Supplementary table S1. Additional genetic syndromes associated with pediatric pulmonary hypertension, 36 of 1475 patients (2.4%). See Table 5 for most common syndromes (214 of 1475 patients, 14.5%).

		WSPH Group		
Syndrome	**Frequency**	**Patient 1**	**Patient 2**	**Patient 3**
ABCA 3 Surfactant Deficiency	1	3.2.3		
Surfactant Type C Deficiency	1	3.7.1		
Adam Oliver	1	1.2		
Alagille	3	1.4.4	1.4.4	3.7.3
Baraitser-Winter	1	1.4.4		
Beckwith-Wiedeman	2	3.7.1	1.4.4	
Congenital muscular dystrophy	1	3.7		
Cornelia De Lange	2	3.7.3	3.7.1	
Dicer 1	1	5.5		
Donnai-Barrow	1	3.7.3		
Eagle-Barrett	1	1.1		
Ehlers-Danlos	1	3.7		
Fryns	2	3.7.3	3.7.3	
Holt-Oram	1	1.4.4		
Jacobsen	2	3.7.1	1.4.4	
Kleefstra	2	1”	1.4.4	
Klippel-Feil	1	3.3		
Klippel-Trenanunay-Weber	1	1.4.4		
Mayer-Rokitansky-Kuster-Hauser	1	1.4.4		
Mowat-Wilson	1	2.3		
Osteogenesis Imperfecta Type III	1	3.7		
Pierre Robin	3	3.7.1	3.7.3	1.4.4
Pompe’s Disease	1	5.3		
Smith-Lemli-Opitz	1	2.4		
Trisomy 6	1	1”		
Turner	1	3.7.3		
Wager	1	1.4.4		
TOTAL	**36**			