Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Orphacode (Orphanet)	Toelichting erkenning
AMC	Achalasia Center, part of Esophageal Center Amsterdam	Sporadic achalasia	ORPHA:930	
AMC	AMC Pulmonary Hypertension Center	Eisenmenger syndrome	ORPHA:97214	
AMC	Amsterdam Centre of congenital malformations	Esophageal atresia	ORPHA:1199	
AMC	Amsterdam Centre of congenital malformations	Chronic intestinal pseudoobstruction	ORPHA:2978	
AMC	Amsterdam Centre of congenital malformations	Sacrococcygeal teratoma	ORPHA:494421	
AMC	Amsterdam Centre of congenital malformations	Anorectal malformation	ORPHA:96346	
AMC	Amsterdam expert center for bronchopulmonary dysplasia	Bronchopulmonary dysplasia	ORPHA:70589	
AMC	Amsterdam Expert Center for Developmental Disorders	Cornelia de Lange syndrome	ORPHA:199	
AMC	Amsterdam Expert Center for Developmental Disorders	Pitt-Hopkins syndrome	ORPHA:2896	
AMC	Amsterdam Expert Center for Developmental Disorders	Marshall-Smith syndrome	ORPHA:561	
AMC	Amsterdam Expert Center for Developmental Disorders	Rubinstein-Taybi syndrome	ORPHA:783	
AMC	Amsterdam Lysosome Center ("Sphinx")	Fabry disease	ORPHA:324	
AMC	Amsterdam Lysosome Center ("Sphinx")	Gaucher disease	ORPHA:355	
AMC	Amsterdam Lysosome Center ("Sphinx")	Mucopolysaccharidosis type 1	ORPHA:579	
AMC	Amsterdam Lysosome Center ("Sphinx")	Mucopolysaccharidosis type 3	ORPHA:581	
AMC	Amsterdam Lysosome Center ("Sphinx")	Mucopolysaccharidosis type 4	ORPHA:582	
AMC	Amsterdam Lysosome Center ("Sphinx")	Niemann-Pick disease type C	ORPHA:646	
AMC	Amsterdam Lysosome Center ("Sphinx")	Lysosomal Disease	ORPHA:68366	
AMC	Amsterdam Lysosome Center ("Sphinx")	Cholesteryl ester storage disease	ORPHA:75234	
AMC	Amsterdam Lysosome Center ("Sphinx")	Niemann-Pick disease type A	ORPHA:77292	
AMC	Amsterdam Lysosome Center ("Sphinx")	Niemann-Pick disease type B	ORPHA:77293	
AMC	Amsterdam Multidisciplinary Lyme borreliosis Center	Lyme disease	ORPHA:91546	
AMC	CAHAL (Center for Congenital Heart Disease Amsterdam-Leiden, adult CHD)	Congenital heart malformation; adult congenital heart disease	ORPHA:88991	
AMC	Center for condylar hyperplasia	Condylaire hyperplasie	ORPHA:477781	
AMC	Center for congenital nevi	Large congenital melanocytic nevus	ORPHA:626	
AMC	Center for hematological immune diseases Amsterdam	Autoimmune hemolytic anemia (AIHA)	ORPHA:98375	
AMC	Center for Idiopathic Nephrotic Syndrome	Primary glomerular disease; Steroid -sensitive and steroid-resistant nephrotic syndrome, incl. congenital nephrotic syndrome, membrano proliferative glomerulonefritis	ORPHA:357502	voor idiopathic nephrotic syndrome
AMC	Center for immune-mediated and genetic cholestasis syndromes	Primary sclerosing cholangitis	ORPHA:171	
AMC	Center for immune-mediated and genetic cholestasis syndromes	Primary biliary cirrhosis	ORPHA:186	
AMC	Center for immune-mediated and genetic cholestasis syndromes	Crigler-Najjar syndrome type 1	ORPHA:79234	
AMC	Center for immune-mediated and genetic cholestasis syndromes	Crigler-Najjar syndrome type 2	ORPHA:79235	
AMC	Center for Osteochondral Defects of the Talus	Osteochondritis van tarsaal/metatarsaal bot	ORPHA:2054	
AMC	Center for Paediatric oncology	soft tissue sarcomas (rhabdomyosarcoma and non rhabdo myosarcoma	ORPHA:3394	voor soft tissue sarcomas
AMC	Center for Paediatric oncology	Osteosarcoma, incl. ewing sarcomas and langer hans cell bone lesions	ORPHA:668	voor osteosarcoma
AMC	Center for rare movement disorders	Primary orthostatic tremor	ORPHA:238606	
AMC	Center for rare movement disorders	Benign adult familial myoclonic epilepsy	ORPHA:86814	
AMC	Centre for Bone Marrow Failure	Congenital neutropenia	ORPHA:101987	
AMC	Centre for Bone Marrow Failure	Rare hematologic disease; incl. hematopoietic stem cell defects resulting in anemia, thrombocytopenia as comprised of medullar aplasia, rare deficiency or constitutively dyserythropoietic anemia, incl Blackfan Diamond, congenital amegakaryocytic thrombocytopenia	ORPHA:68383	voor rare constitutional medullar aplasia
AMC	Centre for Bone Marrow Failure	Shwachman Diamond disease	ORPHA:811	
AMC	Centre for gastroenteropancreatic neuroendocrine tumors	Gastroenteropancreatic endocrine tumor	ORPHA:100092	
AMC	Centre for Genetic Metabolic Diseases Amsterdam	Disorder of phenylalanine metabolism	ORPHA:284814	
AMC	Centre for Genetic Metabolic Diseases Amsterdam	Disorder of galactose metabolism	ORPHA:308467	
AMC	Centre for graft versus host disease Amsterdam	Graft versus host disease (acuut en chronisch)	ORPHA:39812	
AMC	Centre for Hereditary Angioedema	Hereditary angioedema	ORPHA:91378	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Genetic cardiac rhythm disease	ORPHA:101934	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Hypertrophic cardiomyopathy	ORPHA:217569	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Familial dilated cardiomyopathy	ORPHA:217607	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Familial isolated arrhythmogenic right ventricular dysplasia	ORPHA:217656	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Familial dilated cardiomyopathy with conduction defect due to LMNA mutation	ORPHA:300751	
AMC	Centre for Hereditary cardiac rhythm and function disorders	Peripartum cardiomyopathy	ORPHA:563	
AMC	Centre for Immunodeficiencies	Rare immune disease; incl. primary immunodeficiencies	ORPHA:101997	
AMC	Centre for Kawasaki Disease	Kawasaki disease	ORPHA:2331	
AMC	Centre for Marfan syndrome	Marfan syndrome	ORPHA:558	
AMC	Centre for Neuromuscular Diseases	Hereditary motor and sensory neuropathy	ORPHA:166	
AMC	Centre for Neuromuscular Diseases	Poliomyelitis	ORPHA:2912	
AMC	Centre for Neuromuscular Diseases	Chronic inflammatory demyelinating polyneuropathy, incl. Guillain_Barre syndrome, CIDP, MMN	ORPHA:2932 ORPHA:2103 ORPHA:641	
AMC	Centre for Neuromuscular Diseases	Neuromuscular disease	ORPHA:68381	
AMC	Centre for Neuromuscular Diseases	Motor neuron disease; amyotrophic lateral sclerosis, primary sclerosis and progressive muscular atrophy	ORPHA:803 ORPHA:35689 ORPHA:454706	
AMC	Centre for Neuromuscular Diseases	Idiopathic inflammatory myopathy, incl dermatomyositis, polymyositis, necrotizing autoimmune myopathy and inclusion body myositis	ORPHA:98482	
AMC	Centre for Pediatric Rheumatic Diseases	Rare rheumatologic disease; focus on all forms of juvenile idiopathic arthritis, reactive arthritis and all forms of autoinflammatory syndromes	ORPHA:92	voor JIA
AMC	Centre for pediatric thromboembolic events	Rare thrombotic disease of hematologic origin	ORPHA:182054	
AMC	Centre for rare hypothalamic and pituitary diseases	Rare hypothalamic or pituitary disease	ORPHA:181384	
AMC	Centre for rare thyroid diseases	Congenital hypothyroidism	ORPHA:442	
AMC	Centre for Sickle Cell Disease	Hemoglobinopathy; incl Sickle cell disease and alfa or beta thalassemia	ORPHA:68364	
AMC	Centre for Upper GI tumors Amsterdam	Gastro-esophageal tumor	ORPHA:180821	
AMC	Dutch Centre for Peroxisomal disorders	Rhizomelic chondrodysplasia punctata	ORPHA:177	
AMC	Dutch Centre for Peroxisomal disorders	Disorder of peroxisomal alpha- - beta- and omega-oxidation	ORPHA:309810	
AMC	Dutch Centre for Peroxisomal disorders	Peroxisome biogenesis disorder-Zellweger syndrome spectrum	ORPHA:79189	
AMC	Dutch Centre for Peroxisomal disorders	Non-syndromic pontocerebellar hypoplasia	ORPHA:98523	
AMC	Expert Centre for congenital anomalies of the urinary tract EKZ-AMC	Non-syndromic renal or urinary tract malformation; CAKUT	ORPHA:93546	
AMC	Expertise Center Clinical Imunology and Rheumatology-Vasculitis	Vasculitis	ORPHA:52759	
AMC	Expertise center for genetic tumors of the digestive tract	Hyperplastic polyposis syndrome	ORPHA:157798	
AMC	Expertise center for genetic tumors of the digestive tract	Hereditary nonpolyposis colon cancer	ORPHA:443909	
AMC	Expertise center for genetic tumors of the digestive tract	Familial adenomatous polyposis	ORPHA:733	
AMC	Expertise center Vascular medicine	Familial lipoprotein lipase deficiency	ORPHA:309015	
AMC	Expertise center Vascular medicine	Tangier disease	ORPHA:31150	
AMC	Expertise center Vascular medicine	Homozygous familial hypercholesterolemia	ORPHA:391665	
AMC	Gastro-Intestinal Oncology Centre Amsterdam	Rare hepatic and biliary tract tumor; incl gallbladder tumors, Ampulla of Vater carcinoma	ORPHA:101943 ORPHA:300557	
AMC	Gastro-Intestinal Oncology Centre Amsterdam	Pancreatic tumor	ORPHA:424053 ORPHA:424073	voor zeldzame cysteuze pancreas tumoren
AMC	Hemophilia Comprehensive Care Treatment Centre	Rare hemorrhagic disorder due to a coagulation factors defect; hemophilia	ORPHA:448	
AMC	Hemophilia Comprehensive Care Treatment Centre	Rare hemorrhagic disorder due to a coagulation factors defect; von Willebrand Disease	ORPHA:903	
AMC	Intestinal Failure Unit	Short bowel syndrome, also secondary	ORPHA:104008	
AMC	Intestinal Failure Unit	Chronic intestinal failure	ORPHA:294422	
AMC	LYMMCARE Amsterdam (Lymphoma and Myeloma CAre and REsearch)	Lymphoma	ORPHA:223735	

AMC	LYMMCARE Amsterdam (Lymphoma and Myeloma CAre and REsearch)	Multiple myeloma	ORPHA:29073	
AMC	LYMMCARE Amsterdam (Lymphoma and Myeloma CAre and REsearch)	Waldenstrom macroglobulinemia	ORPHA:33226	
AMC	LYMMCARE Amsterdam (Lymphoma and Myeloma CAre and REsearch)	B-cell chronic lymphocytic leukemia	ORPHA:67038	
AMC	Melioidosis Expertise Center	Melioidosis	ORPHA:31202	
AMC	National centre for primary hyperoxaluria	Primary hyperoxaluria	ORPHA:416	
AMC	Solvent Team	Rare intoxication; Chronic Toxic Encephalopathy	ORPHA:108999	voor rare intoxication
AMC	Solvent Team	Lead poisoning; and other orphanet rare intoxications incl. different poisonings		
AMC	Solvent Team	Manganese poisoning		
AMC	Vascular malformations and hemangiomas centre	Diffuse lymphatic malformation	ORPHA:141209	
AMC	Vascular malformations and hemangiomas centre	Mucocutaneous venous malformations	ORPHA:2451	
AMC	Vascular malformations and hemangiomas centre	Dandy-Walker malformation - facial hemangioma; all subs	ORPHA:42775	
AMC	Vascular malformations and hemangiomas centre	Familial multiple nevi flammei	ORPHA:624	
AMC & VUmc	Amsterdam Center for ILD and sarcoïdosis	Sarcoidosis	ORPHA:797	
AMC & VUmc	Amsterdam Centre of congenital malformations	Hirschsprung Disease	ORPHA:388	
AMC & VUmc	Cystic Fibrosis Centre Amsterdam	Cystic fibrosis	ORPHA:586	
AMC & VUmc & LUMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Centrale sereuze chorioretinopathie	ORPHA:443079	
AMC & VUmc & LUMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	ORPHA:71862	
Bartiméus	Bartiméus Diagnostisch Centrum	Congenital stationary night blindness	ORPHA:215	
Bartiméus	Bartiméus Diagnostisch Centrum	Cerebral visual impairment (in children)	ORPHA:447788	
