UMC	Naam van Expertisecentrum	Cluster van / Specifieke aandoening(en)
AMC	Amsterdam Lysosome Center ("Sphinx")	Gaucher disease
		Fabry disease
		Niemann-Pick disease type A
		Niemann-Pick disease type B
		Niemann-Pick disease type C
		Mucopolysaccharidosis type 1
		Mucopolysaccharidosis type 3
		Mucopolysaccharidosis type 4
AMC	Dutch Centre for Peroxisomal disorders	Peroxisome biogenesis disorder-Zellweger syndrome spectrum
		Disorder of peroxisomal alpha- - beta- and omega-oxidation
		Rhizomelic chondrodysplasia punctata
		Non-syndromic pontocerebellar hypoplasia
AMC	Expertise center Vascular medicine	Homozygous familial hypercholesterolemia
		Familial lipoprotein lipase deficiency
		Tangier disease
AMC	Centre for Genetic Metabolic Diseases Amsterdam	Disorder of galactose metabolism
		Disorder of phenylalanine metabolism
AMC	Centre for Neuromuscular Diseases	Neuromuscular disease
		Motor neuron disease; amyotrophic lateral sclerosis, primary sclerosis and progressive muscular atrophy
		Idiopathic inflammatory myopathy, incl dermatomyositis, polymyositis, necrotizing autoimmune myopathy and inclusion body myositis
		Poliomyelitis
		Hereditary motor and sensory neuropathy
		Chronic inflammatory demyelinating polyneuropathy, incl. Guillain_Barre syndrome, CIDP, MMN
AMC	Centre for rare thyroid diseases	Congenital hypothyroidism
AMC	Centre for gastroenteropancreatic neuroendocrine tumors	Gastroenteropancreatic endocrine tumor
AMC	Centre for rare hypothalamic and pituitary diseases	Rare hypothalamic or pituitary disease
AMC	Hemophilia Comprehensive Care Treatment Centre	Rare hemorrhagic disorder due to a coagulation factors defect; hemophilia
		Rare hemorrhagic disorder due to a coagulation factors defect; von Willebrand Disease
AMC	Centre for Sickle Cell Disease	Hemoglobinopathy; incl Sickle cell dieaseand alfa or beta thalassemia.
AMC	Centre for Bone Marrow Failure	Rare constitutional medullar aplasia
		Shwachman Diamond disease
		Congenital neutropenia
AMC	Centre for pediatric thromboembolic events	Rare thrombotic disease of hematologic origin
AMC	Gastro-Intestinal Oncology Centre Amsterdam	Rare cystic pancreas tumors
		Rare hepatic and biliary tract tumor; incl gallbladder tumors, Ampulla of Vater carcinoma
AMC	Centre for Upper GI tumors Amsterdam	Gastro-esophageal tumor
AMC	Intestinal Failure Unit	Short bowel syndrome, also secondary
		Chronic intestinal failure
AMC	Achalasia Center, part of Esophageal Center Amsterdam	Sporadic achalasia
AMC	Centre for cholestatic liver diseases	Primary biliary cirrhosis
		Primary sclerosing cholangitis
AMC	Centre for Immunodeficiencies	Rare immune disease; incl. primary immunodeficiencies
AMC	National centre for primary hyperoxaluria	Primary hyperoxaluria
AMC	Centre for Kawasaki Disease	Kawasaki disease
AMC	Centre for Pediatric Rheumatic Diseases	Juvenile idiopathic arthritis
AMC	Expertise Center Clinical Imunology and Rheumatology-Vasculitis	Vasculitis
AMC	CAHAL (Center for Congenital Heart Disease Amsterdam-Leiden, adult CHD)	Congenital heart malformation; adult congenital heart disease

AMC	Centre for Hereditary cardiac rhythm and function disorders	Familial isolated arrhythmogenic right ventricular dysplasia
		Familial dilated cardiomyopathy with conduction defect due to LMNA mutation
		Genetic cardiac rhythm disease
		Familial dilated cardiomyopathy
		Peripartum cardiomyopathy
AMC	Centre for Marfan syndrome	Marfan syndrome
