



Washington State Newborn Screening Program 2016 Screening and Disorder Summary

2016 Screening Statistics

Washington State Numbers	1 st Qtr	2 nd Qtr	3 rd Qtr	4 th Qtr	Total
Hospital*, Birth Center & Home Births	21,635	23,173	23,027	21,695	90,530
Specimens Tested (most infants have two screens)	41,654	44,112	46,240	41,728	173,734

* Excludes babies born at Bremerton and Whidbey Island Naval hospitals.
These births are no longer excluded effective May 1, 2016

2016 Disorder Statistics

Disorders Diagnosed	Number of Infants				
	1 st Qtr	2 nd Qtr	3 rd Qtr	4 th Qtr	Total
Amino Acid disorders	1	1	2	3	7^a
Biotinidase Deficiency	0	1	0	0	1
Congenital Adrenal Hyperplasia	0	2	3	0	5
Congenital Hypothyroidism	35	22	17	34	108
Cystic Fibrosis	5	2	3	1	11
Fatty Acid Oxidation disorders	2	2	0	3	7^b
Galactosemia	4	0	1	0	5
Organic Acid disorders	0	0	0	1	1^c
Severe Combined Immunodeficiency	0	0	0	0	0
Sickle Cell Disease and Other Clinically Significant Hemoglobinopathies	6	5	5	8	24^d
All Dried Blood Tests Combined	53	35	31	49	169
Early Hearing Loss	14	18	50	55	137
All Disorders Combined	67	53	81	104	306

^a Five infants with phenylketonuria (PKU) and two infants with citrullinemia type I (CIT-I). Does not include one infant with citrin deficiency (a condition not on our screening panel)

^b Four infants with medium-chain acyl-CoA dehydrogenase (MCAD) deficiency, two infants with very-long chain acyl-CoA dehydrogenase (VLCAD) deficiency, and one infant with long-chain L-3-hydroxy acyl-CoA dehydrogenase deficiency (LCHAD)

^c One infant with glutaric acidemia type I (GA-I). Does not include two infants with 3-Methylcrotonyl-CoA carboxylase (3-MCC) deficiency and one infant with cobalamin C deficiency (conditions not on our screening panel)

^d Does not include one infant with other T-cell lymphopenia

Washington State Department of Health • Newborn Screening Program

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