methylmalonic Acidemia [Mutase or Cobalamin Defects]	MUT/CBL	0	3	
Isobutyryl-CoA Dehydrogenase Deficiency	IBD	0	0	
Isovaleryl-CoA Dehydrogenase Deficiency	IVA	0	0	
2-Methylbutyryl-CoA Dehydrogenase Deficiency	2MBG	0	0	
3-Hydroxy-3-Methylglutaryl-CoA Lyase Deficiency	HMG	2	0	
3-Methylcrotonyl-CoA Carboxylase Deficiency	3MCC	3	0	
Multiple Carboxylase Deficiency	MCD	0	0	
3-Methylglutaconyl CoA Hydrastase Deficiency	3MGA	0	0	
Glutaric Aciduria, Type I	GA-1	2	0	
Mitochondrial Acetoacyl CoA Thiolase Deficiency	BKT	0	0	
2-Methyl-3-Hydroxybutyric Acidemia	2M3HBA	0	0	
Malonyl-CoA Decarboxylase Deficiency	MAL	0	0	
<i>Urea Cycle Disorders</i>				7
Citrullinemia I + II	CIT	0	0	
Argininosuccinate Lyase Deficiency	ASA	1	0	
Argininemia	ARG	0	0	
TOTALS		144	149	3264