Bartiméus	Bartiméus Diagnostisch Centrum	Genetic vitreous-retinal disease	ORPHA:98657	
Bartiméus	Bartiméus Diagnostisch Centrum	Oculocutaneous or ocular albinism	ORPHA:98706	
Erasmus MC	Academic Breast Cancer Center	Rare tumor; Hereditary Breast Cancer	ORPHA:227535	
Erasmus MC	Academic Center Kidney & Hypertension	familial cystic renal disease	ORPHA:93587	
Erasmus MC	Academic Center Kidney & Hypertension	rare renal tubular disease	ORPHA:93603	
Erasmus MC	Brain Tumor Center	Glial tumor	ORPHA:182067	
Erasmus MC	Brain Tumor Center	Embryonal tumor of the neuroepithelial tissue	ORPHA:251852	
Erasmus MC	Brain Tumor Center	Primary germ cell tumor of the central nervous system	ORPHA:251995	
Erasmus MC	Brain Tumor Center	Tumor of the meninges	ORPHA:252025	
Erasmus MC	Brain Tumor Center	Hemangioblastoma	ORPHA:252054	
Erasmus MC	Brain Tumor Center	Tumor of cranial and spinal nerves	ORPHA:252057	
Erasmus MC	Brain Tumor Center	Primary central nervous system lymphoma	ORPHA:46135	
Erasmus MC	Brain Tumor Center	Craniopharyngioma	ORPHA:54595	
Erasmus MC	Center for Bronchopulmonary Dysplasia	Bronchopulmonary dysplasia	ORPHA:70589	
Erasmus MC	Center for congenital heart diseases Erasmus MC Rotterdam	Congenital heart malformation: intervention for diseases of D-2-8, D-6-4, D-6-11, D-6-10, D-6-3; A-5-7	ORPHA:88991	
Erasmus MC	Center for Familian and Hereditary Tumors	Lynch Syndrome	ORPHA:144	
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Achromatopsia	ORPHA:49382	
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Retinal dystrophy	ORPHA:71862	
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Retinitis pigmentosa	ORPHA:791	
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Developmental defect of the eye	ORPHA:98553	
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Color-vision disease	ORPHA:98658	
Erasmus MC	Center for inherited cardiovascular diseases	Hypertrophic cardiomyopathy	ORPHA:217569	
Erasmus MC	Center for inherited cardiovascular diseases	Unclassified cardiomyopathy	ORPHA:217678	
Erasmus MC	Center for inherited cardiovascular diseases	Rare cardiac disease; rare familial occurrence of thoracic aortic abnormalities incl. dissection with (un)known genetic cause, e.g. Aneurysm-osteoarthritis syndrome	ORPHA:285014	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Neuronal ceroid lipofuscinosis	ORPHA:216	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Glycoproteinosis	ORPHA:309279	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Disorder of lysosomal amino acid transport	ORPHA:79207	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Mucopolysaccharidosis	ORPHA:79213	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Sphingolipidosis	ORPHA:79225	
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Primary bone dysplasia with defective bone mineralization	ORPHA:93447	
Erasmus MC	Center for Neuro-inflammatory disorders	Limbic encephalitis	ORPHA:163892	
Erasmus MC	Center for Neuro-inflammatory disorders	Inflammatory and autoimmune disease with epilepsy	ORPHA:166484	
Erasmus MC	Center for Neuro-inflammatory disorders	CLIPPERS	ORPHA:284448	
Erasmus MC	Center for Neuro-inflammatory disorders	Paraneoplastic neurologic syndrome	ORPHA:36388	
Erasmus MC	Center for Neuro-inflammatory disorders	Morvan syndrome	ORPHA:83467	
Erasmus MC	Center for Neuro-inflammatory disorders	Isaac syndrome	ORPHA:84142	
Erasmus MC	Center for Neuro-inflammatory disorders	Postinfectious encephalitis	ORPHA:98253	
Erasmus MC	Center for pediatric laryngotracheal stenosis	Congenital subglottic stenosis	ORPHA:141121	
Erasmus MC	Center for pediatric laryngotracheal stenosis	Congenital tracheal stenosis	ORPHA:141127	
Erasmus MC	Center for pediatric laryngotracheal stenosis	Larynго-tracheo-esophageal cleft	ORPHA:2004	
Erasmus MC	Center for Perinatal Psychiatry	Rare disorder related with pregnancy - childbirth and puerperium; Postpartum psychosis	ORPHA:443173	
Erasmus MC	Center for PKU, urea cycle disorders and organic acidurias	Disorder of urea cycle metabolism and ammonia detoxification	ORPHA:79167	
Erasmus MC	Center for PKU, urea cycle disorders and organic acidurias	Disorder of branched-chain amino acid metabolism	ORPHA:79197	
Erasmus MC	Center for pregnancy induced diseases	HELLP-syndrome	ORPHA:244242	
Erasmus MC	Center for pregnancy induced diseases	Pre-eclampsia (< 34 wks)	ORPHA:275555	
Erasmus MC	Center for Rare Systemic Immune Disease	Rare systemic disease; Uveitis, Morbus Behcet, Morbus Sjogren, Systemic sclerosis	ORPHA:98715 ORPHA:117 ORPHA:289390 ORPHA:90291	
Erasmus MC	Center for systemic allergic diseases	Rare immune disease; Systemic mastocytosis	ORPHA:2467	
Erasmus MC	Center of rare skin diseases	Systemic disease with skin involvement; Localized scleroderma and Suppurative hidradenitis	ORPHA:387	voor Supp. hidradenitis
Erasmus MC	Center of rare skin diseases	Netherton syndrome	ORPHA:634	
Erasmus MC	Center of rare skin diseases	Vascular anomaly or angioma	ORPHA:68419	
Erasmus MC	Centre of expertise for Children with Autoimmune Diseases	Juvenile idiopathic arthritis	ORPHA:92	
Erasmus MC	Centre of Expertise for Uveitis	Uveitis	ORPHA:98715	
Erasmus MC	Centre of Oligodontia	Oligodontia	ORPHA:99798	
Erasmus MC	Craniofacial Center	Treacher-Collins syndrome; incl. Nager and Burn-McKeown syn	ORPHA:138050	voor Pierre Robin syndrome associated with branchial archs anomalies
Erasmus MC	Craniofacial Center	Isolated craniosynostosis	ORPHA:139390	
Erasmus MC	Craniofacial Center	Syndromic craniosynostosis; craniofrontonasal syndrome	ORPHA:139393 ORPHA:1520	
Erasmus MC	Craniofacial Center	Cleft lip with or without cleft palate	ORPHA:1991	
Erasmus MC	Craniofacial Center	Cleft palate	ORPHA:2014	
Erasmus MC	Craniofacial Center	Acalvaria; including ossification defects of the skull with or without involvement of skin		
Erasmus MC	Craniofacial Center	Cleft palate; incl cleft palate only and Pierre Robin sequence (eg Stickler, van der Woude, Wolf Hirschhorn)		
Erasmus MC	Craniofacial Center	Facial cleft		
Erasmus MC	Craniofacial Center	Goldenhar syndrome; incl. hemifacial microsomia		
Erasmus MC	Craniofacial Center	Rare bone development disorder; involving craniofacial presentation of fibrous dysplasia, McCune Albright, cherubism, cleidocranial dysplasia		
Erasmus MC	Dutch Porphyria Center	Porphyria	ORPHA:738	
Erasmus MC	Dutch Porphyria Center	Erythropoietic protoporphyria	ORPHA:79278 ORPHA:443197	
Erasmus MC	Dutch Porphyria Center	Acute hepatic porphyria	ORPHA:95157	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Cardiofaciocutaneous syndrome	ORPHA:1340	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Costello syndrome	ORPHA:3071	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Sturge-Weber syndrome	ORPHA:3205	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Neurofibromatosis type 1	ORPHA:636	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Angelman syndrome	ORPHA:72	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Tuberous sclerosis	ORPHA:805	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Fragile X syndrome	ORPHA:908	
Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Central nervous system malformation	ORPHA:98044	
Erasmus MC	Erasmus MC Bone Center	Primary bone dysplasia; incl. a.o. list of 6 specific forms of dysplasia	ORPHA:364526	voor primary bone dysplasia

Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Pancreatic tumor (2x)	ORPHA:1333	voor familiair pancreascarc.
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Autoimmune pancreatitis type 1	ORPHA:280302	
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Autoimmune pancreatitis type 2	ORPHA:280315	
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Congenital pancreatic cyst	ORPHA:313906	
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Recurrent acute pancreatitis	ORPHA:64740	
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Hereditary chronic pancreatitis	ORPHA:676	
Erasmus MC	Erasmus MC Cystic Fibrosis Center	Cystic fibrosis	ORPHA:586	
Erasmus MC	Erasmus MC Leprosy Centre	Leprosy	ORPHA:548	
Erasmus MC	Erasmus MC Liver Center	Rare hepatic and biliary tract tumor	ORPHA:101943	
Erasmus MC	Erasmus MC Liver Center	Adult hepatocellular carcinoma	ORPHA:210159	
Erasmus MC	Erasmus MC Liver Center	Hepatocellular adenoma	ORPHA:54272	
Erasmus MC	Erasmus MC Liver Center	Klatskin tumor	ORPHA:99978	
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Multiple congenital anomalies/dysmorphic syndrome-intellectual disability; Pre-axiale polydactylie, Split hand and foot, Syndactyly	ORPHA:109009	voor Syndrome with limb malformations as a major feature
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Non-syndromic limb malformation	ORPHA:109011	voor Non-syndromic limb malformation
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Dysostosis with limb anomaly as a major feature		
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Multiple congenital anomalies/dysmorphic syndrome without intellectual disability; Radiusdysplasie (Fanconi, TAR etc), Brachydactylie, Holt-Oram/Hart-Hand		
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Non-syndromic limb malformation; Macrodactylie handen voeten geïsoleerd, Hemihypertrofie, Musculaire hyperplasie		
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Syndrome with limb malformations as a major feature; Arthrogryposis, Syndromale polydactylie (o.a. Greig) Transversaal reductie defect		
Erasmus MC	ErasmusMC centre for endocrine disorders	Rare adrenal disease	ORPHA:101954	
Erasmus MC	ErasmusMC centre for endocrine disorders	Rare thyroid disease	ORPHA:101955	
Erasmus MC	ErasmusMC centre for endocrine disorders	Rare hypothalamic or pituitary disease	ORPHA:181384	
Erasmus MC	ErasmusMC centre for endocrine disorders	Endocrine tumor	ORPHA:877	
Erasmus MC	Expert Center Prader Willi syndrome	Chromosomal anomaly; Prader Willi syndrome	ORPHA:739	
Erasmus MC	Expert Center Rare Growth Disorders	Growth hormone insensitivity syndrome	ORPHA:181393	
Erasmus MC	Expert Center Rare Growth Disorders	Silver-Russell syndrome due to maternal uniparental disomy of chromosome 7; also H19 hypomethylation chromosome 11 or unknown	ORPHA:813	
Erasmus MC	Expert Center Rare Growth Disorders	Non-acquired pituitary hormone deficiency; due to known GH gene and yet unknown genetic variants	ORPHA:95488	
Erasmus MC	Expert Center Rare Growth Disorders	Non-acquired pituitary hormone deficiency; not yet known whether or which genes	ORPHA:95488	
Erasmus MC	Expert Center spinal disraphism Rotterdam	Total spina bifida aperta; and occult spinal disraphism and complex congenital anomalies such as VACTERL association or sacrococcygeal disorders	ORPHA:268369	voor spina bifida aperta.
