ANOMALIES DU DEVELOPPEMENT PULMONAIRE

ORPHAcode	Disease ID	Nature	Classification Level	Orphanet classification of rare respiratory diseases (Pat)
ORPHA:97957	12976	Clinical entity	Group of disorders	Respiratory or thoracic malformation
ORPHA:182108	18215	Clinical entity	Group of disorders	Thoracic malformation
ORPHA:1505	537	Clinical entity	Group of disorders	Short rib-polydactyly syndrome
ORPHA:474	283	Clinical entity	Disorder	Jeune syndrome
ORPHA:289	287	Clinical entity	Disorder	Ellis Van Creveld syndrome
ORPHA:1515	1682	Clinical entity	Disorder	Cranioectodermal dysplasia
ORPHA:93268	12211	Clinical entity	Disorder	Short rib-polydactyly syndrome, Beemer-Langer type
ORPHA:93269	12212	Clinical entity	Disorder	Short rib-polydactyly syndrome, Majewski type
ORPHA:93270	12213	Clinical entity	Disorder	Short rib-polydactyly syndrome, Saldino-Noonan type
ORPHA:93271	12214	Clinical entity	Disorder	Short rib-polydactyly syndrome, Verma-Naumoff type
ORPHA:397715	22695	Clinical entity	Disorder	Joubert syndrome with Jeune asphyxiating thoracic dystrophy
ORPHA:498497	25928	Clinical entity	Disorder	Short rib-polydactyly syndrome type 5
ORPHA:957	1278	Clinical entity	Disorder	Acropectorovertebral dysplasia
ORPHA:994	1310	Clinical entity	Disorder	Fetal akinesia deformation sequence
ORPHA:1486	1660	Clinical entity	Disorder	Lethal congenital contracture syndrome type 1
ORPHA:1803	1799	Clinical entity	Disorder	Thoracomelic dysplasia
ORPHA:1861	1838	Clinical entity	Disorder	Thoracic dysplasia-hydrocephalus syndrome
ORPHA:2017	1951	Clinical entity	Disorder	Sternal cleft
ORPHA:2391	2222	Clinical entity	Disorder	Congenitally short costocoracoid ligament
ORPHA:2470	2283	Clinical entity	Disorder	Matthew-Wood syndrome
ORPHA:2753	2509	Clinical entity	Disorder	Orofaciodigital syndrome type 4
ORPHA:3181	2853	Clinical entity	Disorder	Sprengel deformity
ORPHA:3317	2947	Clinical entity	Disorder	Thoracolaryngopelvic dysplasia
ORPHA:73230	11038	Clinical entity	Disorder	Ossification anomalies-psychomotor developmental delay syndrome
ORPHA:464366	24023	Clinical entity	Disorder	NEK9-related lethal skeletal dysplasia
ORPHA:182111	18216	Clinical entity	Group of disorders	Respiratory malformation
ORPHA:774	236	Clinical entity	Disorder	Hereditary hemorrhagic telangiectasia
ORPHA:2414	534	Clinical entity	Disorder	Congenital pulmonary lymphangiectasia
ORPHA:2444	538	Clinical entity	Disorder	Congenital pulmonary airway malformation
ORPHA:280827	20506	Clinical entity	Subtype of disorder	Congenital pulmonary airway malformation type 0
ORPHA:280832	20507	Clinical entity	Subtype of disorder	Congenital pulmonary airway malformation type 1
ORPHA:280840	20508	Clinical entity	Subtype of disorder	Congenital pulmonary airway malformation type 2
ORPHA:280847	20509	Clinical entity	Subtype of disorder	Congenital pulmonary airway malformation type 3
ORPHA:280854	20510	Clinical entity	Subtype of disorder	Congenital pulmonary airway malformation type 4
ORPHA:3346	597	Clinical entity	Disorder	Tracheal agenesis
ORPHA:984	715	Clinical entity	Disorder	Pulmonary agenesis

ORPHA:2357	797	Clinical entity	Disorder	Bronchogenic cyst
ORPHA:2040	798	Clinical entity	Disorder	Congenital respiratory-biliary fistula
ORPHA:1928	802	Clinical entity	Disorder	Congenital lobar emphysema
ORPHA:1120	1400	Clinical entity	Disorder	Lung agenesis-heart defect-thumb anomalies syndrome
ORPHA:2004	1941	Clinical entity	Disorder	Laryngotracheoesophageal cleft
ORPHA:93938	12498	Clinical entity	Subtype of disorder	Laryngotracheoesophageal cleft type 1
