Saving lives with a simple blood spot

Washington State Newborn Screening Program
2015 Screening and Disorder Summary

2015 Screening Statistics					
Washington State Numbers	**1st Qtr**	**2nd Qtr**	**3rd Qtr**	**4th Qtr**	**Total**
Hospital*, Birth Center & Home Births	20,847	22,305	23,479	21,736	**88,367**
Specimens Tested (most infants have two screens)	39,914	42,446	45,105	42,250	**169,715**

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

2015 Disorder Statistics					
Disorders Diagnosed	**Number of Infants**				
	1st Qtr	**2nd Qtr**	**3rd Qtr**	**4th Qtr**	**Total**
Amino Acid disorders	2	4	4	1	**11**[a]
Biotinidase Deficiency	0	1	0	0	**1**
Congenital Adrenal Hyperplasia	0	3	0	3	**6**
Congenital Hypothyroidism	20	17	21	24	**82**
Cystic Fibrosis	1	6	4	1	**12**
Fatty Acid Oxidation disorders	2	2	5	2	**11**[b]
Galactosemia	4	7	4	1	**16**
Organic Acid disorders	0	0	0	3	**3**[c]
Severe Combined Immunodeficiency	0	0	1	0	**1**[d]
Sickle Cell Disease and Other Clinically Significant Hemoglobinopathies	4	8	2	4	**18**
All Dried Blood Tests Combined	**33**	**48**	**41**	**39**	**161**
Early Hearing Loss	**24**	**23**	**38**	**26**	**111**
All Disorders Combined	**57**	**71**	**79**	**65**	**272**

[a] Eight infants with phenylketonuria (PKU) and one infant with maple syrup urine disease (MSUD).
[b] Includes seven infants with medium-chain acyl-CoA dehydrogenase (MCAD) deficiency, one infant with very-long chain acyl-CoA dehydrogenase (VLCAD) deficiency, and one infant with long-chain L-3-hydroxy acyl-CoA dehydrogenase (LCHAD) deficiency.
[c] Does not include one infant with 3-methylcrotonyl CoA carboxylase (3-MCC) deficiency (a condition not on our screening panel).
[d] Does not include three infants with other T-cell lymphopeneas.

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