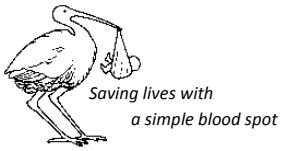

Washington State Newborn Screening Program
2014 Screening and Disorder Summary

2014 Screening Statistics

Washington State Numbers	1st Qtr	2nd Qtr	3rd Qtr	4th Qtr	Total
Hospital*, Birth Center & Home Births	21,163	22,574	23,121	20,540	**87,398**
Specimens Tested (most infants have two screens)	40,572	42,577	44,778	45,965	**173,908**

* Excludes babies born at Bremerton and Whidbey Island Naval hospitals.

2014 Disorder Statistics

Disorders Diagnosed	Number of Infants				
	1st Qtr	2nd Qtr	3rd Qtr	4th Qtr	Total
Amino Acid disorders	2	3	1	0	**6**[a]
Biotinidase Deficiency	0	1	0	0	**1**
Congenital Adrenal Hyperplasia	1	2	0	2	**5**
Congenital Hypothyroidism	15	22	16	37	**113**
Cystic Fibrosis	0	5	4	5	**14**
Fatty Acid Oxidation disorders	0	1	5	3	**10**[b]
Galactosemia	1	1	3	5	**10**
Organic Acid disorders	0	1	0	2	**3**[c]
Severe Combined Immunodeficiency	0	1	0	1	**1**[d]
Sickle Cell Disease and Other Clinically Significant Hemoglobinopathies	3	4	12	7	**26**
All Dried Blood Tests Combined	**22**	**42**	**41**	**62**	**189**
Early Hearing Loss	**7**	**21**	**62**	**58**	**148**
All Disorders Combined	**29**	**63**	**103**	**120**	**337**

[a] Four infants with phenylketonuria (PKU), one infant with argininosuccinic aciduria (ASA) and one infant with maple syrup urine disease (MSUD).
[b] Includes four infants with medium-chain acyl-CoA dehydrogenase (MCAD) deficiency, four infants with very-long chain acyl-CoA dehydrogenase (VLCAD) deficiency, two infants with long-chain L-3-hydroxy acyl-CoA dehydrogenase (LCHAD) deficiency, but excludes one infant with Carnitine Palmitoyltransferase II (CPT-II) deficiency (a condition not on our screening panel).
[c] Includes one infant with Isovaleric acidemia (IVA), two infants with methylmalonic acidemia (MMA), and excludes three infants with 3-methylcrotonyl CoA carboxylase (3-MCC) deficiency (a condition not on our screening panel).
[d] Does not include 3 infants with other T-cell lymphopeneas

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Washington State Department of Health
DOH 951-147 Jun 2015