AMC	National Cornelia de Lange Expert Centre	Cornelia de Lange syndrome
		Rubinstein-Taybi syndrome
		Pitt-Hopkins syndrome
		Marshall-Smith syndrome
AMC	Centre for Hereditary Angioedema	Hereditary angioedema
AMC	Expertise center for genetic tumors of the digestive tract	Familial adenomatous polyposis
		Hereditary nonpolyposis colon cancer
		Hyperplastic polyposis syndrome
AMC & VUmc	Cystic Fibrosis Centre Amsterdam	Cystic fibrosis
AMC	Solvent Team	Rare intoxication
AMC	Center for Idiopathic Nephrotic Syndrome	Idiopathic nephrotic syndrome
AMC	Centre for Crigler-Najjar syndrome	Crigler-Najjar syndrome type 1
		Crigler-Najjar syndrome type 2
AMC	Vascular malformations and hemangiomas centre	Diffuse lymphatic malformation
		Mucocutaneous venous malformations
		Dandy-Walker malformation - facial hemangioma; all subs
		Familial multiple nevi flammei
AMC	AMC Pulmonary Hypertension Center	Eisenmenger syndrome
AMC	Amsterdam Multidisciplinary Lyme borreliosis Center	Lyme disease
AMC	Center for Paediatric oncology	Osteosarcoma
		soft tissue sarcomas
Erasmus MC	Center for Lysosomal and Metabolic Diseases	Mucopolysaccharidosis
		Neuronal ceroid lipofuscinosis
		Glycoproteinosis
		Disorder of lysosomal amino acid transport
		Sphingolipidosis
		Primary bone dysplasia with defective bone mineralization
Erasmus MC	Dutch Porphyria Center	Porphyria
		Erythropoietic protoporphyria
		Acute hepatic porphyria
Erasmus MC	Center urea cycle disorders and organic acidurias	Disorder of urea cycle metabolism and ammonia detoxification
		Disorder of branched-chain amino acid metabolism
Erasmus MC	NeMo, expert centre for Neuromuscular and Mitochondrial Diseases	Mitochondrial disease
Erasmus MC	Center for Neuro-inflammatory disorders	Paraneoplastic neurologic syndrome
		Postinfectious encephalitis
		Limbic encephalitis
		CLIPPERS
		Inflammatory and autoimmune disease with epilepsy
		Morvan syndrome
		Isaac syndrome
Erasmus MC	Pompe Center	Glycogen storage disease due to acid maltase deficiency - infantile onset
		idem - juvenile onset
		idem- adult onset
Erasmus MC	Neuromuscular Center Erasmus MC	Neuromuscular disease
		Guillain-Bar syndrome
		Chronic inflammatory demyelinating polyneuropathy
Erasmus MC	MS center	Neuromyelitis optica
		Multiple sclerosis variant

Erasmus MC	ENCORE - Expertise Center for Neuro-developmental disorders	Neurofibromatosis type 1
		Tuberous sclerosis
		Angelman syndrome
		Fragile X syndrome
		Sturge-Weber syndrome
		Central nervous system malformation
		Cardiofaciocutaneous syndrome
		Costello syndrome
Erasmus MC	Pick Centrum	Behavioral variant of frontotemporal dementia
		Semantic dementia
		Progressive non-fluent aphasia
		Frontotemporal dementia with motor neuron disease
		Classical progressive supranuclear palsy
		Corticobasal degeneration
		Amyotrophic lateral sclerosis-parkinsonism-dementia complex
		Transmissible spongiform encephalopathy
Erasmus MC	Centre of Oligodontia	Oligodontia
Erasmus MC	Center for pediatric laryngotracheal stenosis	Congenital subglottic stenosis
		Laryngo-tracheo-esophageal cleft
		Congenital tracheal stenosis
Erasmus MC	ErasmusMC centre for endocrine disorders	Rare thyroid disease
		Rare hypothalamic or pituitary disease
		Rare adrenal disease
		Endocrine tumor
Erasmus MC	Hemophilia treatment center (volwassen)	Hemophilia
		Von Willebrand disease
		Rare hemorrhagic disorder due to a coagulation factors defect, incl FXIII, FXI, FVII,FV deficiency and alpha2-antiplasmin def.