Erasmus MC	Expertise center DSD	Disorder of sex development	ORPHA:90771	
Erasmus MC	Expertise center Erasmus MC Vascular Genetics	Familial lipoprotein lipase deficiency	ORPHA:309015	
Erasmus MC	Expertise center Erasmus MC Vascular Genetics	Homozygous familial hypercholesterolemia	ORPHA:391665	
Erasmus MC	Expertise Center Rare hemorrhagic disorders (=Hemophilia treatment center, kind)	Rare hemorrhagic disorder due to a coagulation factors defect, incl FXIII, FXI, FVII,FV deficiency and alpha2-antiplasmin def.	ORPHA:248315	
Erasmus MC	Expertise Center Rare hemorrhagic disorders (=Hemophilia treatment center, kind)	Rare hemorrhagic disorder due to a platelet anomaly	ORPHA:248326	
Erasmus MC	Expertise Center Rare hemorrhagic disorders (=Hemophilia treatment center, kind)	Hemophilia	ORPHA:448	
Erasmus MC	Expertise Center Rare hemorrhagic disorders (=Hemophilia treatment center, kind)	Von Willebrand disease	ORPHA:903	
Erasmus MC	Hemophilia treatment center (volwassen)	Rare hemorrhagic disorder due to a coagulation factors defect, incl FXIII, FXI, FVII,FV deficiency and alpha2-antiplasmin def.	ORPHA:248315	
Erasmus MC	Hemophilia treatment center (volwassen)	Rare hemorrhagic disorder due to a platelet anomaly	ORPHA:248326	
Erasmus MC	Hemophilia treatment center (volwassen)	Hemophilia	ORPHA:448	
Erasmus MC	Hemophilia treatment center (volwassen)	Von Willebrand disease	ORPHA:903	
Erasmus MC	Immunodeficiency center	Primary immunodeficiency (*)	ORPHA:101997	
Erasmus MC	Interstitial Lung Disease Centre	Interstitial lung disease; adult	ORPHA:2032	
Erasmus MC	Leukemia and Stem cell transplantation center	Myeloid hemopathy	ORPHA:171895	
Erasmus MC	Leukemia and Stem cell transplantation center	Acute lymphoblastic leukemia (adult)	ORPHA:513	
Erasmus MC	Mesothelioma centre	Malignant peritoneal mesothelioma	ORPHA:168811	
Erasmus MC	Mesothelioma centre	Mesothelioma	ORPHA:50251	
Erasmus MC	Mesothelioma centre	Small cell lung cancer	ORPHA:70573	
Erasmus MC	Mesothelioma centre	Thymoma	ORPHA:99867	
Erasmus MC	MS center	Multiple sclerosis variant	ORPHA:228145	
Erasmus MC	MS center	Neuromyelitis optica	ORPHA:71211	
Erasmus MC	Multiple myeloma treatment center	Multiple myeloma	ORPHA:29073	
Erasmus MC	Multiple myeloma treatment center	Aggressive B-cell non-Hodgkin lymphoma	ORPHA:300846	
Erasmus MC	Multiple myeloma treatment center			
Erasmus MC	Nasal, Paranasal, Ear and Skull base tumors workgroup	Esthesioneuroblastoma	ORPHA:1957	
Erasmus MC	Nasal, Paranasal, Ear and Skull base tumors workgroup	Squamous cell carcinoma of head and neck; tumors arising from the epithelium of the nasal, paranasal and skull base regions	ORPHA:67037	voor tumors arising from the epithelium of the nasal, paranasal and skull base regions
Erasmus MC	NeMo, expert centre for Neuromuscular and Mitochondrial Diseases	Mitochondrial disease	ORPHA:68380	
Erasmus MC	Neuromuscular Center Erasmus MC	Guillain-Bar syndrome	ORPHA:2103	
Erasmus MC	Neuromuscular Center Erasmus MC	Chronic inflammatory demyelinating polyneuropathy	ORPHA:2932	
Erasmus MC	Neuromuscular Center Erasmus MC	Neuromuscular disease	ORPHA:68381	
Erasmus MC	Paediatric Brain Tumour Center	Rare nervous system tumor	ORPHA:98062	
Erasmus MC	Paediatric Brain Tumour Center			
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Esophageal atresia	ORPHA:1199	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Congenital and syndromic diaphragmatic hernia	ORPHA:2140	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Gastroschisis	ORPHA:2368	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Chronic intestinal failure	ORPHA:294422	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Hirschsprung disease	ORPHA:388	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Omphalocele	ORPHA:660	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Anorectal malformation	ORPHA:96346	
Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Intestinal malformation	ORPHA:97945	
Erasmus MC	Pick Centrum	Semantic dementia	ORPHA:100069	
Erasmus MC	Pick Centrum	Progressive non-fluent aphasia	ORPHA:100070	
Erasmus MC	Pick Centrum	Classical progressive supranuclear palsy	ORPHA:240071	
Erasmus MC	Pick Centrum	Behavioral variant of frontotemporal dementia	ORPHA:275864	
Erasmus MC	Pick Centrum	Frontotemporal dementia with motor neuron disease	ORPHA:275872	
Erasmus MC	Pick Centrum	Corticobasal degeneration	ORPHA:454887	
Erasmus MC	Pick Centrum	Transmissible spongiform encephalopathy	ORPHA:56970	
Erasmus MC	Pick Centrum	Amyotrophic lateral sclerosis-parkinsonism-dementia complex	ORPHA:90020	
Erasmus MC	Pompe Center	Glycogen storage disease due to acid maltase deficiency - infantile onset	ORPHA:365	
Erasmus MC	Pompe Center	idem - juvenile onset	ORPHA:365	
Erasmus MC	Pompe Center	idem- adult onset	ORPHA:365	
Erasmus MC	Pulmonary hypertension center	Rare respiratory disease	ORPHA:71198	
Erasmus MC	Rotterdam Head and Neck Tumor work group	Squamous cell carcinoma of head and neck; EBV associated tumor, digestive tumor/Salivary gland tumor, Salivary gland tumor, Cervicofacial lymphatic malformation	ORPHA:98061	voor oral and laryngyal squamous carcinoma
Erasmus MC	Rotterdam Ocular Melanoma Center (ROMC)	Uveal melanoma	ORPHA:39044	
Erasmus MC	Rotterdam Oesophageal and Gastric Cancer Working Group	Esophageal carcinoma; incl. Barrett's oesophagus	ORPHA:70482	

Erasmus MC	Sarcoidosis Centre ErasmusMC	Sarcoidosis	ORPHA:797	
Erasmus MC	Sickle cell center, kind	Beta-thalassemia and related diseases	ORPHA:275749	
Erasmus MC	Sickle cell center, kind	Sickle cell disease and related diseases	ORPHA:275752	
Erasmus MC	Sickle cell center, kind	Hemoglobinopathy	ORPHA:68364	
Erasmus MC	Sickle cell center, volwassen	Alpha-thalassemia and related diseases	ORPHA:275745	
Erasmus MC	Sickle cell center, volwassen	Beta-thalassemia and related diseases	ORPHA:275749	
Erasmus MC	Sickle cell center, volwassen	Sickle cell disease and related diseases	ORPHA:275752	
Erasmus MC	Sickle cell center, volwassen			
Erasmus MC	Soft tissue sarcoma center	Rare soft tissue tumor	ORPHA:71209	
Erasmus MC	Turner Syndrome Center	Turner syndrome	ORPHA:881	
Erasmus MC & LUMC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Constitutional neutropenia	ORPHA:101987	
Erasmus MC & LUMC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Inherited acute myeloid leukemia	ORPHA:319465	
Erasmus MC & LUMC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Myelodysplastic syndrome	ORPHA:52688	
Kempenhaeghe	Center of Sleep Medicine Kempenhaeghe	Sleep disorders	ORPHA:68354	voor rare sleep disorder
LUMC	Alpha1 International Registry (AIR)	Alpha-1-antitrypsin deficiency	ORPHA:60	
LUMC	Autonomic Disease Center	Primary orthostatic hypotension, PAF, MSA	ORPHA:441	voor PAF
LUMC	Bone and soft tissue tumour clinic	Ewing sarcoma	ORPHA:319	
LUMC	Bone and soft tissue tumour clinic	Multiple osteochondromas	ORPHA:321	
LUMC	Bone and soft tissue tumour clinic	Giant cell tumor of bone	ORPHA:363976	
LUMC	Bone and soft tissue tumour clinic	Chondromyxoid fibroma	ORPHA:404507	
LUMC	Bone and soft tissue tumour clinic	Soft tissue sarcoma; Gastrointestinal stromal (cell) tumour	ORPHA:44890	
LUMC	Bone and soft tissue tumour clinic	Chondrosarcoma	ORPHA:55880	
LUMC	Bone and soft tissue tumour clinic	Adamantinoma	ORPHA:55881	
LUMC	Bone and soft tissue tumour clinic	Osteosarcoma	ORPHA:668	
LUMC	Bone and soft tissue tumour clinic	Rare bone tumor	ORPHA:68411	
LUMC	Bone and soft tissue tumour clinic	Rare soft tissue tumor	ORPHA:71209	
LUMC	Center for Bone Quality	Sternocostoclavicular Hyperostosis	ORPHA:178311	
LUMC	Center for Bone Quality	Fibrous dysplasia of bone	ORPHA:249	
LUMC	Center for Bone Quality	Sclerosteosis	ORPHA:3152	
LUMC	Center for Bone Quality	Chronic recurrent multifocal osteomyelitis	ORPHA:324964	
LUMC	Center for Bone Quality	Rare parathyroid disease and phosphocalcic metabolism anomaly	ORPHA:68415	
LUMC	Center for Bone Quality	Primary bone dysplasia with increased bone density	ORPHA:93444	
LUMC	Center for Bone Quality	Primary bone dysplasia with decreased bone density	ORPHA:93446	
LUMC	Center for Bone Quality	Primary bone dysplasia with defective bone mineralization	ORPHA:93447	
LUMC	Center for Bone Quality	Primary bone dysplasia with disorganized development of skeletal components	ORPHA:93450	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Acquired chronic primary adrenal insufficiency	ORPHA:101963	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Parathyroid carcinoma	ORPHA:143	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Thyroid tumor	ORPHA:146 ORPHA:142 ORPHA:1332 ORPHA:97290 ORPHA:319494	ORPHA:319494: zonder MEN type 2
LUMC	Center for Endocrine Tumors Leiden (CETL)	Adrenocortical carcinoma	ORPHA:1501	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Endocrine tumor with other location	ORPHA:181384	voor Rare hypothalamic and pituitary disease (behalve MEN 1 en 2)
LUMC	Center for Endocrine Tumors Leiden (CETL)	Hereditary pheochromocytoma-paraganglioma	ORPHA:29072	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Acromegaly		
LUMC	Center for Endocrine Tumors Leiden (CETL)	Cushing disease		
LUMC	Center for Endocrine Tumors Leiden (CETL)	Non-functioning pituitary adenoma		
LUMC	Center for Endocrine Tumors Leiden (CETL)	Pituitary deficiency		
LUMC	Center for Inherited kidney disease	Autosomal Dominant Medullairy Cystic Kidney Disease	ORPHA:34149	
LUMC	Center for Inherited kidney disease	Autosomal Dominant Polycystic Kidney Disease PKD2 mutation	ORPHA:730	
LUMC	Center for Inherited kidney disease	Autosomal Dominant Polycystic Kidney Disease, PKD1 mutation	ORPHA:730	
LUMC	Center for Inherited kidney disease	Autosomal recessive polycystic kidney disease	ORPHA:731	
LUMC	Center for Narcolepsia	Narcolepsy-cataplexy	ORPHA:2073	
LUMC	Center for Polyomavirus-associated skin infections and cancer (Acronym PASIC)	Virus-associated trichodysplasia spinulosa, usually known as trichodysplasia spinulosa	ORPHA:228379	
LUMC	Center for primary and secondary immunodeficiencies	Combined T and B cell immunodeficiency	ORPHA:101972	
LUMC	Center for primary and secondary immunodeficiencies	Mendelian susceptibility to mycobacterial diseases	ORPHA:748	
LUMC	Center for vestibular schwannoma and NF2	Vestibularis schwannoom	ORPHA:252175	
LUMC	Center of expertise Coffin-Siris syndrome	Coffin-Siris syndroom	ORPHA:1465	
LUMC	Cerebral Hereditary Angiopathy Center	CADASIL, and RVCL - HCHWA-D	ORPHA:136 ORPHA:247691 ORPHA:100006	
LUMC	Clinic for Lupus-, Vasculitis- and Complement-mediated systemic diseases	Anti-neutrophil cytoplasmic antibody-associated vasculitis	ORPHA:156152	
LUMC	Clinic for Lupus-, Vasculitis- and Complement-mediated systemic diseases	Immunoglobulin-mediated membranoproliferative glomerulonephritis	ORPHA:329903	
LUMC	Clinic for Lupus-, Vasculitis- and Complement-mediated systemic diseases	C3 glomerulonephritis	ORPHA:329931	
LUMC	Clinic for Lupus-, Vasculitis- and Complement-mediated systemic diseases	Pauci-immune glomerulonephritis	ORPHA:93126	
LUMC	Congenital Heart malformations (CAHAL pediatric)	Rare cardiac rhythm disease; non-genetic	ORPHA:218439	
LUMC	Congenital Heart malformations (CAHAL pediatric)	Congenital heart malformation; pediatric	ORPHA:88991	
LUMC	Expert center for aplastic anemia	Idiopathic aplastic anemia	ORPHA:88	
LUMC	Expert center for cutaneous lymphomas	Primary cutaneous lymphoma	ORPHA:542	
LUMC	Expert center for familial cutanous melanoma	Familial melanoma; incl. FAMMM syndr. and FAMMMPC syndr.	ORPHA:618 ORPHA:404560	
LUMC	Expert Center for hematopoietic stem cell transplantation and T-cell immunotherapy	Acute lymfoblastic leukemia	ORPHA:513	
LUMC	Expert Center for hematopoietic stem cell transplantation and T-cell immunotherapy	Acute myeloid leukemia	ORPHA:519	
LUMC	Expert Center for hematopoietic stem cell transplantation and T-cell immunotherapy	Plasma cell tumor	ORPHA:98282	
LUMC	Expert center for hemoglobinopathies	Hemoglobinopathy, incl alpha- beta-thalassemia, Sickle Cell Disease, HbS and Hb variants	ORPHA:68364	
LUMC	Expert center for pediatric stem cell transplantation	Combined T and B cell immunodeficiency, mainly SCID and ICF syndr	ORPHA:101972 ORPHA:2268	
LUMC	Expert center for pediatric stem cell transplantation	Primary immunodeficiency due to a defect in adaptive immunity	ORPHA:179006	
LUMC	Expert center for pediatric stem cell transplantation	Acute graft versus host disease	ORPHA:99920	
LUMC	Expertise Center Fetal medicine	Hydrops fetalis	ORPHA:1041	
LUMC	Expertise Center Fetal medicine	Non-syndromic respiratory or mediastinal malformation	ORPHA:108993	
LUMC	Expertise Center Fetal medicine	Hemolytic disease due to fetomaternal alloimmunization	ORPHA:275938	
LUMC	Expertise Center Fetal medicine	Fetal parvovirus syndrome	ORPHA:295	
