Disorders	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014
2-Methylbutyryl-CoA Dehydrogenase Deficiency	*	*	0	1	0	0	0	1	0	1
3-Methylcrotonyl-CoA Carboxylase Deficiency (3-MCC)	*	*	0	0	1	0	0	2	0	0
Biotindinase Deficiency	0	0	1	0	1	0	0	0	0	0
Carnitine Transporter Defect	0	0	1	2	3	0	2	1	1	1
Citrullinemia	1	0	0	0	0	0	0	0	0	0
Congenital Adrenal Hyperplasia	4	1	1	7	5	3	5	4	2	5
Congenital Hypothyroidism	25	23	18	16	30	33	29	32	36	33
Critical Congenital Heart Disease	*	*	*	*	*	*	*	3	2	2
Cystic Fibrosis	*	*	*	7	23	16	19	14	12	19
Galactosemia	1	1	2	0	0	4	0	3	1	2
Glutaric Acidemia	*	0	1	1	0	0	1	0	0	0
Hearing Loss		47	55	43	29	31	68	60	54	Pending
Homocystinuria	0	0	1	1	0	0	0	0	0	0
Isovaleric Acidemia	*	0	0	0	0	0	0	0	0	0
Long chain Acyl-CoA Dehydrogenase Deficiency (LCAD)	*	*	0	0	0	0	0	0	1	0
Maple Syrup Urine Disease	0	0	0	0	0	0	0	0	1	0
Medium chain Acyl-CoA Dehydrogenase Deficiency (MCAD)	1	5	3	3	6	5	5	4	3	3
Methylmalonic Acidemia (MMA)	1	0	0	2	2	0	1	1	1	1
Phenylketonuria	2	5	6	3	4	3	5	2	4	4
Propionic Acidemia	0	0	0	0	0	1	0	0	0	1
Secondary Congenital Heart Disease	*	*	*	*	*	*	*	5	3	1
Sickle Cell Disease	34	56	51	57	62	56	66	52	53	58
Sickle Cell Trait	1622	1754	2038	1818	1940	1860	1835	1866	1817	1468
Trifunctional Protein Deficiency	*	*	0	0	1	0	0	0	0	0
Tyrosinemia	-	-	-	-	-	1	0	0	0	0
Very long chain Acyl-CoA Dehydrogenase Deficiency (VLCAD)	*	*	1	0	0	0	1	1	1	2