		Rare hemorrhagic disorder due to a platelet anomaly
Erasmus MC	Expertise Center Rare hemorrhagic disorders (=Hemophilia treatment center, kind)	Hemophilia
		Von Willebrand disease
		Rare hemorrhagic disorder due to a coagulation factors defect, incl FXIII, FXI, FVII,FV deficiency and alpha2-antiplasmin def.
		Rare hemorrhagic disorder due to a platelet anomaly
Erasmus MC	Sickle cell center, volwassen	Sickle cell disease and related diseases
		Beta-thalassemia and related diseases
		Alpha-thalassemia and related diseases
Erasmus MC	Sickle cell center, kind	Sickle cell disease and related diseases
		Beta-thalassemia and related diseases
		Hemoglobinopathy
Erasmus MC	Leukemia and Stem cell transplantation center	Myeloid hemopathy
		Acute lymphoblastic leukemia (adult)
Erasmus MC	Multiple myeloma treatment center	Multiple myeloma
		Aggressive B-cell non-Hodgkin lymphoma
Erasmus MC	Paediatric Brain Tumour Center	Rare nervous system tumor
Erasmus MC	Brain Tumor Center	Glial tumor
		Tumor of the meninges
		Primary central nervous system lymphoma
		Primary germ cell tumor of the central nervous system
		Embryonal tumor of the neuroepithelial tissue
		Hemangioblastoma
		Craniopharyngioma
		Tumor of cranial and spinal nerves
Erasmus MC	Academic Breast Cancer Center	Hereditary Breast Cancer
Erasmus MC	Nasal, Paranasal, Ear and Skull base tumors workgroup	Tumors arising from the epithelium of the nasal, paranasal and skull base regions
		Esthesioneuroblastoma
Erasmus MC	Rotterdam Head and Neck Tumor work group	Oral and laryngyal squamous carcinoma
Erasmus MC	Soft tissue sarcoma center	Rare soft tissue tumor

Erasmus MC	Pediatric Surgical Centre for Anatomical Congenital Malformations	Congenital and syndromic diaphragmatic hernia
		Esophageal atresia
		Hirschsprung disease
		Anorectal malformation
		Omphalocele
		Gastroschisis
		Intestinal malformation
		Chronic intestinal failure
Erasmus MC	Erasmus MC Center for Pancreatic Diseases	Hereditary chronic pancreatitis
		Recurrent acute pancreatitis
		Autoimmune pancreatitis type 1
		Autoimmune pancreatitis type 2
		Familiar pancreascarcinoma
		Congenital pancreatic cyst
Erasmus MC	Rotterdam Oesophageal and Gastric Cancer Working Group	Esophageal carcinoma; incl. Barrett's oesophagus
Erasmus MC	Erasmus MC Liver Center	Hepatocellular adenoma
		Adult hepatocellular carcinoma
		Klatskin tumor
Erasmus MC	Immunodeficiency center	Primary immunodeficiency
Erasmus MC	Center for systemic allergic diseases	Systemic mastocytosis
Erasmus MC	Center of rare skin diseases	Netherton syndrome
		Suppurative hidradenitis
		Vascular anomaly or angioma
Erasmus MC	Centre of expertise for Children with Autoimmune Diseases	Juvenile idiopathic arthritis
Erasmus MC	Center for Rare Systemic Immune Disease	Rare systemic disease; Uveitis, Morbus Behcet, Morbus Sjogren, Systemic sclerosis
Erasmus MC	Turner Syndrome Center	Turner syndrome
Erasmus MC	Dutch Craniofacial Center	Isolated craniosynostosis
		Syndromic craniosynostosis; craniofrontonasal syndrome
		Pierre Robin syndrome associated with branchial archs anomalies
Erasmus MC	ErasmusMC Center for Congenital Hand and Upper Limb Malformations	Syndrome with limb malformations as a major feature
		Non-syndromic limb malformation
Erasmus MC	Expertise center DSD	Disorder of sex development
Erasmus MC	Expert Center spinal disraphism Rotterdam	Spina bifida aperta
Erasmus MC	Center for inherited cardiovascular diseases	Hypertrophic cardiomyopathy
		Unclassified cardiomyopathy