ORPHA:93939	12499	Clinical entity	Subtype of disorder	Laryngotracheoesophageal cleft type 2
ORPHA:93940	12500	Clinical entity	Subtype of disorder	Laryngotracheoesophageal cleft type 3
ORPHA:93941	12501	Clinical entity	Subtype of disorder	Laryngotracheoesophageal cleft type 4
ORPHA:280205	20437	Clinical entity	Subtype of disorder	Laryngotracheoesophageal cleft type 0
ORPHA:2257	2125	Clinical entity	Disorder	Primary pulmonary hypoplasia
ORPHA:2407	2233	Clinical entity	Disorder	LOC syndrome
ORPHA:2470	2283	Clinical entity	Disorder	Matthew-Wood syndrome
ORPHA:3035	2729	Clinical entity	Disorder	Growth delay-hydrocephaly-lung hypoplasia syndrome
ORPHA:3317	2947	Clinical entity	Disorder	Thoracolaryngopelvic dysplasia
ORPHA:1132	3428	Clinical entity	Group of disorders	Aortic arch defects
ORPHA:99075	14092	Clinical entity	Disorder	Encircling double aortic arch
ORPHA:99076	14093	Clinical entity	Disorder	Persistent fifth aortic arch
ORPHA:99077	14094	Clinical entity	Disorder	Kommerell diverticulum
ORPHA:99078	14095	Clinical entity	Disorder	Neuhauser anomaly
ORPHA:99079	14096	Clinical entity	Disorder	Cervical aortic arch
ORPHA:99081	14098	Clinical entity	Disorder	Right aortic arch
ORPHA:99082	14099	Clinical entity	Disorder	Dysphagia lusoria
ORPHA:3161	3457	Clinical entity	Disorder	Congenital pulmonary sequestration
ORPHA:280802	20503	Clinical entity	Subtype of disorder	Intralobar congenital pulmonary sequestration
ORPHA:280811	20504	Clinical entity	Subtype of disorder	Extralobar congenital pulmonary sequestration
ORPHA:280821	20505	Clinical entity	Subtype of disorder	Communicating congenital bronchopulmonary-foregut malformation
ORPHA:70589	10960	Clinical entity	Disorder	Bronchopulmonary dysplasia
ORPHA:95430	12589	Clinical entity	Disorder	Congenital tracheomalacia
ORPHA:411501	23013	Clinical entity	Disorder	Williams-Campbell syndrome
ORPHA:454750	23661	Clinical entity	Disorder	Isolated tracheoesophageal fistula

HYPERTENSION PULMONAIRE

ORPHAcode	Disease ID	Nature	Classification Level	Orphanet classification of rare respiratory diseases (Pat)
ORPHA:71198	10976	Clinical entity	Group of disorders	Rare pulmonary hypertension
ORPHA:70591	10962	Clinical entity	Disorder	**Chronic thromboembolic pulmonary hypertension**
ORPHA:182090	18210	Clinical entity	Group of disorders	Pulmonary arterial hypertension
ORPHA:422	3444	Clinical entity	Disorder	**Idiopathic/heritable pulmonary arterial hypertension**
ORPHA:275766	20327	Clinical entity	Subtype of disorder	**Idiopathic pulmonary arterial hypertension**
ORPHA:275777	20328	Clinical entity	Subtype of disorder	**Heritable pulmonary arterial hypertension**
ORPHA:275786	20329	Clinical entity	Group of disorders	Drug- or toxin-induced pulmonary arterial hypertension
ORPHA:275791	20330	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with another disease
ORPHA:275798	20331	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with connective tissue disease
ORPHA:275803	20332	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with congenital heart disease
ORPHA:97214	12855	Clinical entity	Disorder	**Eisenmenger syndrome**
ORPHA:275808	20333	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with HIV infection
ORPHA:275813	20334	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with portal hypertension
ORPHA:275823	20335	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with schistosomiasis
ORPHA:275828	20336	Clinical entity	Group of disorders	Pulmonary arterial hypertension associated with chronic hemolytic anemia
ORPHA:275837	20337	Clinical entity	Group of disorders	Pulmonary hypertension owing to lung disease and/or hypoxia
