Newborn Screening Program - 2015 Annual Report
Jan 1-Dec 31 2015
New York State Department of Health
Wadsworth Center
Biggs Laboratory
Albany, NY

Specimens Received	
Initial Valid	226,900
Initial Invalid	10,601
Total Newborns	237,501
Repeat Specimens	28,036
Total Specimens	265,537

Screened Disorders	Analytes	Referrals	# Confirmed with Disease	
Endocrine Disorders				
Congenital Adrenal Hyperplasia	17-hydroxyprogesterone	257	Congenital adrenal hyperplasia - 21-Hydroxylase deficiency	15
			Congenital adrenal hyperplasia, other enzyme deficiency	1
Congenital Hypothyroidism	Thyroxine	756	Primary Congenital Hypothyroidism	91
			Secondary Congenital Hypothyroidism	8
			Other	291
Hemoglobin Disorders				
Hemoglobin Disorders	Hemoglobin SS	134	Hemoglobin S + S (sickle cell) disease	127
	Hemoglobin SC	76	Hemoglobin S + C disease	76
	Hemoglobin CC	33	Hemoglobin C + C disease	26
	Other Hemoglobins	66	Other Hemoglobinopathies	51
Infectious Disease				
HIV	HIV-1 Antibodies	401	Confirmed by diagnosis developed by the AIDS Institute	
Amino Acid Disorders				
Maple Syrup Urine Disease	Leucine	5	Maple Syrup Urine disease	1
			Hydroxyprolinemia	0
Homocystinuria	Methionine	3	Homocystinuria	0
			Hypermethioninemia	0
Phenylketonuria	Phenylalanine	24	Phenylketonuria (PKU)	5
			Hyperphenylalaninemia	17
Tyrosinemia Type I	Succinylacetone	0	Tyrosinemia Type 1	0
Tyrosinemia Type II, III	Tyrosine	8	Tyrosinemia Type 2	0
			Tyrosinemia Type 3	0
Fatty Acid Oxidation Disorders				
Carnitine uptake defect	Free Carnitine (C0), Total Acylcarnitines (SUM AC)	20	Carnitine uptake defect (CUD)	2
Carnitine palmitoyltransferase 1 deficiency	C0/(C16 + C18)	0	Carnitine palmitoyltransferase 1 (CPT1) deficiency	0
Carnitine palmitoyltransferase 2 deficiency/Carnitine/acylcarnitine translocase (CACT) deficiency	Hexadecanoylcarnitine (C16), Octadecenoylcarnitine (C18:1)	10	Carnitine palmitoyltransferase 2 (CPT2) deficiency	1
2,4-Dienoyl-CoA reductase deficiency	Decadienoylcarnitine (C10:2)	0	2,4-Dienoyl-CoA (2,4Di) reductase deficiency	0
Long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency/Trifunctional protein deficiency	Hydroxyhexadecanoylcarnitine (C16OH), Hydroxyoctadecenoylcarnitine (C18:1OH)	3	Long-chain 3-hydroxyacyl-CoA dehydrogenase (LCHAD) deficiency	0
			Trifunctional protein (TFP) deficiency	0
Multiple acyl-CoA dehydrogenase deficiency/ Medium-chain acyl-CoA dehydrogenase deficiency/Medium-chain 3-keto acyl-CoA thiolase deficiency	Hexanoylcarnitine (C6), Octanoylcarnitine (C8)	11	Medium-chain acyl-CoA dehydrogenase (MCAD) deficiency	4
			Multiple acyl-CoA dehydrogenase (MAD) deficiency - glutaric acidemia type II (GA-II)	0
			Medium-chain 3-keto acyl-CoA thiolase (MCKAT) deficiency	0
Very long-chain acyl-CoA dehydrogenase deficiency	Tetradecanoylcarnitine (C14), Tetradecenoylcarnitine (C14:1)	7	Very long-chain acyl-CoA dehydrogenase (VLCAD) deficiency	2