		Rare cardiac disease; rare familial occurrence of thoracic aortic abnormalities incl. dissection with (un)known genetic cause, e.g. Aneurysm-osteoarthritis syndrome
Erasmus MC	Center for congenital heart diseases Erasmus MC Rotterdam	Congenital heart malformation
Erasmus MC	Expert Center Prader Willi syndrome	Chromosomal anomaly; Prader Willi syndrome
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
		Non-acquired pituitary hormone deficiency; not yet known whether or which genes
		Non-acquired pituitary hormone deficiency; due to known GH gene and yet unknown genetic variants
		Growth hormone insensitivity syndrome
Erasmus MC	Expertise center Erasmus MC Vascular Genetics	Homozygous familial hypercholesterolemia
		Familial lipoprotein lipase deficiency
Erasmus MC	Erasmus MC Cystic Fibrosis Center	Cystic fibrosis
Erasmus MC	Pulmonary hypertension center	Rare respiratory disease

Erasmus MC	Interstitial Lung Disease Centre	Interstitial lung disease; adult
Erasmus MC	Sarcoidosis Centre ErasmusMC	Sarcoidosis
Erasmus MC	Mesothelioma centre	Mesothelioma
Erasmus MC	Center for Bronchopulmonary Dysplasia	Bronchopulmonary dysplasia
Erasmus MC	Erasmus MC Bone Center	Primary bone dysplasia
Erasmus MC	Center for Perinatal Psychiatry	Postpartum psychosis
Erasmus MC	Erasmus MC Leprosy Centre	Leprosy
Erasmus MC	Center for Pediatric Hematological Malignancies	Acute lymphoblastic leukemia
		Lymphoma; Hodgkin and non-Hodgkin Lymphoma
		Chronic myeloid leukemia
		Myelodysplastic syndromes
		Juvenile myelomonocytic leukemia
		Acute myeloid leukemia
LUMC	Center for Bone Quality	Sclerosteosis
		Primary bone dysplasia with decreased bone density
		Primary bone dysplasia with defective bone mineralization
		Fibrous dysplasia of bone
		Sternocostoclavicular Hyperostosis
		Rare parathyroid disease and phosphocalcic metabolism anomaly
LUMC	Nerve Centre	Nerve lesion
LUMC	Neuromuscular Center LUMC	Neuromuscular disease
		Duchenne en Becker muscular dystrophy
		Acquired neuromuscular junction disease
		Facioscapulohumeral dystrophy
		Oculopharyngeal muscular dystrophy
		Inclusion body myositis
LUMC	Huntington Disease Center Leiden	Huntington disease
LUMC	Cerebral Hereditary Angiopathy Center	CADASIL, and RVCL - HCHWA-D
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.
LUMC	Autonomic Disease Center	Pure autonomic failure, PAF
LUMC	Gender Clinic Leiden, WAKZ-Curium LUMC	Gender dysphoria
LUMC	Expertise center for monogenic diabetes mellitus	MODY syndrome
LUMC	Expertise center for lipodystrophy	Primary lipodystrophy
LUMC	Center for Endocrine Tumors Leiden (CETL)	Rare hypothalamic and pituitary disease (behalve MEN 1 en 2)
		Hereditary pheochromocytoma-paraganglioma
		Thyroid tumor
		Adrenocortical carcinoma
		Parathyroid carcinoma
LUMC	Expertise Center Genetics of growth	Growth disorders with a height <-3 SDS
		Growth disorders with a height >3 SDS
		Leri-Weill dyschondrosteosis/syndrome
		Disorders in the GH-IGF1 axis and signaling pathways
		IGSF1 deficiency syndrome
LUMC	Hemophilia treatment centre LUMC-Haga	Hemophilia A
		Hemophilia B
		Von Willebrand disease
		Acquired hemophilia
		Acquired von Willebrand syndrome
LUMC	Expert center for aplastic anemia	Idiopathic aplastic anemia
LUMC	Expert center for hemoglobinopathies	Hemoglobinopathy, incl alpha- beta-thalassemia, Sickle Cell Disease, HbS and Hb variants
LUMC	Expert center for pediatric stem cell transplantation	Combined T and B cell immunodeficiency, mainly SCID and ICF syndr
