



Washington State Newborn Screening Program 2017 Screening and Disorder Summary

2017 Screening Statistics

Washington State Numbers	1 st Qtr	2 nd Qtr	3 rd Qtr	4 th Qtr	Total
Hospital, Birth Center & Home Births	21,045	22,421	23,071	21,278	87,815
Specimens Tested (most infants have two screens)	41,009	43,279	44,245	41,167	169,700

2017 Disorder Statistics

Disorders Diagnosed	Number of Infants				
	1 st Qtr	2 nd Qtr	3 rd Qtr	4 th Qtr	Total
Amino Acid disorders	2	2	0	0	4^a
Biotinidase Deficiency	0	1	0	0	1
Congenital Adrenal Hyperplasia	1	2	2	2	7
Congenital Hypothyroidism	32	25	26	22	105
Cystic Fibrosis	4	2	2	8	16
Fatty Acid Oxidation disorders	4	1	1	3	9^b
Galactosemia	1	0	1	2	4
Organic Acid disorders	3	1	1	1	6^c
Severe Combined Immunodeficiency	1	0	0	3	4^d
Sickle Cell Disease and Other Clinically Significant Hemoglobinopathies	4	10	9	2	25
All Dried Blood Tests Combined	52	44	41	42	178
Early Hearing Loss	16	32	34	65	147
All Disorders Combined	68	73	75	107	325

^a Three infants with phenylketonuria (PKU) and one infant with argininosuccinic aciduria (ASA). Excludes one infant with Citrin deficiency a condition not on our screening panel.

^b Eight infants with medium-chain acyl-CoA dehydrogenase (MCAD) deficiency and one infant with trifunctional protein (TFP) deficiency.

^c One infant with glutaric acidemia type 1 (GA-1), two infants with isovaleric acidemia (IVA), and three infants with methylmalonic acidemia (MMA). Excludes five infants with 3-methylcrotonyl carboxylase (3-MCC), one infant with 2-methyl butyryl-CoA dehydrogenase (2-MBDH) deficiency, and one infant with glutaric acidemia type-II (GA-II), conditions not on our screening panel.

^d Excludes three infants with Di George syndrome, one infant with Noonan syndrome, one infant with Jacobsen's syndrome, one infant with Soto's syndrome, one infant with CHARGE syndrome and three infants with other T-Cell lymphopenias - conditions not on our screening panel.