LUMC	Expertise Center Fetal medicine	Fetal and neonatal alloimmune thrombocytopenia	ORPHA:853	
LUMC	Expertise Center Fetal medicine	Congenital heart malformation; fetal cardiac interventions	ORPHA:88991	
LUMC	Expertise Center Fetal medicine	Posterior urethral valve; Lower Urinary Tract Obstruction	ORPHA:93110	
LUMC	Expertise Center Fetal medicine	Twin to twin transfusion syndrome	ORPHA:95431	
LUMC	Expertise center for lipodystrophy	Berardinelli-Seip congenital lipodystrophy	ORPHA:90970	voor primary lipodystrophy
LUMC	Expertise center for lipodystrophy	Acquired generalized lipodystrophy, also called Lawrence-Seip		
LUMC	Expertise center for lipodystrophy	Familial partial lipodystrophy - bberling type		
LUMC	Expertise center for lipodystrophy	Familial partial lipodystrophy - Dunnigan type		
LUMC	Expertise center for lipodystrophy	Familial partial lipodystrophy associated with PLIN1 mutations		
LUMC	Expertise center for lipodystrophy	Familial partial lipodystrophy associated with PPARG mutations		
LUMC	Expertise center for lipodystrophy	Familial partial lipodystrophy due to AKT2 mutations		
LUMC	Expertise center for monogenic diabetes mellitus	MODY syndrome	ORPHA:552	
LUMC	Expertise Center Genetics of growth	Growth disorders with a height <-3 SDS	ORPHA:139021	

LUMC	Expertise Center Genetics of growth	Disorders in the GH-IGF1 axis and signaling pathways	ORPHA:181393	
LUMC	Expertise Center Genetics of growth	Leri-Weill dyschondrosteosis/syndrome	ORPHA:240	
LUMC	Expertise Center Genetics of growth	IGSF1 deficiency syndrome	ORPHA:329235	
LUMC	Expertise Center Genetics of growth	Growth disorders with a height >3 SDS	ORPHA:93460	
LUMC	Expertise Center Pediatric ophthalmology	Retinopathy of prematurity	ORPHA:90050	
LUMC	Expertise Center Rare autoinflammatory diseases	Juvenile rheumatoid factor-negative polyarthritis with anti-nuclear antibodies	ORPHA:247854	
LUMC	Expertise Center Rare autoinflammatory diseases	Systemic sclerosis	ORPHA:90291	
LUMC	Familial and hereditary breast cancer center	Hereditary breast and ovarian cancer syndrome	ORPHA:145	
LUMC	Familial and hereditary breast cancer center	Hereditary breast cancer	ORPHA:227535	
LUMC	Female Cancer Center - Leiden (FCC-L)	Rare ovarian cancer	ORPHA:213500	
LUMC	Female Cancer Center - Leiden (FCC-L)	Rare cancer of the corpus uteri	ORPHA:213569	
LUMC	Female Cancer Center - Leiden (FCC-L)	Rare cancer of the cervix uteri	ORPHA:213761	
LUMC	Female Cancer Center - Leiden (FCC-L)	Rare vulvovaginal tumor; Vulvar cancer	ORPHA:494418 ORPHA:137583	
LUMC	Gender Clinic Leiden, WAKZ-Curium LUMC	Rare endocrine disease, gender dysphoria	ORPHA:459690	voor gender dysphoria
LUMC	Headache Center LUMC	Rare headache; Sporadic/Familial Hemiplegic Migraine, Visual Snow, SUNCT, Cluster Headache, Trigeminus Neuralgia, Hemicrania Continua, Paroxysmal Hemicrania, TAC nao, Hypnic Headache.	ORPHA:569 ORPHA:420556 ORPHA:57145 ORPHA:221091 ORPHA:443070 ORPHA:157835 ORPHA:157843 ORPHA:276429	
LUMC	Hemophilia treatment centre LUMC-Haga	Acquired hemophilia	ORPHA:73274	
LUMC	Hemophilia treatment centre LUMC-Haga	Von Willebrand disease	ORPHA:903	
LUMC	Hemophilia treatment centre LUMC-Haga	Hemophilia A	ORPHA:98878	
LUMC	Hemophilia treatment centre LUMC-Haga	Hemophilia B	ORPHA:98879	
LUMC	Hemophilia treatment centre LUMC-Haga	Acquired von Willebrand syndrome	ORPHA:99147	
LUMC	Hereditairy bowel cancer centre	Hereditary nonpolyposis colon cancer; Lynch caused by MLH1 or MSH2 mutation	ORPHA:144	
LUMC	Hereditairy bowel cancer centre	Hereditary nonpolyposis colon cancer; Lynch caused by MSH6 or PMS2 mutation	ORPHA:144	
LUMC	Hereditairy bowel cancer centre	MUTYH-related attenuated familial adenomatous polyposis	ORPHA:247798	
LUMC	Hereditairy bowel cancer centre	Familial adenomatous polyposis; APC associated polyposis	ORPHA:733 ORPHA:247806	
LUMC	Huntington Disease Center Leiden	Huntington disease	ORPHA:399	
LUMC	Leiden Ocular Oncology Center	Uveal melanoma	ORPHA:39044	
LUMC	Marfan-FTAAD Clinic	Marfan syndrome; including neonatal Marfan syndrome, FTAAD	ORPHA:558 ORPHA:284979 ORPHA:91387	
LUMC	Nerve Centre	Rare neurologic disease, nerve lesion	ORPHA:439202	
LUMC	Neuromuscular Center LUMC	Skeletal muscle disease	ORPHA:262	voor Duchenne en Becker muscular dystrophy
LUMC	Neuromuscular Center LUMC	Facioscapulohumeral dystrophy	ORPHA:269	
LUMC	Neuromuscular Center LUMC	Oculopharyngeal muscular dystrophy	ORPHA:270	
LUMC	Neuromuscular Center LUMC	Inclusion body myositis	ORPHA:611	
LUMC	Neuromuscular Center LUMC	Neuromuscular disease	ORPHA:68381	
LUMC	Neuromuscular Center LUMC	Acquired neuromuscular junction disease	ORPHA:98494	
LUMC	Prenatal and congenital infections by cytomegalovirus and parvovirus B19	Infectious embryofetopathy; Congenital CMV inf. and fetal and congenital parvovirus B19 inf.	ORPHA:294 ORPHA:295	
LUMC & AMC & VUmc	Center for Hereditary Retinal Diseases Leiden Amsterdam	Centrale sereuze chorioretinopathie	ORPHA:443079	
LUMC & AMC & VUmc	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	ORPHA:71862	
LUMC & Erasmus MC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Constitutional neutropenia	ORPHA:101987	
LUMC & Erasmus MC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Inherited acute myeloid leukemia	ORPHA:319465	
LUMC & Erasmus MC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Myelodysplastic syndrome	ORPHA:52688	
MUMC+	Academic Center for Epilepsy	Epilepsy syndrome	ORPHA:166463	
MUMC+	Academic Center for Epilepsy	Early infantile epileptic encephalopathy	ORPHA:1934	
MUMC+	Academic Center for Epilepsy	Continuous spikes and waves during sleep, epileptic encephalopathy	ORPHA:725	
MUMC+	Center for Endocrine tumors	Endocrine tumor with other location; Papillary and follicular thyroid carcinoma, medullary thyroid carcinoma, anaplastic thyroid carcinoma	ORPHA:100088	voor thyroid carcinoma
MUMC+	Center for Genodermatoses	Birt-Hogg-Du syndrome	ORPHA:122	
MUMC+	Center for Genodermatoses	Inherited ichthyosis	ORPHA:183435	
MUMC+	Center for Genodermatoses	Basal cell nevus syndrome	ORPHA:377	
MUMC+	Center for Genodermatoses	Other sporadic genodermatoses	ORPHA:68346	voor rare genetic skin disease
MUMC+	Center for Genodermatoses	Erythrokeratoderma	ORPHA:79355	
MUMC+	Center for Genodermatoses	Hereditary palmoplantar keratoderma	ORPHA:79357	
MUMC+	Center for Genodermatoses	Lymphedema	ORPHA:79383	
MUMC+	Division of Balance Disorders	Idiopathic Bilateral Vestibulopathy	ORPHA:171684	
MUMC+	Expertise Center Hereditary Tumors	Hereditary breast and ovarian cancer	ORPHA:145	
MUMC+	Expertise center Cardiogenetics MUMC+	Brugada syndrome	ORPHA:130	
MUMC+	Expertise center Cardiogenetics MUMC+	Dilated cardiomyopathy	ORPHA:217604	
MUMC+	Expertise center Cardiogenetics MUMC+	Idiopathic ventricular fibrillation - not Brugada type	ORPHA:228140	
MUMC+	Expertise center Cardiogenetics MUMC+	Arrhythmogenic right ventricular dysplasia	ORPHA:247	
MUMC+	Expertise center Cardiogenetics MUMC+	Familial long QT syndrome	ORPHA:768	
MUMC+	Expertise center Cardiogenetics MUMC+	Rare familial disorder with hypertrophic cardiomyopathy	ORPHA:99739	
MUMC+	Expertise Center Cerebral Palsy	Spastic diplegia - infantile type. 1. spastic unilateral cerebral palsy 2. dyskinetic cerebral palsy		
MUMC+	Expertise Center Galactosemia	Galactosemia	ORPHA:352	
MUMC+	Expertise Center Huntington's disease	Huntington disease	ORPHA:399	
MUMC+	Expertise Center Hyperostosis of the skull	Osteopetrosis	ORPHA:2781	
MUMC+	Expertise Center Neural tube defects	Neural tube defect	ORPHA:3388	
MUMC+	Expertise center Pulmonary hypertension	Pulmonary hypertension with unclear multifactorial mechanism; auto immune mechanisms in PH, and right ventricular failure	ORPHA:275844	
MUMC+	Expertise center Rare syndromes and cognitive disorders	Kabuki syndrome	ORPHA:2322	
MUMC+	Expertise center Rare syndromes and cognitive disorders	Rare developmental defect during embryogenesis	ORPHA:68341 ORPHA:68335	
MUMC+	Expertise center Rare syndromes and cognitive disorders	Rett Syndrome	ORPHA:778	
MUMC+	Gastro-intestinal center Maastricht	Rare hepatic and biliary tract tumor, incl. Cholangiocarcinoma Carcinoma of the gallbladder, Carcinoma of the ampulla of Vater, Hepatocellular carcinoma, Fibrolamellar hepatocellular carcinoma, Hepatocellular adenoma	ORPHA:56044 ORPHA:88673	voor biliary tract carcinoom en hepatho-cellular carcinoma
MUMC+	Limburg renal registry	EGPA, GPA and MPA	ORPHA:183 ORPHA:900 ORPHA:727	
MUMC+	Limburg renal registry	Cryoglobulinemic vasculitis	ORPHA:91138	
MUMC+	Lung cancer center Maastricht	Small cell lung cancer	ORPHA:70573	
MUMC+	Maastricht Gynaecological Oncology Center	Rare ovarian cancer; epithelial and non- epithelial, also tumor of Fallopian tubes	ORPHA:180220	
MUMC+	Maastricht Gynaecological Oncology Center	Rare cancer of the corpus uteri	ORPHA:213569	
MUMC+	Maastricht Head & Neck Cancer Center	Rare otorhinolaryngologic tumor; as a group	ORPHA:98061	
MUMC+	Maastricht Soft Tissue Tumor Center	Rare soft tissue tumor; long list, together all soft tissue sarcomas in adults	ORPHA:71209	
MUMC+	MUMC Mediastinal tumors	Thymic tumor	ORPHA:100100	

MUMC+	Neuroendocrine tumours Center	Rare tumor; Neuro-endocrine tumor en carcinoid syndrome, Multiple endocrine neoplasia, Bronchial neuroendocrine tumor, Gastroenteropancreatic neuroendocrine tumor, Merkelcell carcinoma, Thymic neuroendocrine tumor	ORPHA:97287 ORPHA:79140	voor bronchial NET en Merkelcell carcinoma
MUMC+	Neuromuscular Centre MUMC+	Myotonic dystrophy	ORPHA:206647	
MUMC+	Neuromuscular Centre MUMC+	Duchenne and Becker muscular dystrophy	ORPHA:262	
MUMC+	Neuromuscular Centre MUMC+	Sodium channelopathy-related small fiber neuropathy	ORPHA:306577	
MUMC+	Neuromuscular Centre MUMC+	Neuromuscular disease	ORPHA:68381	
MUMC+	Neuro-oncologie centrum Maastricht	Glial tumor	ORPHA:182067	
MUMC+	Neuro-oncologie centrum Maastricht	Tumor of cranial and spinal nerves	ORPHA:252057	
MUMC+ & Radboudumc	Marfan and related disorders policlinic	Marfan syndrome	ORPHA:558	
MUMC+ & Radboudumc	Marfan and related disorders policlinic	Loeys-Dietz syndrome	ORPHA:60030	
MUMC+ & Radboudumc	Marfan and related disorders policlinic	Familial thoracic aortic aneurysm and aortic dissection	ORPHA:91387	
NKI-AvL	Center of rare GI tumours	Rare gastroesophageal tumor	ORPHA:180821	voor carcinoma of esophagus, incl. junction carcinoma ORPHA 70482
NKI-AvL	Center of rare GI tumours	Epithelial tumor of anal cancer	ORPHA:424010	
NKI-AvL	Center of rare head and neck tumours	Rare otorhinolaryngologic disease;Tumours of the salivary glands	ORPHA:276142	
NKI-AvL	Center of rare head and neck tumours	Rare otorhinolaryngologic disease; Tumours of the head and neck	ORPHA:290849	
NKI-AvL	Centre for rare nervous system tumor	Glial tumor	ORPHA:182067	
NKI-AvL	Expert Center for Hereditary Cancer	Hereditary breast and ovarian cancer syndrome	ORPHA:145	
NKI-AvL	Expert Center for Hereditary Cancer	Li-Fraumeni syndrome	ORPHA:524	
NKI-AvL	Expert Center of familial GI tumours	Familial gastric cancer	ORPHA:423776	
NKI-AvL	Expert Center of familial GI tumours	Genetic digestive tract tumor; 1. Hereditary nonpolyposis colon cancer 2. Familial adenomatous polyposis 3. Attenuated Familial adenomatous polyposis 4. Hereditary mixed poliposis syndrome	ORPHA:443909 ORPHA:733 ORPHA:220460 ORPHA:157794	
NKI-AvL	Expert centre for rare urological diseases	Testicular non seminomatous germ cell tumor; testicular germ cell tumors consist of testicular seminomatous germ cell tumor, testicular non-seminomatous germ cell tumor and spermatocytic seminoma	ORPHA:363472	voor testicular cancer
NKI-AvL	Expert centre for rare urological diseases	Squamous cell carcinoma of penis	ORPHA:398043	voor penile cancer
NKI-AvL	Expert Centre of rare thoracic tumours	Rare respiratory tumor; Rare pleural malignancies	ORPHA:50251	voor mesothelioom
NKI-AvL	Rare Skin Cancer Center	Merkel Cell carcinoma	ORPHA:79140	
NKI-AvL	Sarcoma Expertise Centre Amsterdam	Soft tissue sarcoma	ORPHA:3394	
NKI-AvL	Sarcoma Expertise Centre Amsterdam	Rare soft tissue tumor; Gastrointestinal stromal tumor (GIST)	ORPHA:44890	
NKI-AvL& UMCU	Expert Center of Neuroendocriene carcinomas	Gastroenteropancreatic endocrine tumor	ORPHA:100092	
PMC	Princess Máxima Center for rare paediatric tumors	rare tumor, pediatric	ORPHA:98057	
Radboudumc	Mycology reference center	Aspergillosis	ORPHA:1163	
Radboudumc	Mycology reference center	Aspergillosis; chronic aspergillosis and ABPA	ORPHA:1163 ORPHA:1164	
Radboudumc	Mycology reference center	Chronic mucocutaneous candidiasis	ORPHA:1334	
Radboudumc	Mycology reference center	Rare mycosis; in patients with hyper IgE syndrome	ORPHA:163591	
Radboudumc	Mycology reference center	Rare mycosis; in patients with chronic granulomatous dis.		
