

Kids lung register disease classifications Category

Kids lung register disease classifications Subcategory

< All >	< All >
A1 - DPLD-Diffuse developmental disorders	ACDnoMisalign. pulm.veins +anophthalmia, cong. HeartD, Diaph.Hernia, Lung Hypoplasia, Men.Retard.
A1 - DPLD-Diffuse developmental disorders	Acinar dysplasia
A1 - DPLD-Diffuse developmental disorders	Alveolar capillary dysplasia with misalignment pulmonary vein
A1 - DPLD-Diffuse developmental disorders	ACD no misalignment
A1 - DPLD-Diffuse developmental disorders	ACD+MPV+FOXF1 mutation
A1 - DPLD-Diffuse developmental disorders	Congenital alveolar dysplasia
A1 - DPLD-Diffuse developmental disorders	Alveolar Dysgenesis/Primary Pulmonary Hypoplasia
A2 - DPLD-Growth abnormalities deficient alveolarisation	Intrauterine growth retardation (alcohol)
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with thoracic dystrophies
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with neuromuscular dysfunction
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with thoracic mass
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with oligohydramnion
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with polyhydramnion
A2 - DPLD-Growth abnormalities deficient alveolarisation	Pulmonary hypoplasia associated with diaphragmatic hernia
A2 - DPLD-Growth abnormalities deficient alveolarisation	Related to chromosomal disorders
A2 - DPLD-Growth abnormalities deficient alveolarisation	Related to congenital heart disease
A2 - DPLD-Growth abnormalities deficient alveolarisation	Related to preterm birth (BPD-cLDI)
A2 - DPLD-Growth abnormalities deficient alveolarisation	Related to preterm birth (Wilson Mikity, new BPD)
A2 - DPLD-Growth abnormalities deficient alveolarisation	Chronic tachypnoe of infancy (CTI)
A3 - DPLD-Infant conditions of undefined etiology	Neuroendocrine cell hyperplasia of infancy (NEHI)
A3 - DPLD-Infant conditions of undefined etiology	Chronic tachypnoe of infancy (CTI), usual
A3 - DPLD-Infant conditions of undefined etiology	Chronic tachypnoe of infancy (CTI), aberrant
A3 - DPLD-Infant conditions of undefined etiology	Pulmonary interstitial glyco-genosis (PIG)
A3 - DPLD-Infant conditions of undefined etiology	Pulmonary interstitial glyco-genosis (PIG) primary
A3 - DPLD-Infant conditions of undefined etiology	Pulmonary interstitial glyco-genosis (PIG) associated with other lung diseases
A4 - DPLD-related to alveolar surfactant region	ABCA3 mutations 1
A4 - DPLD-related to alveolar surfactant region	ABCA3 mutations 2
A4 - DPLD-related to alveolar surfactant region	Acute Fibrinous and Organizing Pneumonia
A4 - DPLD-related to alveolar surfactant region	Alveolar microlithiasis
A4 - DPLD-related to alveolar surfactant region	Chronic pneumonitis of infancy (CPI)
A4 - DPLD-related to alveolar surfactant region	Combined pulmonary fibrosis and emphysema
A4 - DPLD-related to alveolar surfactant region	Cryptogenic Organizing Pneumonia (Bronchiolitis Obliterans Organizing Pneumonia)
A4 - DPLD-related to alveolar surfactant region	Diffuse Alveolar Damage and Acute Interstitial Pneumonia
A4 - DPLD-related to alveolar surfactant region	DIP
A4 - DPLD-related to alveolar surfactant region	Lipoidpneumonitis, Cholesterol pneumonia
A4 - DPLD-related to alveolar surfactant region	Nrx21 gene defect
A4 - DPLD-related to alveolar surfactant region	Nonspecific interstitial pneumonia (NSIP)
A4 - DPLD-related to alveolar surfactant region	NSIP, cellular
A4 - DPLD-related to alveolar surfactant region	NSIP, fibrotic
A4 - DPLD-related to alveolar surfactant region	NSIP, DIP, PAP pattern
A4 - DPLD-related to alveolar surfactant region	NSIP +/- DIP +/- PAP pattern
A4 - DPLD-related to alveolar surfactant region	NSIP, DIP pattern
A4 - DPLD-related to alveolar surfactant region	NSIP+PAP+Microvasulopathie
A4 - DPLD-related to alveolar surfactant region	NSIP, PAP pattern
A4 - DPLD-related to alveolar surfactant region	PAP histopath +
A4 - DPLD-related to alveolar surfactant region	PAP, adult NO GMCSF autoantibodies
A4 - DPLD-related to alveolar surfactant region	PAP, adult with GMCSF autoantibodies
A4 - DPLD-related to alveolar surfactant region	PAP+DIP+cholesterin granulomas
A4 - DPLD-related to alveolar surfactant region	PAP, GATA2 mutation
A4 - DPLD-related to alveolar surfactant region	PAP, GM-CSF-RA Mutation
A4 - DPLD-related to alveolar surfactant region	PAP, hematol
A4 - DPLD-related to alveolar surfactant region	PAP, juvenile
A4 - DPLD-related to alveolar surfactant region	PAP, juvenile Reunion
A4 - DPLD-related to alveolar surfactant region	PAP, MARS mutation