ORPHA:275844	20338	Clinical entity	Group of disorders	Pulmonary hypertension with unclear multifactorial mechanism
ORPHA:275853	20339	Clinical entity	Group of disorders	Syndrome with pulmonary hypertension as a major feature
ORPHA:52047	10685	Clinical entity	Disorder	**Braddock syndrome**
ORPHA:363694	22340	Clinical entity	Disorder	**Hyperuricemia-pulmonary hypertension-renal failure-alkalosis syndrome**
ORPHA:431353	23248	Clinical entity	Group of disorders	Pulmonary veno-occlusive disease and/or pulmonary capillary haemangiomatosis
ORPHA:31837	9801	Clinical entity	Disorder	**Pulmonary venoocclusive disease**

MALADIES PULMONAIRES RARES

ORPHAcode	Disease ID	Nature	Classification Level	Orphanet classification of rare respiratory diseases (Pat)
ORPHA:101944	14880	Clinical entity	Group of disorders	Rare pulmonary disease
ORPHA:586	49	Clinical entity	Disorder	Cystic fibrosis
ORPHA:60	194	Clinical entity	Disorder	Alpha-1-antitrypsin deficiency
ORPHA:2140	506	Clinical entity	Disorder	Congenital diaphragmatic hernia
ORPHA:2903	564	Clinical entity	Disorder	Familial spontaneous pneumothorax
ORPHA:244	665	Clinical entity	Disorder	Primary ciliary dyskinesia
ORPHA:1164	744	Clinical entity	Disorder	Allergic bronchopulmonary aspergillosis
ORPHA:3389	863	Clinical entity	Disorder	Tuberculosis
ORPHA:661	2497	Clinical entity	Disorder	Ondine syndrome
ORPHA:3471	3050	Clinical entity	Disorder	Young syndrome
ORPHA:3167	3091	Clinical entity	Disorder	Siegler-Brewer-Carey syndrome
ORPHA:1303	7030	Clinical entity	Disorder	Bronchiolitis obliterans with obstructive pulmonary disease
ORPHA:3348	7032	Clinical entity	Disorder	Tracheobronchopathia osteochondroplastica
ORPHA:36238	10413	Clinical entity	Disorder	Staphylococcal necrotizing pneumonia
ORPHA:60026	10798	Clinical entity	Disorder	Pulmonary nodular lymphoid hyperplasia
ORPHA:60032	10801	Clinical entity	Disorder	Recurrent respiratory papillomatosis
ORPHA:60033	10802	Clinical entity	Disorder	Idiopathic bronchiectasis
ORPHA:70588	10959	Clinical entity	Disorder	Meconium aspiration syndrome
ORPHA:70590	10961	Clinical entity	Disorder	Infantile apnea
ORPHA:99981	14554	Clinical entity	Subtype of disorder	Apnea of prematurity
ORPHA:90060	11951	Clinical entity	Disorder	Diffuse alveolar hemorrhage
ORPHA:90066	11957	Clinical entity	Disorder	Pneumonia caused by Pseudomonas aeruginosa infection
ORPHA:90291	12002	Clinical entity	Disorder	Systemic sclerosis
ORPHA:220393	18905	Clinical entity	Subtype of disorder	Diffuse cutaneous systemic sclerosis
ORPHA:220402	18906	Clinical entity	Subtype of disorder	Limited cutaneous systemic sclerosis
ORPHA:220407	18907	Clinical entity	Subtype of disorder	Limited systemic sclerosis
ORPHA:99803	14376	Clinical entity	Disorder	Haddad syndrome
ORPHA:140896	16993	Clinical entity	Disorder	Severe acute respiratory syndrome
ORPHA:168593	17779	Clinical entity	Disorder	Sudden infant death-dysgenesis of the testes syndrome
ORPHA:171700	17929	Clinical entity	Disorder	Diffuse panbronchiolitis
ORPHA:178320	18013	Clinical entity	Disorder	Acute lung injury
ORPHA:182095	18211	Clinical entity	Group of disorders	Interstitial lung disease
ORPHA:264656	20104	Clinical entity	Group of disorders	Interstitial lung disease specific to childhood
ORPHA:264665	20106	Clinical entity	Group of disorders	Primary interstitial lung disease specific to childhood
ORPHA:137631	16696	Clinical entity	Disorder	Lung fibrosis-immunodeficiency-46,XX gonadal dysgenesis syndrome
ORPHA:264670	20107	Clinical entity	Group of disorders	Primary interstitial lung disease specific to childhood due to alveolar structure disorder