Radboudumc	(Pediatric) urology center	Agenesis and aplasia of uterine body; Cloacal anomalies and anal atresia in combination with uronary tract anomalies	ORPHA:180142	
Radboudumc	(Pediatric) urology center	Non-syndromic urogenital tract malformation of male and female; e.g. in spina bifida	ORPHA:182124	
Radboudumc	(Pediatric) urology center	Bladder exstrophy; incl. cloacal exstrophy and epispadias	ORPHA:322	
Radboudumc	(Pediatric) urology center	Posterior urethral valve	ORPHA:93110	
Radboudumc	(Pediatric) urology center	Posterior hypospadias	ORPHA:95706	
Radboudumc	Center for Head and Neck Oncology	Malignant epithelial tumor of the salivary glands	ORPHA:276145	
Radboudumc	Center for Head and Neck Oncology	Squamous cell carcinoma of head and neck	ORPHA:98061	
Radboudumc	Center for rare CNS and retinal vascular disease	Acquired aneurysmal subarachnoid hemorrhage	ORPHA:90065	
Radboudumc	Centre for genetic movement disorders	Rare hereditary ataxia, mainly autosomal dominant and recessive cerebellar ataxias	ORPHA:183518	
Radboudumc	Centre for genetic movement disorders	Hereditary spastic paraplegia	ORPHA:685	
Radboudumc	Centre for genetic movement disorders	Mainly Sjogren-Larsson syndrome, GLUT1 deficiency syndrome, and disorders of dopamine metabolism	ORPHA:816 ORPHA:71277 ORPHA:79169	
Radboudumc	Centre for genetic neurodevelopmental disorders	PTEN hamartoma tumor syndrome	ORPHA:201	voor Cowden syndrome
Radboudumc	Centre for genetic neurodevelopmental disorders	KBG syndrome	ORPHA:2332	
Radboudumc	Centre for genetic neurodevelopmental disorders	Kleefstra syndrome due to a point mutation	ORPHA:261652	
Radboudumc	Centre for genetic neurodevelopmental disorders	Multiple congenital anomalies/dysmorphic syndrome - variable intellectual disability	ORPHA:648	voor Noonan syndrome
Radboudumc	Centre for genetic neurodevelopmental disorders	Syndromic obesity	ORPHA:739	voor PWS i.p.v. Syndr. obesity
Radboudumc	Centre for genetic neurodevelopmental disorders	Multiple congenital anomalies/dysmorphic syndrome-intellectual disability	ORPHA:96169	voor Koolen -de Vries syndrome
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Vascular tumor; incl complicated hemangiomas	ORPHA:211237	
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Venous malformation	ORPHA:211252	
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Lymphatic system malformation	ORPHA:211255	
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Arteriovenous malformation	ORPHA:211266	
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Complex - combined vascular malformation; incl Klippel-Trenaunay-syndrome	ORPHA:211277	
Radboudumc	Centre for thyroid carcinomas	Thyroid tumor	ORPHA:100087	
Radboudumc	Centre of Paroxysmal Nocturnal hemoglobinuria	Paroxysmal nocturnal hemoglobinuria	ORPHA:447	
Radboudumc	Cleft (lip and) palate center Nijmegen	Cleft palate; cleft hard and / or soft palate	ORPHA:101023 ORPHA:99772	
Radboudumc	Cleft (lip and) palate center Nijmegen	Oculo-auriculo-vertebral spectrum (=hemifaciale microsomia)	ORPHA:141132	
Radboudumc	Cleft (lip and) palate center Nijmegen	Cleft palate; Submucosal cleft palate	ORPHA:155878	
Radboudumc	Cleft (lip and) palate center Nijmegen	Cleft palate; cleft lip/palate	ORPHA:199306	
Radboudumc	Craniofacial team Nijmegen	Craniosynostosis	ORPHA:1531	
Radboudumc	Craniofacial team Nijmegen	Cranial malformation	ORPHA:98038	
Radboudumc	Haemophilia treatment centre	Rare coagulation disorder; hemophilia	ORPHA:448	
Radboudumc	Haemophilia treatment centre	Rare coagulation disorder; von Willebrand Disease	ORPHA:903	
Radboudumc	Haemophilia treatment centre	Rare coagulation disorder; other	ORPHA:98429	
Radboudumc	Hearing & Genes Centre	Rare genetic deafness	ORPHA:68361	
Radboudumc	Hearing & Genes Centre	Usher syndrome	ORPHA:886	
Radboudumc	Hereditary cancer centre	Hereditary nonpolyposis colon cancer; Lynch Syndrome	ORPHA:144	
Radboudumc	Hereditary cancer centre	Hereditary breast and ovarian cancer syndrome; BRCA mutation carriers	ORPHA:145	
Radboudumc	Hereditary cancer centre	APC-related attenuated familial adenomatous polyposis	ORPHA:247806	
Radboudumc	Hereditary cancer centre	Familial gastric cancer, incl her. diffuse GC	ORPHA:423776	
Radboudumc	Liver cyst center	Isolated polycystic liver disease	ORPHA:2924	
Radboudumc	Neuromuscular Centre	Myotonic dystrophy	ORPHA:206647	
Radboudumc	Neuromuscular Centre	Non-dystrophic myopathy	ORPHA:206656	
Radboudumc	Neuromuscular Centre	Congenital myotonia and paramyotonia congentia	ORPHA:206973 ORPHA:684	
Radboudumc	Neuromuscular Centre	Duchenne and Becker muscular dystrophy	ORPHA:262	
Radboudumc	Neuromuscular Centre	Facioscapulohumeral dystrophy	ORPHA:269	
Radboudumc	Neuromuscular Centre	Oculopharyngeal muscular dystrophy	ORPHA:270	
Radboudumc	Neuromuscular Centre	Neuralgic amyotrophy	ORPHA:2901	
Radboudumc	Neuromuscular Centre	Neuromuscular disease	ORPHA:68381	
Radboudumc	Neuromuscular Centre	Idiopathic inflammatory myopathy	ORPHA:98482	
Radboudumc	Nijmegen Center for Disorders of Glycosylation (NCDG)	Congenital disorder of glycosylation	ORPHA:137	
Radboudumc	Nijmegen centre for mitochondrial disorders	Mitochondrial disease	ORPHA:68380	
Radboudumc	Ophthalmogenetic center	Choroideremia	ORPHA:180	

Radboudumc	Ophthalmogenetic center	Retinal dystrophy; central serous retinopathy	ORPHA:443079	
Radboudumc	Ophthalmogenetic center	Leber congenital amaurosis	ORPHA:65	
Radboudumc	Ophthalmogenetic center	Stargardt disease and other ABCA4-related diseases	ORPHA:827 ORPHA:1872 ORPHA:791	
Radboudumc	Ophthalmogenetic center	Genetic vitreous-retinal disease	ORPHA:98657	
Radboudumc	Radboud Adrenal Centre	Adrenal/paraganglial tumor, incl Von Hippel Lindau and MEN-2 syndrome	ORPHA:100091	
Radboudumc	Radboud Adrenal Centre	Adrenal/paraganglial tumor; except catecholamines, aldosterone or cortisol producing . Incl. incidentalomas and carcinomas	ORPHA:100091	
Radboudumc	Radboud Adrenal Centre	Primary adrenal insufficiency	ORPHA:101958	
Radboudumc	Radboud Adrenal Centre	Adrenogenital syndrome	ORPHA:181412	
Radboudumc	Radboud Adrenal Centre	Rare primary hyperaldosteronism	ORPHA:181415	
Radboudumc	Radboud Adrenal Centre	Cushing syndrome	ORPHA:553	
Radboudumc	Radboud Center for Congenital Diaphragmatic Hernia and neonatal pulmonary hypertension	Congenital alveolo-capillary dysplasia	ORPHA:210122	
Radboudumc	Radboud Center for Congenital Diaphragmatic Hernia and neonatal pulmonary hypertension	Congenital diaphragmatic hernia; non- syndromic and syndromic	ORPHA:2140	
Radboudumc	Radboud Center for Congenital Diaphragmatic Hernia and neonatal pulmonary hypertension	Pulmonary hypertension owing to lung disease and/or hypoxia; Persistent Pulmonary Hypertension of the Newborn	ORPHA:275837	
Radboudumc	Radboud Center for iron disorders	Sideroblastic anemia	ORPHA:1047	
Radboudumc	Radboud Center for iron disorders	Constitutional dyserythropoietic anemia, mainly type I-IV	ORPHA:293830	
Radboudumc	Radboud Center for iron disorders	Disorder of iron metabolism and transport, focus on hereditary hemochromatosis, FTH1-related iron overload, congenital atransferrinemia, microcytic anemia with iron overload and aceruloplasminemia	ORPHA:309842 ORPHA:247790	
Radboudumc	Radboud Center for iron disorders	Constitutional anemia due to iron metabolism disorder	ORPHA:98360	
Radboudumc	Radboud Center Renal Disorders	Cystinosis	ORPHA:213	
Radboudumc	Radboud Center Renal Disorders	Renal or urinary tract malformation	ORPHA:93545	
Radboudumc	Radboud Center Renal Disorders	Glomerular disease	ORPHA:93548	
Radboudumc	Radboud Center Renal Disorders	Rare renal disease; Thrombotic microangiopathy (level ontbrak in form voor ref)	ORPHA:93573	
Radboudumc	Radboud Center Renal Disorders	Familial cystic renal disease; all cystic kidney dis. in children, incl. ciliopathies/nephronophtises	ORPHA:93587	
Radboudumc	Radboud Center Renal Disorders	Rare renal tubular disease	ORPHA:93603	
Radboudumc	Radboud Centre for vulvar and cervix cancer and Dutch Mole Registry	Vulvar intraepithelial neoplasia; rare vulvar cancers and VIN	ORPHA:137583	
Radboudumc	Radboud Centre for vulvar and cervix cancer and Dutch Mole Registry	Gestational trophoblastic neoplasm; all different subgroups	ORPHA:59305	
Radboudumc	Radboud DSD centre	Turner syndrome	ORPHA:881	
Radboudumc	Radboud DSD centre	46 -XX disorder of sex development induced by fetal androgens excess	ORPHA:90776	
Radboudumc	Radboud DSD centre	46 -XY disorder of sex development	ORPHA:98085	
Radboudumc	Radboud Intestinal failure Unit	Chronic intestinal failure	ORPHA:294422	
Radboudumc	Radboud Pituitary Center	Pituitary deficiency; also incl. all hypothalamic and pituitary diseases resulting in pituitary def.	ORPHA:101957	
Radboudumc	Radboud Pituitary Center	Prolactinoma; also incl. other rare types of functioning pituitary tumor	ORPHA:2965 ORPHA:314753	
Radboudumc	Radboud Pituitary Center	Cushing disease	ORPHA:96253	
Radboudumc	Radboud Pituitary Center	Somatotropic adenoma	ORPHA:96256	
Radboudumc	Radboud Sarcoma Center	Bone sarcoma; incl. bone and soft tissue tumors and GIST (gastointestinal stromal tumors)	ORPHA:223727 ORPHA:68411 ORPHA:71209 ORPHA:44890	
Radboudumc	Radboud Skull base centre	Rare tumor; different very rare skull base tumors	ORPHA:252164 ORPHA:252175 ORPHA:94080 ORPHA:29072 ORPHA:276627 ORPHA:324299	
Radboudumc	Radboud Skull base centre	Rare nervous system tumor; acoustic neuroma; cerebello pontine angle tumor; including NF2 patients	ORPHA:252175	excl. NF2
Radboudumc	Radboud Skull base centre	Tumor of endocrine glands; Hereditary pheochromocytoma-paraganglioma	ORPHA:29072	
Radboudumc	Radboud University Medical Centre CF centre	Cystic fibrosis	ORPHA:586	
Radboudumc	Radboudumc center for congenital disorders of dental development	Schöpf-Schulz-Passarge syndrome	ORPHA:50944	
Radboudumc	Radboudumc center for congenital disorders of dental development	Oligodontia	ORPHA:99798	
Radboudumc	Radboudumc center for facial palsy	Paralytic facial malformation	ORPHA:156224	
Radboudumc	Radboudumc Center for male infertility	Male infertility due to gonadal dysgenesis or sperm disorder	ORPHA:399764	
Radboudumc	Radboudumc Center for male infertility	Male infertility due to obstructive azoospermia	ORPHA:98343	
Radboudumc	Radboudumc Center for male infertility	Rare idiopathic male infertility	ORPHA:98345	
Radboudumc	Radboudumc Center for Pulmonary Hypertension	Idiopathic pulmonary arterial hypertension; all kinds of PH	ORPHA:71198 ORPHA:275766	