		Primary immunodeficiency due to a defect in adaptive immunity

		Acute graft versus host disease
LUMC	Bone and soft tissue tumour clinic	Soft tissue sarcoma; Gastrointestinal stromal (cell) tumour
LUMC	Bone and soft tissue tumour clinic	Multiple osteochondromas
		Adamantinoma
		Rare bone tumor
		Chondromyxoid fibroma
		Osteosarcoma
		Ewing sarcoma
		Chondrosarcoma
		Giant cell tumor of bone
		Rare soft tissue tumor
LUMC	Leiden Ocular Oncology Center	Uveal melanoma
LUMC	Female Cancer Center - Leiden (FCC-L)	Rare cancer of the cervix uteri
		Rare vulvovaginal tumor; Vulvar cancer
		Rare cancer of the corpus uteri
		Rare ovarian cancer
LUMC	Expertise Center Pediatric ophthalmology	Retinopathy of prematurity
LUMC	Expertise Center Fetal medicine	Hemolytic disease due to fetomaternal alloimmunization
		Twin to twin transfusion syndrome
		Fetal and neonatal alloimmune thrombocytopenia
		Fetal parvovirus syndrome
		Hydrops fetalis
		Congenital heart malformation; fetal cardiac interventions
		Posterior urethral valve; Lower Urinary Tract Obstruction
		Non-syndromic respiratory or mediastinal malformation
LUMC	Prenatal and congenital infections by cytomegalovirus and parvovirus B19	Infectious embryofetopathy; Congenital CMV inf. and fetal and congenital parvovirus B19 inf.
LUMC	Expert center for cutaneous lymphomas	Primary cutaneous lymphoma
LUMC	Expertise Center Rare autoinflammatory diseases	Systemic sclerosis
		Juvenile rheumatoid factor-negative polyarthritis with anti-nuclear antibodies
LUMC	Congenital Heart malformations (CAHAL pediatric)	Congenital heart malformation; pediatric
		Rare cardiac rhythm disease; non-genetic
LUMC	Marfan-FTAAD Clinic	Marfan syndrome; including neonatal Marfan syndrome, FTAAD
LUMC	Hereditairy bowel cancer centre	Hereditary nonpolyposis colon cancer; Lynch caused by MLH1 or MSH2 mutation
		Hereditary nonpolyposis colon cancer; Lynch caused by MSH6 or PMS2 mutation
		MUTYH-related attenuated familial adenomatous polyposis
		Familial adenomatous polyposis; APC associated polyposis
LUMC	Center for Complement mediated renal disease	C3 glomerulonephritis
LUMC	Center for Inherited kidney disease	Autosomal recessive polycystic kidney disease
		Autosomal Dominant Polycystic Kidney Disease, PKD1 mutation
		Autosomal Dominant Polycystic Kidney Disease PKD2 mutation
		Autosomal Dominant Medullairy Cystic Kidney Disease
LUMC	Alpha1 International Registry (AIR)	Alpha-1-antitrypsin deficiency
LUMC	Center for Narcolepsia	Narcolepsy-cataplexy
LUMC	Expert center for familial cutanous melanoma	Familial melanoma; incl. FAMMM syndrome and FAMMMPC syndrome
MUMC+	Expertise Center Galactosemia	Galactosemia
MUMC+	Expertise Center Hyperostosis of the skull	Osteopetrosis
MUMC+	Expertise Center Huntington's disease	Huntington disease
MUMC+	Neuromuscular Centre MUMC+	Neuromuscular disease
		Myotonic dystrophy
		Sodium channelopathy-related small fiber neuropathy
		Duchenne and Becker muscular dystrophy

MUMC+	Expertise Center Neural tube defects	Neural tube defect
MUMC+	Academic Center for Epilepsy	Epilepsy syndrome
		Continuous spikes and waves during sleep, epileptic encephalopathy
		Early infantile epileptic encephalopathy
MUMC+	Expertise Center Cerebral Palsy	Spastic diplegia - infantile type. 1. spastic unilateral cerebral palsy 2. dyskinetic cerebral palsy
MUMC+	Expertise Center Hereditary Tumors	Hereditary breast and ovarian cancer
MUMC+	Expertise Center Genodermatoses	Inherited ichthyosis
		Erythrokeratoderma
		Lymphedema
		Birt-Hogg-Du syndrome