ORPHA:70587	10958	Clinical entity	Disorder	Infant acute respiratory distress syndrome
ORPHA:100049	14622	Clinical entity	Group of disorders	Primary interstitial lung disease specific to childhood due to pulmonary surfactant protein anomalies
ORPHA:209905	18666	Clinical entity	Disorder	Brain-lung-thyroid syndrome
ORPHA:217563	18862	Clinical entity	Disorder	Neonatal acute respiratory distress due to SP-B deficiency
ORPHA:264675	20108	Clinical entity	Disorder	Hereditary pulmonary alveolar proteinosis
ORPHA:440402	23404	Clinical entity	Disorder	Interstitial lung disease due to ABCA3 deficiency
ORPHA:440427	23408	Clinical entity	Disorder	Severe early-onset pulmonary alveolar proteinosis due to MARS deficiency
ORPHA:572428	28746	Clinical entity	Disorder	Infantile-onset pulmonary alveolar proteinosis-hypogammaglobulinemia
ORPHA:306504	21219	Clinical entity	Disorder	Junctional epidermolysis bullosa with respiratory and renal involvement
ORPHA:264683	20109	Clinical entity	Group of disorders	Primary interstitial lung disease specific to childhood due to alveolar vascular disorder
ORPHA:2414	534	Clinical entity	Disorder	Congenital pulmonary lymphangiectasia
ORPHA:662	6520	Clinical entity	Disorder	Yellow nail syndrome
ORPHA:210122	18684	Clinical entity	Disorder	Congenital alveolar capillary dysplasia
ORPHA:264688	20110	Clinical entity	Disorder	Congenital chylothorax
ORPHA:264691	20111	Clinical entity	Disorder	Isolated pulmonary capillaritis
ORPHA:352629	22087	Clinical entity	Disorder	16q24.1 microdeletion syndrome
ORPHA:264694	20112	Clinical entity	Group of disorders	Interstitial lung disease specific to infancy
ORPHA:91359	12133	Clinical entity	Disorder	Chronic pneumonitis of infancy

ORPHA:217557	18860	Clinical entity	Disorder	Pulmonary interstitial glycogenosis
ORPHA:217560	18861	Clinical entity	Disorder	Neuroendocrine cell hyperplasia of infancy
ORPHA:264699	20113	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to childhood associated with a systemic disease
ORPHA:264704	20114	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to childhood associated with a connective tissue disease
ORPHA:92	720	Clinical entity	Group of disorders	Juvenile idiopathic arthritis
ORPHA:85410	11710	Clinical entity	Disorder	Oligoarticular juvenile idiopathic arthritis
ORPHA:247839	19555	Clinical entity	Subtype of disorder	OBSOLETE: Oligoarticular juvenile idiopathic arthritis with anti-nuclear antibodies
ORPHA:247846	19556	Clinical entity	Subtype of disorder	OBSOLETE: Oligoarticular juvenile idiopathic arthritis without anti-nuclear antibodies
ORPHA:85414	11711	Clinical entity	Disorder	Systemic-onset juvenile idiopathic arthritis
ORPHA:85436	11713	Clinical entity	Disorder	Psoriasis-related juvenile idiopathic arthritis
ORPHA:85438	11715	Clinical entity	Disorder	Enthesitis-related juvenile idiopathic arthritis
ORPHA:91140	12117	Clinical entity	Disorder	Unspecified juvenile idiopathic arthritis
ORPHA:404580	22944	Clinical entity	Group of disorders	Polyarticular juvenile idiopathic arthritis
ORPHA:85408	11709	Clinical entity	Disorder	Rheumatoid factor-negative polyarticular juvenile idiopathic arthritis
ORPHA:247854	19557	Clinical entity	Subtype of disorder	OBSOLETE: Rheumatoid factor-negative juvenile idiopathic arthritis with anti-nuclear antibodies
ORPHA:247861	19558	Clinical entity	Subtype of disorder	OBSOLETE: Rheumatoid factor-negative juvenile idiopathic arthritis without anti-nuclear antibodies