Radboudumc	Radboudumc center for systemic autoimmune diseases	Eosinophilic fasciitis	ORPHA:3165	
Radboudumc	Radboudumc center for systemic autoimmune diseases	Localized scleroderma	ORPHA:90289	
Radboudumc	Radboudumc center for systemic autoimmune diseases	Systemic sclerosis	ORPHA:90291	
Radboudumc	Radboudumc Expertise Center for immunodeficiency and autoinflammation	Immunodeficiency due to a complement cascade protein anomaly	ORPHA:101992	
Radboudumc	Radboudumc Expertise Center for immunodeficiency and autoinflammation	Primary immunodeficiency	ORPHA:101997	
Radboudumc	Radboudumc Expertise Center for immunodeficiency and autoinflammation	Autoinflammatory syndrome with immune deficiency	ORPHA:290839	
Radboudumc	Radboudumc Expertise Center for immunodeficiency and autoinflammation	Mevalonate kinase deficiency = Hyper IgD Syndrome (HIDS)	ORPHA:309025	
Radboudumc	Radboudumc Expertise Center for immunodeficiency and autoinflammation	Schnitzler syndrome	ORPHA:37748	
Radboudumc	Radboudumc Neuro-oncological center	Tumor of the neuroepithelial tissue	ORPHA:251558	
Radboudumc	Radboudumc Neuro-oncological center	Tumor of the meninges	ORPHA:252025	voor primary melanocytic tumor of the CNS
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Anal fistula	ORPHA:228113	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Gastroschisis	ORPHA:2368	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Caudal regression sequence	ORPHA:3027	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Hirschsprung disease	ORPHA:388	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	High anorectal malformation	ORPHA:557	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Omphalocele	ORPHA:660	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	VACTERL/VATER association	ORPHA:887	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Cloacal exstrophy	ORPHA:93929	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Intestinal malformation	ORPHA:97945	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Intermediate anorectal malformation		
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	Low anorectal malformation		
Radboudumc & MUMC+	Marfan and related disorders policlinic	Marfan syndrome	ORPHA:558	
Radboudumc & MUMC+	Marfan and related disorders policlinic	Loeys-Dietz syndrome	ORPHA:60030	

Radboudumc & MUMC+	Marfan and related disorders policlinic	Familial thoracic aortic aneurysm and aortic dissection	ORPHA:91387	
Radboudumc/UCCZ Dekkerswald	Center for Mycobacterial diseases	Tuberculosis; also incl. nontuberculous mycobacterial inf.	ORPHA:3389 ORPHA:411703	
STZ - St. Antonius Ziekenhuis, Nieuwegein	St. Antonius Oesofagus Centrum	Esophageal carcinoma	ORPHA:70482	
STZ- St Elisabeth Hospital Tilburg	Neuro-oncology Center Tilburg	Glial tumor	ORPHA:182067	
STZ-Albert Schweitzer hospital, Dordrecht	Centre of expertise Retroperitoneal Fibrosis	Retroperitoneal fibrosis	ORPHA:49041	
STZ-CWZ Nijmegen	Center for Cerebrotendinous xanthomatosis	Cerebrotendinous xanthomatosis	ORPHA:909	
STZ-CWZ Nijmegen	Malignant Hyperthermia investigation unit Nijmegen	Malignant hyperthermia	ORPHA:423	
STZ-Jeroen Bosch Hospital	Center for Primary immunodeficiencies	Transient hypogammaglobulinemia of infancy	ORPHA:101977	Voorlopig t/m sep 2017, voor Immuno-deficiency predominantly affecting antibody production, mainly the various types of unclassified antibody deficiency
STZ-Maxima Medisch Centrum	Center for Adrenal Tumors	Adrenocortical carcinoma; also 'Catecholamine-producing tumor'and the 'Adrenal incidentaloma'	ORPHA:1501	alleen voor Adrenocortical carcinoma
STZ-Maxima Medisch Centrum	SolviMáx, Center of Excellence for Abdominal Wall and Groin Pain	Acquired peripheral neuropathy; Anterior cutaneous nerve syndrome (ACNES)	ORPHA:51890	alleen voor ACNES
STZ-Medisch Centrum Haaglanden-Bronovo-Nebo	Center for Neuro-oncology The Hague	Rare nervous system tumor	ORPHA:182067	voor gliomen
STZ-OLVG	Center for HME-MO (Hereditaire Multipele exostosen-Multipele Osteochondromen)	Multiple osteochondromas; Hereditary Multiple Exostoses - Multiple Osteochondromas	ORPHA:321	
STZ-OLVG	EC for interstitial lungdiseases OLVG	Idiopathic pulmonary fibrosis; incl. idiopathic interstitial pneumonia	ORPHA:98300	voor IPF en IIP
STZ-OLVG		Exposure-related interstitial lung disease; in its broadest sense	ORPHA:264984	
STZ-St. Antonius Ziekenhuis	Center for Pulmonary vascular diseases	Idiopathic pulmonary arterial hypertension	ORPHA:275766	
STZ-St. Antonius Ziekenhuis	Center for Pulmonary vascular diseases	Chronic thromboembolic pulmonary hypertension	ORPHA:70591	
STZ-St. Antonius Ziekenhuis	Center for Pulmonary vascular diseases	Hereditary hemorrhagic telangiectasia	ORPHA:774	
STZ-St. Antonius Ziekenhuis	Interstitial Lung Diseases Center of Excellence	Interstitial lung disease	ORPHA:182095	
STZ-St. Antonius Ziekenhuis	Interstitial Lung Diseases Center of Excellence	Idiopathic pulmonary fibrosis (IPF)	ORPHA:2032	
STZ-St. Antonius Ziekenhuis	Interstitial Lung Diseases Center of Excellence	Hypersensitivity pneumonitis (PH)	ORPHA:31740	
STZ-St. Antonius Ziekenhuis	Interstitial Lung Diseases Center of Excellence	Sarcoidosis	ORPHA:797	
SZT-Maasstad Hospital	Burn Centre Maasstad Hospital, in cooperation with Burn Centre Red Cross Hospital and Martini Hospital (ADBC: Association of Dutch Burn Care Centers)	Toxic epidermal necrolysis	ORPHA:537	
SZT-Medisch Spectrum Twente	Gastrointestinal ischemia Centre	Celiac trunk compression syndrome	ORPHA:293208	
SZT-St. Elisabeth Ziekenhuis	Neurovascular Center Tilburg	Neurovascular malformation	ORPHA:102006	
SZT-St. Elisabeth Ziekenhuis	Neurovascular Center Tilburg	Acquired aneurysmal subarachnoid hemorrhage	ORPHA:90065	
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Rare acquired eye disease; Herpes simplex virus keratitis, stromal, neurtrophic and endotheliitis	ORPHA:137586 ORPHA:137596	voor alle genoemde aandoeningen
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Uveal melanoma	ORPHA:39044	
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Central serous chorioretinopathy	ORPHA:443079	
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Rare genetic eye disease; Retinal Dystrophies	ORPHA:71862	
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Uveitis	ORPHA:98715	
UMCG	Adrenal centre UMCG	Rare primary hyperaldosteronism	ORPHA:181415	
UMCG	Adrenal centre UMCG	Catecholamine-producing tumor, incl. pheochromocytoma/ paraganglioma and non-secreting head and neck paragangliomas	ORPHA:717 ORPHA:94080	
UMCG	Brain Centre Rudolf Magnus, Neuromuscular Diseases		ORPHA:206701 ORPHA:53739 ORPHA:70 ORPHA:211037	voor Spinal muscular atrophy
UMCG	Center for Blistering Diseases	Inherited epidermolysis bullosa	ORPHA:79361	
UMCG	Center for Blistering Diseases	Autoimmune bullous skin disease; all forms of pemphigus and pemfigoid	ORPHA:79669	
UMCG	Center for rare inherited inborn errors of metabolism	Molybdenum cofactor deficientie type A	ORPHA:308386	
UMCG	Centre for Necrotizing Enterocolitis	Necrotiserende Enterocolitis	ORPHA:391673	
UMCG	Centre of expertise for choledochal malformations	Choledochal cyst	ORPHA:480501	
UMCG	Centre of familial tumors	Inherited cancer-predisposing syndrome; incl VHL, MEN1, MEN2 and familial paraganglioma/PCC.	ORPHA:140162 ORPHA:29072	
UMCG	Clinic for Connective tissue disorders	Marfan syndrome	ORPHA:558	
UMCG	Clinic for rare chromosome disorders	Autosomal anomaly; wide diversity of chromosomal deletions and duplications (and not the more common trisomies)	ORPHA:98127	
UMCG	Cystic Fibrosis centre Groningen	Cystic fibrosis	ORPHA:586	
UMCG	Dutch expertise centre for lympho-vascular medicine	Primary lymphedema	ORPHA:77240	
UMCG	Dutch expertise centre for lympho-vascular medicine	Syndromic lymphedema	ORPHA:89832	
UMCG	Expert Center for Children and Adults with Pulmonary Hypertension	Pulmonary arterial hypertension	ORPHA:182090	
UMCG	Expert Center for Children and Adults with rare Congenital Heart Diseases	Congenital heart malformation	ORPHA:88991	
UMCG	Expert Center head- and neck oncology	Squamous cell carcinoma of head and neck	ORPHA:98061	
UMCG	Expert Center neuro-oncology in adults	Glial tumor	ORPHA:182067	
UMCG	Expert centre for carcinoid / neuroendocrine carcinoma's (NEC)	Carcinoid tumor and carcinoid syndrome	ORPHA:100093	
UMCG	Expert Centre for Cardiogenetics	Genetic cardiac rhythm disease	ORPHA:101934	
UMCG	Expert Centre for Cardiogenetics	Cardiomyopathy	ORPHA:167848	
UMCG	Expert Centre for Cardiogenetics	Familial isolated arrhythmogenic ventricular dysplasia - biventricular form	ORPHA:293899	
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease due to glucose-6-phosphatase deficiency type a	ORPHA:364	
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease due to glycogen debranching enzyme deficiency	ORPHA:366	
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease due to liver glycogen phosphorylase deficiency	ORPHA:369	
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease	ORPHA:79201	
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease due to glucose-6-phosphatase deficiency type b		
UMCG	Expert centre for M(C)ADD	Multiple acyl-CoA dehydrogenation deficiency - severe neonatal type	ORPHA:26791	
UMCG	Expert centre for M(C)ADD	Medium chain acyl-CoA dehydrogenase deficiency	ORPHA:42	
UMCG	Expert centre for M(C)ADD	Multiple acyl-CoA dehydrogenation deficiency - mild type		
UMCG	Expert centre for movement disorders in adults and children	Paroxysmal dyskinesia	ORPHA:1431	
UMCG	Expert centre for movement disorders in adults and children	Inherited congenital spastic tetraplegia	ORPHA:210141	
UMCG	Expert centre for movement disorders in adults and children	Rare myoclonus; myoclonus dystonia GOSR2 and FCMTE	ORPHA:280620 ORPHA:86814	

UMCG	Expert centre for movement disorders in adults and children	Hyperekplexia	ORPHA:306773	
UMCG	Expert centre for movement disorders in adults and children	Neurodegeneration with brain iron accumulation; a.o. PKAN	ORPHA:385	
UMCG	Expert centre for movement disorders in adults and children	Rare choreic movement disorder, huntington's disease	ORPHA:399	
UMCG	Expert centre for movement disorders in adults and children	Rare dystonia: myoclonus, focal,generalised, dopa responsive	ORPHA:68363	
UMCG	Expert centre for movement disorders in adults and children	Psychogenic movement disorders	ORPHA:71519	
UMCG	Expert centre for movement disorders in adults and children	Autosomal dominant cerebellar ataxia, incl recessive ataxias	ORPHA:99 ORPHA:1172	