Screened Disorders	Analytes	Referrals	# Confirmed with Disease	
Short-chain acyl-CoA dehydrogenase deficiency	Butyrylcarnitine (C4)	24	Short-chain acyl-CoA dehydrogenase (SCAD) deficiency	11
			Isobutyryl-CoA dehydrogenase (IBCD) deficiency	2
Medium/short-chain hydroxyl CoA dehydrogenase deficiency	Hydroxybutyrylcarnitine (C4OH), Hydroxyhexanoylcarnitine (C6OH)	2	Medium/short-chain hydroxyl CoA dehydrogenase (M/SCHAD) deficiency	0
Organic Acid Disorders				
Mitochondrial acetoacetyl-CoA thiolase deficiency/2-Methyl-3-hydroxybutyryl-CoA-dehydrogenase deficiency	Tiglylcarnitine (C5:1)	0	Mitochondrial acetoacetyl-CoA thiolase deficiency - beta-ketothiolase (BKT) deficiency	0
			2-Methyl-3-hydroxybutyryl-CoA-dehydrogenase (MHBD) deficiency	0
Glutaryl-CoA dehydrogenase deficiency	Glutarylcarnitine (C5DC)	4	Glutaryl-CoA dehydrogenase deficiency - glutaric aciduria (GA-I)	2
Isovaleryl CoA dehydrogenase deficiency/2-methylbutyryl-CoA dehydrogenase deficiency	Isovalerylcarnitine (C5)	4	Isovaleryl CoA dehydrogenase deficiency - isovaleric acidemia (IVA)	2
			2-Methylbutyrylglycinuria (2MBG) - 2-methylbutyryl-CoA dehydrogenase (2MBCD) deficiency - short/branched chain acyl-CoA dehydrogenase (SBCAD) deficiency	0
3-Methylcrotonyl-CoA carboxylase deficiency/2-Methyl-3-hydroxybutyryl-CoA dehydrogenase deficiency/3-Methylglutaconic aciduria	Hydroxyisovalerylcarnitine (C5OH)	39	3-Methylcrotonyl-CoA carboxylase (3MCC) deficiency	13
			3-Hydroxy-3-methylglutaryl-CoA lyase (HMG) deficiency	0
			2-Methyl-3-hydroxybutyryl-CoA dehydrogenase (MHBD) deficiency - 2-Methyl-3-hydroxybutric acidemia (2M3HBA)	0
			3-Methylglutaconic aciduria (3MGA)	1
Malonyl-CoA decarboxylase deficiency	Malonylcarnitine (C3DC)	1	Malonyl-CoA decarboxylase deficiency - Malonic Aciduria (MA)	1
Propionyl-CoA carboxylase deficiency/Methylmalonyl-CoA mutase deficiency	Propionylcarnitine (C3), Methylmalonylcarnitine (C4DC)	43	Propionyl-CoA carboxylase deficiency (PA)	1
			Methylmalonyl-CoA mutase deficiency (MMA)	0
			Cobalamin A/B deficiency	1
			Cobalamin C/D/F deficiency	0
			Multiple Carboxylase deficiency	0
Urea Cycle Disorders				
Argininosuccinic aciduria/Citrullinemia	Citrulline	4	Argininosuccinic aciduria	0
			Citrullinemia	1
Argininemia	Arginine	1	Argininemia	1
Lysosomal Storage Disorders				
Krabbe Disease	Galactocerebrosidase	47	Krabbe disease possible late onset	1
Pompe Disease	Alpha-glucosidase deficiency	45	Infantile-onset Pompe Disease	1
			Possible late-onset Pompe disease	20
Other Genetic Conditions				
Adrenoleukodystrophy	C26:0 Lysophosphatidylcholine (C26:0 LPC)	22	Male with X-linked Adrenoleukodystrophy (X-ALD)	5
			Female carrier of X-ALD	6
			Zellweger Syndrome	1
			Other Peroxisomal Biogenesis Disorder	0
Biotinidase Deficiency	Biotinidase	6	Biotinidase Deficiency	5
Cystic Fibrosis	Immunoreactive Trypsin	838	Cystic Fibrosis	30
Galactosemia	Galactose Transferase	5	Galactosemia	4
Severe Combined Immunodeficiency (SCID)	T-cell receptor excision circles	93	Classic SCID	5
			Leaky SCID	0
			Variant SCID	6
Total		2992		838

* Infants classified as confirmed for Krabbe disease include those at high risk for disease based on confirmatory enzyme activity testing