		Rare genetic skin disease
MUMC+	Expertise center Rare syndromes and cognitive disorders	Rare developmental defect during embryogenesis
		Kabuki syndrome
		Rett Syndrome
MUMC+ & Radboudumc	Marfan and related disorders policlinic	Marfan syndrome
		Loeys-Dietz syndrome
		Familial thoracic aortic aneurysm and aortic dissection
MUMC+	Expertise center Cardiogenetics MUMC+	Rare familial disorder with hypertrophic cardiomyopathy
		Dilated cardiomyopathy
		Arrhythmogenic right ventricular dysplasia
		Familial long QT syndrome
		Idiopathic ventricular fibrillation - not Brugada type
		Brugada syndrome
MUMC+	Expertise center Pulmonary hypertension	Pulmonary hypertension with unclear multifactorial mechanism; auto immune mechanisms in PH, and right ventricular failure
MUMC+	Limburg renal registry	EGPA, GPA and MPA
		Cryoglobulinemic vasculitis
MUMC+	Gastro-intestinal center Maastricht	Biliary tract carcinoma en Hepathocellular carcinoma
MUMC+	Maastricht Soft Tissue Tumor Center	Rare soft tissue tumor; long list, together all soft tissue sarcomas in adults
MUMC+	Maastricht Gynaecological Oncology Center	Rare ovarian cancer; epithelial and non- epithelial, also tumor of Fallopian tubes
		Rare cancer of the corpus uteri
MUMC+	Center for Endocrine tumors	Thyroid carcinoma
MUMC+	Neuroendocrine tumours Center	Bronchial NET en Merkelcell carcinoma
MUMC+	Neuro-oncologie centrum Maastricht	Glial tumor
		Tumor of cranial and spinal nerves
MUMC+	Maastricht Head & Neck Cancer Center	Rare otorhinolaryngologic tumor; as a group
MUMC+	Lung cancer center maastricht	Small cell lung cancer
Radboudumc	Nijmegen centre for mitochondrial disorders	Mitochondrial disease
Radboudumc	Nijmegen Center for Disorders of Glycosylation (NCDG)	Congenital disorder of glycosylation
Radboudumc	Centre for genetic movement disorders	Rare hereditary ataxia, mainly autosomal dominant and recessive cerebellar ataxias
		Hereditary spastic paraplegia
		Mainly Sjogren-Larsson syndrome, GLUT1 deficiency syndrome, and disorders of dopamine metabolism
Radboudumc	Neuromuscular Centre	Neuromuscular disease
		Facioscapulohumeral dystrophy
		Oculopharyngeal muscular dystrophy
		Duchenne and Becker muscular dystrophy
		Myotonic dystrophy
		Neuralgic amyotrophy
		Idiopathic inflammatory myopathy
		Congenital myotonia and paramyotonia congentia
		Non-dystrophic myopathy

Radboudumc	Center for rare CNS and retinal vascular disease	Acquired aneurysmal subarachnoid hemorrhage
Radboudumc	Hearing & Genes Centre	Rare genetic deafness
		Usher syndrome
Radboudumc	Radboud Adrenal Centre	Cushing syndrome
		Adrenogenital syndrome
		Rare primary hyperaldosteronism
		Primary adrenal insufficiency
		Adrenal/paraganglial tumor, incl Von Hippel Lindau and MEN-2 syndrome
		Adrenal/paraganglial tumor; except catecholamines, aldosterone or cortisol producing . Incl. incidentalomas and carcinomas
Radboudumc	Centre of Paroxysmal Nocturnal hemoglobinuria	Paroxysmal nocturnal hemoglobinuria
Radboudumc	Haemophilia treatment centre	Rare coagulation disorder; hemophilia
		Rare coagulation disorder; von Willebrand Disease
		Rare coagulation disorder; other
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
		Sideroblastic anemia
		Constitutional anemia due to iron metabolism disorder
		Constitutional dyserythropoietic anemia, mainly type I-IV
Radboudumc	Hereditary cancer centre	Hereditary breast and ovarian cancer syndrome; BRCA mutation carriers
		Hereditary nonpolyposis colon cancer; Lynch Syndrome
		Familial gastric cancer, incl her. diffuse GC