UMCG	Expert centre for movement disorders in adults and children	Neurometabolic disease, related tot movement disorders		
UMCG	Expert centre for Phenylketonuria (PKU) and Tyrosinemia type I	Disorder of phenylalanin or tyrosine metabolism	ORPHA:79190	
UMCG	Expert centre for serine deficiencies	Neurometabolic disorder due to serine deficiency	ORPHA:35705	
UMCG	Expert centre for systemic vasculitis	Granulomatosis with polyangiitis	ORPHA:52759	voor vasculitis
UMCG	Expert centre for systemic vasculitis	Eosinophilic granulomatosis with polyangiitis		
UMCG	Expert centre for systemic vasculitis	Giant cell arteritis		
UMCG	Expert centre for systemic vasculitis	Microscopic polyangiitis		
UMCG	Expert centre for systemic vasculitis	Takayasu arteritis		
UMCG	Expert centre mastocytosis Netherlands (ECMN)	Mastocytosis	ORPHA:98292	
UMCG	Expert centre Sjögren syndrome	Tumor of hematopoietic and lymphoid tissues; MALT lymphoma associated with Sjögren's disease	nn	
UMCG	Expert centre Sjögren syndrome	Systemic autoimmune disease; Sjögren (including MALT lymphoma)	ORPHA:289390	
UMCG	Expertise Center for Polycystic Kidney Diseases	Familial cystic renal disease	ORPHA:93587	
UMCG	Expertise Center Germcell tumors	Testicular germ cell tumor	ORPHA:363504	
UMCG	Expertise Center Groningen Papilloma studies	Recurrent respiratory papillomatosis	ORPHA:60032	
UMCG	Expertise Center Gyneco-oncology UMCG	Rare cancer of the cervix uteri; incl. squamous cell carcinoma	ORPHA:213761	
UMCG	Expertise Center Gyneco-oncology UMCG	Malignant epithelial tumor of ovary; different types adenocarcinoma	ORPHA:398934	
UMCG	Expertise Center Gyneco-oncology UMCG	Rare vulvovaginal tumor; squamous cell carcinoma of vulva	ORPHA:494448	
UMCG	Expertise Center pediatric liver disease, pediatric liver surgery and pediatric liver transplantation	Biliary atresia	ORPHA:30391	
UMCG	Expertise Center Soft tissue and bone tumors	Soft tissue sarcomas	ORPHA:3394	
UMCG	Expertise Center Soft tissue and bone tumors	Osteosarcoma	ORPHA:668	
UMCG	Familial Breast Ovarian Cancer Clinic	Hereditary breast and ovarian cancer syndrome; BRCA1 BRCA2	ORPHA:145	
UMCG	Familial Colorectal Cancer Clinic	Hereditary nonpolyposis colon cancer	ORPHA:443909	
UMCG	Familial Colorectal Cancer Clinic	Familial adenomatous polyposis; incl FAP and MUTYH	ORPHA:733 ORPHA:220460	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Senile systemic amyloidosis	ORPHA:330001	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Primary systemic amyloidosis	ORPHA:85443	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Secondary amyloidosis	ORPHA:85445	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Transthyretin-related familial amyloid cardiomyopathy	ORPHA:85451	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Familial amyloid polyneuropathy	ORPHA:93560 ORPHA:238269 ORPHA:439232 ORPHA:85448 ORPHA:271861	
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Primary localized amyloidosis		
UMCG	Hemophilia Treatment Centre UMCG	Rare hemorrhagic disorder; hemophilia	ORPHA:448	
UMCG	Hemophilia Treatment Centre UMCG	Rare hemorrhagic disorder: other allied bleeding disorders	ORPHA:68334	
UMCG	Hemophilia Treatment Centre UMCG	Rare hemorrhagic disorder; von Willebrand Disease	ORPHA:903	
UMCG	Neurovascular Team UMCG	Cerebral malformation, intracranial dural AV-fistula	ORPHA:97339	
UMCG	Neurovascular Team UMCG	Cerebral malformation, brainstem cavernomas		
UMCG	Neurovascular Team UMCG	Cerebral malformation, proliferative angiopathy		
UMCG	Neurovascular Team UMCG	Spinal arteriovenous shunts		
UMCG	Paediatric centre for Rheumatologic and immunologic diseases	Juvenile idiopathic arthritis	ORPHA:92	
UMCG	Paediatric Colorectal Expertise Center Groningen	Hirschsprung disease	ORPHA:388	
UMCG	Pediatric Neuro-oncology Team	Medulloblastoma; and PNET	ORPHA:616 ORPHA:251870	
UMCG	Pediatric Neuro-oncology Team	Rare nervous system tumor; pediatric CNS tumours	ORPHA:98062	voor rare nervous system tumor
UMCG	Pediatric Oncology group UMCG	Tumor of hematopoietic and lymphoid tissues	ORPHA:171895 ORPHA:223735	
UMCG	Small bowel rehabilitation and transplant centre	Chronic intestinal failure	ORPHA:294422	
UMCG	The multidisciplinary CHARGE clinic	CHARGE syndrome	ORPHA:138	
UMCG	Thyroid cancer centre	Thyroid tumor	ORPHA:100087	
UMCG	Tuberculosis centre Beatrixoord	Tuberculosis	ORPHA:3389	
UMCG	UMCG Pituitary Center	Rare hypothalamic or pituitary disease	ORPHA:99408 ORPHA:101957	voor rare pituitary disease
UMCG	UMCG/ oesophageal/ gastric cancer tumorgroup	Esophageal adenocarcinoma	ORPHA:99976	
UMCG	Paediatric Colorectal Expertise Center Groningen	Anorectal malformation	ORPHA:96346	
UMCU	Expert Centre Hereditary and congenital nephrologic and urologic disorders	Non-syndromic renal or urinary tract malformation; CAKUT	ORPHA:93546	
UMCU	Expert Centre Hereditary and congenital nephrologic and urologic disorders	Rare renal disease; congenital or inherited renal or urinary tract disease	ORPHA:93547	
UMCU	Expert Centre Hereditary and congenital nephrologic and urologic disorders	Familial cystic renal disease; nephronophthisis, as feature of i.e. Joubert and Meckel or isolated or part of other ciliopathies.	ORPHA:93587 ORPHA:65 ORPHA:156162	
UMCU	Brain Centre Rudolf Magnus, Neuromuscular Diseases	Neuromuscular disease; ALS, PLS, Progressive Spinal Muscular Atrophy and polyneuropathy	ORPHA:68381	voor Neuro-muscular disease
UMCU	Center for Rare Ear and Hearing Diseases	Middle ear anomaly	ORPHA:164004	
UMCU	Center for Refractory Pediatric Epilepsy	Rare epilepsy	ORPHA:101998	
UMCU	Center for Refractory Pediatric Epilepsy	Genetic causes of refractory pediatric epilepsy	ORPHA:166463	voor Epilepsy syndrome
UMCU	Center for Refractory Pediatric Epilepsy	Continuous spikes and waves during sleep and ESES	ORPHA:725	
UMCU	Center for Refractory Pediatric Epilepsy	Tuberous sclerosis	ORPHA:805	
UMCU	Center Inherited Metabolic Diseases	Disorder of fatty acid oxidation and ketone body metabolism	ORPHA:79174	
UMCU	Center Inherited Metabolic Diseases	Disorder of pyridoxine metabolism	ORPHA:79192	
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	Hemifaciale microsomie & Microtie	ORPHA:141136 ORPHA:83463	
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	Cleft/lip palate	ORPHA:199306	
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	22q11.2 deletion syndrome	ORPHA:567	
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	Isolated Pierre Robin syndrome	ORPHA:718	
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	Rare odontologic disease; oligodontia	ORPHA:99798	
UMCU	Centre for rare tumors	Thyroid tumor	ORPHA:100087	
UMCU	Centre for rare tumors	Inherited cancer-predisposing syndrome	ORPHA:652	voor Men 1
UMCU	Centre for rare tumors	Multiple endocrine neoplasia type 2A; incl. fam medullary thyroid carcinoma, MEN2B and sporadic medulary thyroid carcinoma	ORPHA:653 ORPHA:1332	
UMCU	Centre for rare tumors	Von Hippel-Lindau disease	ORPHA:892	
UMCU	Children's Heartcenter WKZ	Congenital heart malformation	ORPHA:88991	
UMCU	Clinic for Tuberous Sclerosis Complex	Rare genetic neurological disorder; Tubererous Sclerosis Complex	ORPHA:805	
UMCU	Cystic Fibrosis Clinic	Cystic fibrosis	ORPHA:586	
UMCU	Expertise Center Rare GI and hepatic diseases	Progressive familial intrahepatic cholestasis	ORPHA:172	
UMCU	Expertise Center Rare GI and hepatic diseases	Intractable diarrhea of infancy; due to genetic defects	ORPHA:363300	
UMCU	Expertise Center Rare GI and hepatic diseases	Wilson disease	ORPHA:905	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare anemia	ORPHA:108997	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder; Congenital and acquired platelet disorders	ORPHA:248326	

UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder due to a coagulation factors defect; deficiency of factor II/ V/ VII/X/XI	ORPHA:325 ORPHA:326 ORPHA:327 ORPHA:328 ORPHA:329	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder due to a coagulation factors defect; hemophilia	ORPHA:448	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare constitutional medullar aplasia; Fanconi, Diamond-Blackfan anemia and congenital neutropenia	ORPHA:68383 ORPHA:101987	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder due to a coagulation factors defect; antiplasmin deficiency	ORPHA:79	
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder due to a coagulation factors defect; von Willebrand Disease	ORPHA:903	
UMCU	Expertise centre for malignant hematology	Myeloproliferative neoplasm ; Polycytemia vera (PV) and Essential thrombocythemia (ET)	ORPHA:29073	voor Multiple myeloma
UMCU	Expertise centre for malignant hematology	Primary central nervous system lymphoma	ORPHA:513 ORPHA:519	voor acute lymphoblastic and myeloid leukemia
UMCU	Expertise centre for malignant hematology	Myeloid hemopathy, ao CMMol / Myelofibrosis/ eosinophilic disorders	ORPHA:547	voor Non Hodgkin lymphoma
UMCU	Expertise centre for malignant hematology	Acute lymphoblastic leukemia		
UMCU	Expertise centre for malignant hematology	Lymphoma; NHL, Hodgkin, CLL, Waldenstrom, hairycell and other lymfoproliferative diseases		
UMCU	Expertise centre for malignant hematology	Myeloid hemopathy; AML/RAEB-T, RAEB, Myelodysplasia and remaining diseases		
UMCU	Expertise centre for malignant hematology	Myeloid hemopathy; HES and Langerhanscel histiocytose		
UMCU	Expertise centre for malignant hematology	Myeloproliferative neoplasm; CML		
UMCU	Expertise centre for malignant hematology	Plasma cell tumor; MM/ Amyloidosis and Monoclonal gammopathy(MGUS)		
UMCU	Expertise centre for malignant hematology	Systemic mastocytosis		
UMCU	Expertise centre for primary immunodeficiences	Primary immunodeficiency due to a defect in adaptive immunity; B and T cell immunodeficiencies: SCID, CID	ORPHA:101972	
UMCU	Expertise centre for primary immunodeficiences	Immunodeficiency predominantly affecting antibody production; incl. CVID, XLA, other types of complete agammaglobulinemia	ORPHA:101977	
UMCU	Expertise centre for primary immunodeficiences	Primary immunodeficiency due to a defect in innate immunity	ORPHA:101988	
UMCU	Expertise centre for primary immunodeficiences	Primary hemophagocytic lymphohistiocytosis	ORPHA:158038	
UMCU	Expertise centre for primary immunodeficiences	Graft versus host disease	ORPHA:39812	
UMCU	Expertise centre for primary immunodeficiences	Immunodeficiency predominantly affecting antibody production; incl. SADNI, IgG subclass- and IgA def.		