ORPHA:85435	11712	Clinical entity	Disorder	Rheumatoid factor-positive polyarticular juvenile idiopathic arthritis
ORPHA:93552	12394	Clinical entity	Disorder	Pediatric systemic lupus erythematosus
ORPHA:93568	12410	Clinical entity	Disorder	Juvenile polymyositis
ORPHA:93672	12464	Clinical entity	Disorder	Juvenile dermatomyositis
ORPHA:264709	20115	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to childhood associated with a systemic vasculitis
ORPHA:761	749	Clinical entity	Disorder	Immunoglobulin A vasculitis
ORPHA:264714	20116	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to childhood associated with a granulomatous disease
ORPHA:379	176	Clinical entity	Disorder	Chronic granulomatous disease
ORPHA:90340	12018	Clinical entity	Disorder	Blau syndrome
ORPHA:264719	20117	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to childhood associated with a metabolic disease
ORPHA:405	3600	Clinical entity	Disorder	Familial hypocalciuric hypercalcemia
ORPHA:93372	12300	Clinical entity	Subtype of disorder	Familial hypocalciuric hypercalcemia type 1
ORPHA:101049	14760	Clinical entity	Subtype of disorder	Familial hypocalciuric hypercalcemia type 2
ORPHA:101050	14761	Clinical entity	Subtype of disorder	Familial hypocalciuric hypercalcemia type 3
ORPHA:77260	11103	Clinical entity	Subtype of disorder	Gaucher disease type 2
ORPHA:77261	11104	Clinical entity	Subtype of disorder	Gaucher disease type 3
ORPHA:77293	11106	Clinical entity	Disorder	Niemann-Pick disease type B
ORPHA:231500	19195	Clinical entity	Subtype of disorder	Hermansky-Pudlak syndrome with pulmonary fibrosis
ORPHA:444092	23481	Clinical entity	Disorder	Autoimmune interstitial lung disease-arthritis syndrome
ORPHA:264735	20119	Clinical entity	Group of disorders	Interstitial lung disease specific to adulthood
ORPHA:264740	20120	Clinical entity	Group of disorders	Primary interstitial lung disease specific to adulthood
ORPHA:538	3386	Clinical entity	Disorder	Lymphangioleiomyomatosis
ORPHA:747	3482	Clinical entity	Disorder	Autoimmune pulmonary alveolar proteinosis
ORPHA:122	8627	Clinical entity	Disorder	Birt-Hogg-Dubé syndrome
ORPHA:70578	10954	Clinical entity	Disorder	Adult acute respiratory distress syndrome
ORPHA:98300	13317	Clinical entity	Group of disorders	Idiopathic interstitial pneumonia
ORPHA:2032	7029	Clinical entity	Disorder	Idiopathic pulmonary fibrosis
ORPHA:1302	7034	Clinical entity	Disorder	Cryptogenic organizing pneumonia
ORPHA:79126	11153	Clinical entity	Disorder	Acute interstitial pneumonia
ORPHA:79127	11154	Clinical entity	Disorder	Respiratory bronchiolitis-interstitial lung disease syndrome
ORPHA:79128	11155	Clinical entity	Disorder	Lymphoid interstitial pneumonia
ORPHA:91364	12134	Clinical entity	Disorder	Non-specific interstitial pneumonia
ORPHA:98852	13869	Clinical entity	Disorder	Desquamative interstitial pneumonia
ORPHA:300564	21131	Clinical entity	Disorder	Combined pulmonary fibrosis-emphysema syndrome
ORPHA:494428	25673	Clinical entity	Disorder	Idiopathic pleuroparenchymal fibroelastosis
ORPHA:182098	18212	Clinical entity	Group of disorders	Pneumoconiosis
ORPHA:2302	865	Clinical entity	Disorder	Asbestos intoxication
ORPHA:133	1061	Clinical entity	Disorder	Chronic beryllium disease
ORPHA:210136	18687	Clinical entity	Disorder	Pulmonary fibrosis-hepatic hyperplasia-bone marrow hypoplasia syndrome
ORPHA:221043	18932	Clinical entity	Disorder	Hereditary fibrosing poikiloderma-tendon contractures-myopathy-pulmonary fibrosis syndrome
ORPHA:264745	20121	Clinical entity	Group of disorders	Secondary interstitial lung disease specific to adulthood associated with a systemic disease