A4 - DPLD-related to alveolar surfactant region	PAP, Lysinuric proteinuria
A4 - DPLD-related to alveolar surfactant region	PAP, neonatal
A4 - DPLD-related to alveolar surfactant region	PAP, secondary to associated disease
A4 - DPLD-related to alveolar surfactant region	Respiratory Bronchiolitis-Interstitial Lung Disease
A4 - DPLD-related to alveolar surfactant region	Surfactant protein B mutations
A4 - DPLD-related to alveolar surfactant region	Surfactant protein C mutations
A4 - DPLD-related to alveolar surfactant region	Usual interstitial pneumonitis
Ax - DPLD-unclear RDS in the mature neonate	Familial
Ax - DPLD-unclear RDS in the mature neonate	No or very low SP-C biochemically
Ax - DPLD-unclear RDS in the mature neonate	No SP-B biochemically
Ax - DPLD-unclear RDS in the mature neonate	Pulmonary hypertension
Ay - DPLD-unclear RDS in the almost (30-36 wks) mature neonate	Familial
Ay - DPLD-unclear RDS in the almost (30-36 wks) mature neonate	No or very low SP-C biochemically
Ay - DPLD-unclear RDS in the almost (30-36 wks) mature neonate	No SP-B biochemically
Ay - DPLD-unclear RDS in the almost (30-36 wks) mature neonate	Pulmonary hypertension
B1 - DPLD-related to systemic disease processes	Achondroplasia, zB Cartilage-Hair Hypoplasia
B1 - DPLD-related to systemic disease processes	Acute idiopathic DAH of infancy
B1 - DPLD-related to systemic disease processes	Alagille Syndrome (arteriohepatic dysplasia)
B1 - DPLD-related to systemic disease processes	Antibasement Membrane Antibody Disease (Good pasture's Syndrome)
B1 - DPLD-related to systemic disease processes	Antisynthetase Syndrome
B1 - DPLD-related to systemic disease processes	Behcet's Syndrome
B1 - DPLD-related to systemic disease processes	Birt-Hogg-Dube Syndrome
B1 - DPLD-related to systemic disease processes	Blau Syndrome (polyarthritis, uveitis, papuloerythematous rash) + rarely lung
B1 - DPLD-related to systemic disease processes	Cantu Syndrome (Hypertrichosis, facial, Osteochondrodysplasia, PHT)
B1 - DPLD-related to systemic disease processes	Central hypoventilation syndrome (Ondine)
B1 - DPLD-related to systemic disease processes	Celiac disease
B1 - DPLD-related to systemic disease processes	Celiac disease + pulm hemorrhage (Lane-Hamilton)
B1 - DPLD-related to systemic disease processes	Congenital muscle disease
B1 - DPLD-related to systemic disease processes	Cryoglobulemic vasculitis
B1 - DPLD-related to systemic disease processes	Diffuse alveolar hemorrhage due to vasculitic disorders
B1 - DPLD-related to systemic disease processes	Diffuse alveolar hemorrhage due to COPA mutations
B1 - DPLD-related to systemic disease processes	Heiner syndrome (milk induced)
B1 - DPLD-related to systemic disease processes	Idiopathic pulmonary capillaritis
B1 - DPLD-related to systemic disease processes	DIP - pattern
B1 - DPLD-related to systemic disease processes	Disseminated Visceral Giant Cell Angiitis
B1 - DPLD-related to systemic disease processes	EGPA -- Eosinophilic granulomatosis with polyangiitis (Churg Strauss)
B1 - DPLD-related to systemic disease processes	Erdheim-Chester Disease
B1 - DPLD-related to systemic disease processes	Familial dysautonomia (Chromosome 9q31 encoding ICAP)
B1 - DPLD-related to systemic disease processes	Familial ILD, growth defic., hepatopathy, normal psychomotor development, triventricular hydrocephalus
B1 - DPLD-related to systemic disease processes	Filamin A Mutation
B1 - DPLD-related to systemic disease processes	Giant Cell Arteritis
B1 - DPLD-related to systemic disease processes	GPA -- Granulomatosis with polyangiitis (Wegener)
B1 - DPLD-related to systemic disease processes	Hermansky-Pudlak Syndrome
B1 - DPLD-related to systemic disease processes	Hoyerall Hreidasson Syndrome (Dyskeratosis congenita)
B1 - DPLD-related to systemic disease processes	Hypercalcaemic hypercalcaemia-DPLD
B1 - DPLD-related to systemic disease processes	IgA vasculitis (Schoenlein-Hennoch)
B1 - DPLD-related to systemic disease processes	Immune-mediated/collagen vascular disorders





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Paracoccidioides Brasiliensis  
Paragonimiasis  
Parainfluenza  
Pneumocystis Carinii Pneumonia  
Pseudoallescheria Boydii  
Pseudomonas Aeruginosa  
Respiratory Syncytial Virus  
Rhodococcus Equi and Malakoplakia  
Schistosomiasis  
Sporothrix Schenkii  
Staphylococcus Sp  
Streptococcus Pneumoniae  
Streptococcus Sp (Nonpneumococcal)  
Strongyloidiasis  
Toxoplasmosis  
Tropheryma Whipplei (Whipple's Disease)  
Varicella-Zoster  
Yersinia Pestis  
Zygomycetes  
acute Pneumonia