UMCU	Expertise centre systemic autoimmune diseases	Periodic fever syndrome, incl. CAPS, FMF, behcet, Traps, Pfapa	ORPHA:101995 ORPHA:117	
UMCU	Expertise centre systemic autoimmune diseases	Systemic autoimmune disease; extraglandular manifestations in Sjogren	ORPHA:289390	
UMCU	Expertise centre systemic autoimmune diseases	Vasculitis	ORPHA:52759	
UMCU	Expertise centre systemic autoimmune diseases	Systemic sclerosis	ORPHA:90291	
UMCU	Expertise centre systemic autoimmune diseases	Juvenile idiopathic arthritis	ORPHA:92	
UMCU	Expertise centre systemic autoimmune diseases	Juvenile dermatomyositis	ORPHA:93672	
UMCU	Expertise centre systemic autoimmune diseases	Rare coagulation disorder; Antiphospholipid syndrome		
UMCU	Multidisciplinary Center for Limb Reduction Defects	Rare bone disease; m.n. primaire skeletdysplasieen en dysostoses, ook osteogenesis imperfecta en achondroplasie	ORPHA:364526 ORPHA:364559	
UMCU	Multidisciplinary Center for Limb Reduction Defects	Non-syndromic limb reduction defects	ORPHA:93457	
UMCU	UMCU ophthalmology uveitisgroup	Intermediate uveitis; (non) infectious	ORPHA:279914	
UMCU	UMCU ophthalmology uveitisgroup	Anterior uveitis; (non) infectious	ORPHA:279922 ORPHA:306648	
UMCU	UMCU ophthalmology uveitisgroup	Posterior uveitis; (non) infectious and in syst dis.	ORPHA:280892	
UMCU	UMCU ophthalmology uveitisgroup	Systemic diseases with panuveitis	ORPHA:280933	
UMCU	UMCU ophthalmology uveitisgroup	Rare inflammatory eye disease		
UMCU	WKZ center for congenital malformations	Esophageal atresia	ORPHA:1199	
UMCU	WKZ center for congenital malformations	Hirschsprung Disease	ORPHA:388	
UMCU	WKZ center for congenital malformations	Anorectal malformation	ORPHA:96346	
UMCU	Center for inherited cardiovascular disease	Genetic cardiac rhytm disease	ORPHA:101934	
UMCU	Center for inherited cardiovascular disease	Cardiomyopathy	ORPHA:167848	
UMCU	Center for inherited cardiovascular disease	Rare genetic vascular disease	ORPHA:233655	
UMCU	Center for Neonatal Neurology	Hypoxic ischemic brain injury	ORPHA:137577	
UMCU	Center for Neonatal Neurology	Periventricular leukomalacia	ORPHA:171676	
UMCU	Center for Neonatal Neurology	Pediatric arterial ischemic stroke	ORPHA:439175	
UMCU	Center of vascular anomalies Utrecht	PHACE syndrome	ORPHA:211243 ORPHA:211237	voor vascular anomaly
UMCU	Center of vascular anomalies Utrecht	Diffuse neonatal hemangiomatosis		
UMCU	Center of vascular anomalies Utrecht	Kaposiform hemangioendothelioma		
UMCU	Center of vascular anomalies Utrecht	Laryngotracheal angioma		
UMCU	Center of vascular anomalies Utrecht	Macrocystic lymphatic malformation		
UMCU	Center of vascular anomalies Utrecht	Microcystic lymphatic malformation		
UMCU	Center of vascular anomalies Utrecht	Mucocutaneous venous malformations		
UMCU	Center of vascular anomalies Utrecht	PELVIS syndrome		
UMCU	Center of vascular anomalies Utrecht	Rapidly involuting congenital hemangioma; RICH and NICH		
UMCU	Center of vascular anomalies Utrecht	Vascular malformation		
UMCU	Centre of expertise for extracranial carotid artery aneurysms (ECAA)	Extracraniele carotis aneurysma	ORPHA:494424	
UMCU	Dutch National Expertise Center for Pseudoxanthoma elasticum (DNECP)	Pseudoxanthoma elasticum-	ORPHA:758	
UMCU	Head and Neck Working Group Utrecht	Nasopharyngeal carcinoma	ORPHA:150	
UMCU	Head and Neck Working Group Utrecht	Rare tumors of salivary glands	ORPHA:276142	
UMCU	Head and Neck Working Group Utrecht	Squamous cell carcinoma of the oral tongue	ORPHA:457252	
UMCU	Head and Neck Working Group Utrecht	Squamous cell carcinoma of head and neck	ORPHA:98061	
UMCU	Mobility Clinic	Osteochondritits Dissecans	ORPHA:2764	
UMCU	Sylvia Toth Center for Multi-disciplinary follow up of Lysosomal Storage Disorders, University Medical Center Utrecht	Hurler disease	ORPHA:93473	voor Hurler disease
UMCU	UMCU Neuro-Oncology Center	Glial tumor	ORPHA:182067	
UMCU	Utrecht Center for Keratoconus and Corneal Dystrophies	Keratoconus	ORPHA:156071	
UMCU & NKI-AvL	Expert Center of Neuroendocriene carcinomas	Gastroenteropancreatic endocrine tumor		
VUmc	Birt-Hogg-Dubé task force	Inherited renal cell cancer-predisposing syndrome; Birt-Hogg-Dubé syndrome	ORPHA:122	
VUmc	Brain Tumor Center Amsterdam	Rare tumor; Brain Tumors	ORPHA:98062	
VUmc	Celiac disease center	Short Bowel Syndrome	ORPHA:104008	
VUmc	Celiac disease center	Autoimmune hepatitis	ORPHA:2137	
VUmc	Celiac disease center	Refractory celiac disease	ORPHA:398063	
VUmc	Celiac disease center	Enteropathy Associated T-cell Lymphoma	ORPHA:86880	
VUmc	Center for Childhood White Matter Disorders	Rare neurologic disease	ORPHA:68356 ORPHA:171676 ORPHA:83629	
VUmc	Center for pediatric oral and maxillofacial surgery	Rare odontologic disease, Robin sequence	ORPHA:718	voor Robin seq.
VUmc	Center for pediatric rehabilitation medicine	Spastic diplegia - infantile type		
VUmc	Center for rare haematologic cancers	Myeloid hemopathy	ORPHA:171895	
VUmc	Center for rare haematologic cancers	Lymphoid hemopathy	ORPHA:171898	
VUmc	Center for rare haematologic cancers	Acute myeloid leukemia	ORPHA:29073	voor Multiple myeloma
VUmc	Center for rare haematologic cancers	Hodgkin lymphoma	ORPHA:513 ORPHA:519	voor acute lymphoblastic and myeloid leukemia
VUmc	Center for rare haematologic cancers	Chronic Myeloid Leukemia (al erkend onder cluster NHL)	ORPHA:521	
VUmc	Center for rare haematologic cancers	Myelodysplastic syndromes (al erkend onder cluster NHL)	ORPHA:52688	
VUmc	Center for rare haematologic cancers	Chronic myeloid leukemia	ORPHA:547	voor Non Hodgkin lymphoma
VUmc	Center for rare haematologic cancers	Primary Myelofibrosis	ORPHA:824	
VUmc	Center for rare haematologic cancers	Hodgkin Lymphoma (al erkend onder cluster AL&ML)	ORPHA:98293	

VUmc	Center for rare haematologic cancers	Acute lymphoblastic leukemia		
VUmc	Center for rare haematologic cancers	Aggressive B-cell non-Hodgkin lymphoma		
VUmc	Center for rare haematologic cancers	Enteropathy-associated T-cell lymphoma		
VUmc	Center for rare haematologic cancers	Langerhans cell histiocytosis		
VUmc	Center for rare haematologic cancers	Multiple myeloma; incl Amyloidosis, Castlemans disease and POEMS		
VUmc	Center for rare haematologic cancers	Myelodysplastic syndromes		
VUmc	Center for rare haematologic cancers	Myelofibrosis with myeloid metaplasia		
VUmc	Center on Atypical Sex or Gender development	Gender dysphoria	ORPHA:459690	
VUmc	Center on Atypical Sex or Gender development	Disorder of sex development	ORPHA:90771	
VUmc	Centre for Genetic Metabolic Diseases Amsterdam (CGMA) - VUmc specific	Disorder of creatine biosynthesis	ORPHA:79172	
VUmc	Centre for Systemic Sclerosis and Systemic Lupus Erythematosus, embedded in Amsterdam Rheumatology and immunology Centre	Systemic sclerosis	ORPHA:90291	
VUmc	Cleft Lip and Palate Team	Rare developmental defect during embryogenesis; cleft lip and palate	ORPHA:2014 ORPHA:1991	
VUmc	Down Center the Netherlands, location West	Down syndrome	ORPHA:870	
VUmc	Dutch Center for Oral Medicine and Oral Pathology	Squamous cell carcinoma of head and neck; Ameloblastomas and Keratocystic odontogenic tumors, Gorlin-Goltz syndrome	ORPHA:314419 ORPHA:447777	
VUmc	Dutch Retinoblastoma Center	Retinoblastoma	ORPHA:790	
VUmc	Expertcenter for Fybrodysplasia Ossificans Progressiva	Fibrodysplasia ossificans progressiva	ORPHA:337	
VUmc	Expertise Center for Osteogenesis Imperfecta	Primary bone dysplasia with decreased bone density	ORPHA:93446	
VUmc	Expertise Center Head and Neck tumors	Nasopharyngeal carcinoma	ORPHA:150	
VUmc	Expertise Center Head and Neck tumors	Hereditary pheochromocytoma-paraganglioma	ORPHA:29072	
VUmc	Expertise Center Head and Neck tumors	Rare head and neck tumor	ORPHA:290849	
VUmc	Expertise Center Head and Neck tumors	Squamous cell carcinoma of the oral cavity and lip	ORPHA:502369	
VUmc	Expertise Center Neurophthalmology	Idiopathic intracranial hypertension; loss of vision due to IIH	ORPHA:238624	
VUmc	Expertise Center Neurophthalmology	Acute zonal occult outer retinopathy	ORPHA:284454	
VUmc	Expertise Center Neurophthalmology	Paraneoplastic neurologic syndrome; optic neuropathies relevant to DD of CRION, RION and NMO-SD	ORPHA:36388	
VUmc	Expertise Center Neurophthalmology	Neuromyelitis optica: autoimmune optic neuropathies incl. NMO, CRION, RION, ION, MSON	ORPHA:71211 ORPHA:499085 ORPHA:499096 ORPHA:71505 ORPHA:420556	
VUmc	Expertise Center Neurophthalmology	Adult-onset myasthenia gravis; ocular MG		
VUmc	Expertise Center Preeclampsia from origin to healhy aging	Preeclampsia	ORPHA:275555	
VUmc	Fetal Akinesia Deformation Sequence Centre	neuromusculaire aandoeningen, in het bijzonder Foetale Akinesie	ORPHA:994	voor FADS
VUmc	PCD-center Vumc	Primary ciliary dyskinesia	ORPHA:244	
VUmc	Pediatric nephrology centre	Idiopathic nephrotic syndrome	ORPHA:357502	
VUmc	Pediatric nephrology centre	Non-syndromic renal or urinary tract malformation; CAKUT	ORPHA:93546	
VUmc	VUmc multidisciplinary (genetic) breast cancer team	Rare genetic tumor; Hereditary breast cancer, CHEK2*1100delC related	ORPHA:227535	voor hereditary breast cancer
VUmc	VUMC PH Centre	Chronic thromboembolic pulmonary hypertension	ORPHA:70591	
VUmc	VUMC PH Centre	Rare pulmonary hypertension; many subforms of PH; associated: collagen vascular disease, Hereditary PH, IPAH, type III WHO associated with emfysema.	ORPHA:71198	
VUmc & AMC	Amsterdam Centre of congenital malformations	Esophageal atresia	ORPHA:1199	
VUmc & AMC	Amsterdam Centre of congenital malformations	Chronic intestinal pseudoobstruction	ORPHA:2978	
VUmc & AMC	Amsterdam Centre of congenital malformations	Hirschsprung Disease	ORPHA:388	
VUmc & AMC	Amsterdam Centre of congenital malformations	Sacrococcygeal teratoma	ORPHA:494421	
VUmc & AMC	Amsterdam Centre of congenital malformations	Sarcoidosis	ORPHA:797	
VUmc & AMC	Amsterdam Centre of congenital malformations	Anorectal malformation	ORPHA:96346	
Vumc & AMC	Cystic Fibrosis Centre Amsterdam	Cystic Fibrosis	ORPHA:586	
VUmc & LUMC & AMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Centrale sereuze chorioretinopathie	ORPHA:443079	
VUmc & LUMC & AMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	ORPHA:71862	