		APC-related attenuated familial adenomatous polyposis
Radboudumc	Centre for thyroid carcinomas	Thyroid tumor
Radboudumc	Radboud Skull base centre	Rare tumor; different very rare skull base tumors
		Rare nervous system tumor; acoustic neuroma; cerebello pontine angle tumor; excl. NF2 patients
		Tumor of endocrine glands; Hereditary pheochromocytoma-paraganglioma
Radboudumc	Center for colorectal surgery	High anorectal malformation
		Intermediate anorectal malformation
		Low anorectal malformation
		Anal fistula
		VACTERL/VATER association
		Caudal regression sequence
		Hirschsprung disease
		Cloacal exstrophy
Radboudumc	Ophthalmogenetic center	Genetic vitreous-retinal disease
		Choroideremia
		Stargardt disease and other ABCA4-related diseases
		Retinal dystrophy; central serous retinopathy
		Leber congenital amaurosis
Radboudumc	Liver cyst center	Isolated polycystic liver disease
Radboudumc	Nijmegen Center for immunodeficiency and autoinflammation	Primary immunodeficiency
		Autoinflammatory syndrome with immune deficiency
		Immunodeficiency due to a complement cascade protein anomaly
		Schnitzler syndrome
		Mevalonate kinase deficiency = Hyper IgD Syndrome (HIDS)
Radboudumc	Mycology reference center	Aspergillosis; chronic aspergillosis and ABPA
		Chronic mucocutaneous candidiasis
		Rare mycosis; in patients with hyper IgE syndrome
		Rare mycosis; in patients with chronic granulomatous dis.
		Aspergillosis
Radboudumc	Radboud Center Renal Disorders	Glomerular disease
		Rare renal tubular disease
		Familial cystic renal disease; all cystic kidney dis. in children, incl. ciliopathies/nephronophtises
		Renal or urinary tract malformation
		Cystinosis
		Thrombotic microangiopathy
Radboudumc	Centre for genetic neurodevelopmental disorders	Kleefstra syndrome due to a point mutation
		Koolen-de Vries syndrome

		Cowden syndrome
		KBG syndrome
		Noonan syndrome
		Prader Willi Syndrome
Radboudumc	Craniofacial team Nijmegen	Cranial malformation
		Craniosynostosis
Radboudumc	Radboud DSD centre	46 -XX disorder of sex development induced by fetal androgens excess
		46 -XY disorder of sex development
		Turner syndrome
Radboudumc & MUMC+	Marfan and related disorders policlinic	Marfan syndrome
		Loeys-Dietz syndrome
		Familial thoracic aortic aneurysm and aortic dissection
Radboudumc	Centre for Hemangiomas and Congenital Vascular Anomalies Nijmegen (HECOVAN)	Vascular tumor; incl complicated hemangiomas
		Venous malformation
		Lymphatic system malformation
		Arteriovenous malformation
		Complex - combined vascular malformation; incl Klippel-Trenaunay-syndrome
Radboudumc	(Pediatric) urology center	Bladder exstrophy; incl. cloacal exstrophy and epispadias
		Agenesis and aplasia of uterine body; Cloacal anomalies and anal atresia in combination with uronary tract anomalies
		Non-syndromic urogenital tract malformation of male and female; e.g. in spina bifida
		Posterior urethral valve
		Posterior hypospadias
Radboudumc	Radboud Intestinal failure Unit	Chronic intestinal failure
Radboudumc	Cleft (lip and) palate center Nijmegen	Cleft palate; cleft hard and / or soft palate
		Cleft palate; cleft lip/palate
		Cleft palate; Submucosal cleft palate
		Oculo-auriculo-vertebral spectrum (=hemifaciale microsomia)
Radboudumc	Radboudumc Center for Pulmonary Hypertension	Idiopathic pulmonary arterial hypertension; all kinds of PH
Radboudumc/UCCZ Dekkerswald	Center for Mycobacterial diseases	Tuberculosis; also incl. nontuberculous mycobacterial inf.