ORPHA:797	735	Clinical entity	Disorder	Sarcoidosis
ORPHA:81	8611	Clinical entity	Disorder	Antisynthetase syndrome
ORPHA:264757	20123	Clinical entity	Group of disorders	Interstitial lung disease in childhood and adulthood
ORPHA:264762	20124	Clinical entity	Group of disorders	Primary interstitial lung disease in childhood and adulthood
ORPHA:99931	14504	Clinical entity	Disorder	Idiopathic pulmonary hemosiderosis
ORPHA:264930	20125	Clinical entity	Group of disorders	Primary interstitial lung disease in childhood and adulthood due to alveolar structure disorder
ORPHA:217566	18863	Clinical entity	Disorder	Chronic respiratory distress with surfactant metabolism deficiency
ORPHA:440392	23403	Clinical entity	Disorder	Interstitial lung disease due to SP-C deficiency
ORPHA:264935	20126	Clinical entity	Group of disorders	Primary interstitial lung disease in childhood and adulthood due to alveolar vascular disorder
ORPHA:182101	18213	Clinical entity	Group of disorders	Idiopathic eosinophilic pneumonia
ORPHA:2902	7033	Clinical entity	Disorder	Idiopathic chronic eosinophilic pneumonia
ORPHA:724	8741	Clinical entity	Disorder	Idiopathic acute eosinophilic pneumonia
ORPHA:264944	20127	Clinical entity	Group of disorders	Secondary interstitial lung disease in childhood and adulthood
ORPHA:99930	14503	Clinical entity	Disorder	Secondary pulmonary hemosiderosis
ORPHA:99932	14505	Clinical entity	Subtype of disorder	Heiner syndrome
ORPHA:264949	20128	Clinical entity	Group of disorders	Secondary interstitial lung disease in childhood and adulthood associated with a systemic disease
ORPHA:375	747	Clinical entity	Disorder	Anti-glomerular basement membrane disease
ORPHA:182104	18214	Clinical entity	Group of disorders	Secondary interstitial lung disease in childhood and adulthood associated with a connective tissue disease
ORPHA:809	3631	Clinical entity	Disorder	Mixed connective tissue disease
ORPHA:264968	20130	Clinical entity	Group of disorders	Secondary interstitial lung disease in childhood and adulthood associated with a metabolic disease
ORPHA:646	853	Clinical entity	Disorder	Niemann-Pick disease type C
ORPHA:216972	18801	Clinical entity	Subtype of disorder	Niemann-Pick disease type C, severe perinatal form
ORPHA:216975	18802	Clinical entity	Subtype of disorder	Niemann-Pick disease type C, severe early infantile neurologic onset
ORPHA:216978	18803	Clinical entity	Subtype of disorder	Niemann-Pick disease type C, late infantile neurologic onset
ORPHA:216981	18804	Clinical entity	Subtype of disorder	Niemann-Pick disease type C, juvenile neurologic onset
ORPHA:216986	18805	Clinical entity	Subtype of disorder	Niemann-Pick disease type C, adult neurologic onset
ORPHA:77259	11102	Clinical entity	Subtype of disorder	Gaucher disease type 1
ORPHA:264973	20131	Clinical entity	Group of disorders	Secondary interstitial lung disease in childhood and adulthood associated with a systemic vasculitis
ORPHA:91138	12115	Clinical entity	Disorder	Cryoglobulinemic vasculitis
ORPHA:93554	12396	Clinical entity	Subtype of disorder	Mixed cryoglobulinemia type II
ORPHA:93555	12397	Clinical entity	Subtype of disorder	Mixed cryoglobulinemia type III
ORPHA:156152	17099	Clinical entity	Group of disorders	Anti-neutrophil cytoplasmic antibody-associated vasculitis
ORPHA:183	745	Clinical entity	Disorder	Eosinophilic granulomatosis with polyangiitis
ORPHA:727	753	Clinical entity	Disorder	Microscopic polyangiitis
ORPHA:900	759	Clinical entity	Disorder	Granulomatosis with polyangiitis
ORPHA:264984	20133	Clinical entity	Group of disorders	Exposure-related interstitial lung disease
ORPHA:31740	9794	Clinical entity	Group of disorders	Hypersensitivity pneumonitis
ORPHA:99907	14480	Clinical entity	Disorder	House allergic alveolitis
ORPHA:99909	14482	Clinical entity	Group of disorders	Occupational allergic alveolitis
ORPHA:99906	14479	Clinical entity	Disorder	Farmer's lung disease
ORPHA:99908	14481	Clinical entity	Disorder	Pigeon-breeder lung disease
ORPHA:264978	20132	Clinical entity	Disorder	Drug or radiation exposure-related interstitial lung disease
ORPHA:420259	23102	Clinical entity	Disorder	Secondary pulmonary alveolar proteinosis
ORPHA:217080	18823	Clinical entity	Disorder	Pulmonary fungal infections in patients deemed at risk
ORPHA:228426	19132	Clinical entity	Disorder	Syndromic multisystem autoimmune disease due to Itch deficiency
ORPHA:247522	19523	Clinical entity	Disorder	Primary ciliary dyskinesia-retinitis pigmentosa syndrome
ORPHA:293987	20900	Clinical entity	Disorder	Rapid-onset childhood obesity-hypothalamic dysfunction-hypoventilation-autonomic dysregulation syndrome
ORPHA:330012	21958	Clinical entity	Disorder	High altitude pulmonary edema
ORPHA:411703	23030	Clinical entity	Disorder	Pulmonary non-tuberculous mycobacterial infection
ORPHA:439881	23394	Clinical entity	Disorder	Plastic bronchitis
ORPHA:449266	23612	Clinical entity	Disorder	Pleural empyema
ORPHA:454836	23665	Clinical entity	Disorder	Avian influenza
ORPHA:505248	26330	Clinical entity	Disorder	Mucopolysaccharidosis-like syndrome with congenital heart defects and hematopoietic disorders
ORPHA:505395	26354	Clinical entity	Disorder	Ventilator-induced diaphragmatic dysfunction
ORPHA:576074	28838	Clinical entity	Disorder	Middle East respiratory syndrome
ORPHA:289877	20771	Clinical entity	Disorder	Transient hyperammonemia of the newborn

MALADIES RESPIRATOIRES RARES

ORPHAcode	Disease ID	Nature	Classification Level	
ORPHA:97955	12974	Clinical entity	Group of disorders	Rare respiratory disease
ORPHA:3347	2967	Clinical entity	Disorder	Mounier-Kühn syndrome
ORPHA:60025	10797	Clinical entity	Disorder	Pulmonary alveolar microlithiasis
ORPHA:63999	10831	Clinical entity	Disorder	IgG4-related mediastinitis

TUMEURS RESPIRATOIRES RARES

ORPHAcode	Disease ID	Nature	Classification Level	Orphanet classification of rare respiratory diseases (Pat)
ORPHA:98060	13078	Clinical entity	Group of disorders	Rare respiratory tumor
ORPHA:150	1063	Clinical entity	Disorder	Nasopharyngeal carcinoma
ORPHA:101945	14881	Clinical entity	Group of disorders	Rare bronchopulmonary tumor
ORPHA:2420	752	Clinical entity	Disorder	Primary pulmonary lymphoma
ORPHA:50251	10645	Clinical entity	Disorder	Pleural mesothelioma
ORPHA:64741	10845	Clinical entity	Disorder	Pulmonary blastoma
ORPHA:64742	10846	Clinical entity	Disorder	Pleuropulmonary blastoma
ORPHA:99933	14506	Clinical entity	Subtype of disorder	Pleuropulmonary blastoma type 1
ORPHA:99934	14507	Clinical entity	Subtype of disorder	Pleuropulmonary blastoma type 2
ORPHA:99935	14508	Clinical entity	Subtype of disorder	Pleuropulmonary blastoma type 3
ORPHA:284343	20598	Clinical entity	Subtype of disorder	Pleuropulmonary blastoma familial tumor susceptibility syndrome
ORPHA:284362	20601	Clinical entity	Subtype of disorder	Fetal lung interstitial tumor
ORPHA:70573	10953	Clinical entity	Disorder	Small cell lung cancer
ORPHA:97287	12881	Clinical entity	Disorder	Bronchial neuroendocrine tumor
ORPHA:284395	20604	Clinical entity	Disorder	Well-differentiated fetal adenocarcinoma of the lung
ORPHA:466962	24185	Clinical entity	Disorder	SMARCA4-deficient sarcoma of thorax