Radboudumc	Radboud Center for Congenital Diaphragmatic Hernia and neonatal pulmonary hypertension	Congenital diaphragmatic hernia; non- syndromic and syndromic
		Pulmonary hypertension owing to lung disease and/or hypoxia; Persistent Pulmonary Hypertension of the newborn
Radboudumc	Radboud Pituitary Center	Cushing disease
		Somatotropic adenoma
		Prolactinoma; also incl. other rare types of functioning pituitary tumor
		Pituitary deficiency; also incl. all hypothalamic and pituitary diseases resulting in pituitary def.
Radboudumc	Radboud Sarcoma Center	Bone sarcoma; incl. bone and soft tissue tumors and GIST (gastointestinal stromal tumors)
Radboudumc	Radboud Centre for vulvar and cervix cancer and Dutch Mole Registry	Gestational trophoblastic neoplasm; all different subgroups
		Vulvar intraepithelial neoplasia; rare vulvar cancers and VIN
Radboudumc	Radboudumc Neuro-oncological center	Tumor of the neuroepithelial tissue
		Primary melanocytic tumor of the CNS
Radboudumc	Center for Head and Neck Oncology	Squamous cell carcinoma of head and neck
		Malignant epithelial tumor of the salivary glands
UMCG	Expert centre for Phenylketonuria (PKU) and Tyrosinemia type I	Disorder of phenylalanin or tyrosine metabolism
UMCG	Expert centre for hepatic Glycogen Storage Diseases	Glycogen storage disease due to glucose-6-phosphatase deficiency type a
		Glycogen storage disease due to glucose-6-phosphatase deficiency type b
		Glycogen storage disease due to glycogen debranching enzyme deficiency
		Glycogen storage disease
		Glycogen storage disease due to liver glycogen phosphorylase deficiency
UMCG	Expert centre for M(C)ADD	Medium chain acyl-CoA dehydrogenase deficiency

		Multiple acyl-CoA dehydrogenation deficiency - severe neonatal type
		Multiple acyl-CoA dehydrogenation deficiency - mild type
UMCG	Expert centre for serine deficiencies	Neurometabolic disorder due to serine deficiency
UMCG	Neurovascular Team UMCG	Cerebral malformation, intracranial dural AV-fistula
		Spinal arteriovenous shunts
		Cerebral malformation, brainstem cavernomas
		Cerebral malformation, proliferative angiopathy
UMCG	Expert centre for movement disorders in adults and children	Rare dystonia: myoclonus, focal,generalised, dopa responsive
		Hyperekplexia
		Neurodegeneration with brain iron accumulation; a.o. PKAN
		Rare myoclonus; myoclonus dystonia GOSR2 and FCMTE
		Psychogenic movement disorders
		Autosomal dominant cerebellar ataxia, incl recessive ataxias
		Paroxysmal dyskinesia
		Inherited congenital spastic tetraplegia
		Rare choreic movement disorder, huntington's disease
		Neurometabolic disease, related tot movement disorders
UMCG	Expertise Center Groningen Papilloma studies	Recurrent respiratory papillomatosis
UMCG	Adrenal centre UMCG	Catecholamine-producing tumor, incl. pheochromocytoma/ paraganglioma and non-secreting head and neck paragangliomas
		Rare primary hyperaldosteronism
UMCG	Hemophilia Treatment Centre UMCG	Rare hemorrhagic disorder; hemophilia
		Rare hemorrhagic disorder; von Willebrand Disease
		Rare hemorrhagic disorder: other allied bleeding disorders
UMCG	Expert centre mastocytosis Netherlands (ECMN)	Mastocytosis
UMCG	Expert Center head- and neck oncology	Squamous cell carcinoma of head and neck
UMCG	Expert Center neuro-oncology in adults	Glial tumor
UMCG	Expert centre for carcinoid / neuroendocrine carcinoma's (NEC)	Carcinoid tumor and carcinoid syndrome
UMCG	Centre of familial tumors	Inherited cancer-predisposing syndrome; incl VHL, MEN1, MEN2 and familial paraganglioma/PCC.
UMCG	Thyroid cancer centre	Thyroid tumor
UMCG	Familial Breast Ovarian Cancer Clinic	Hereditary breast and ovarian cancer syndrome; BRCA1 BRCA2
UMCG	Familial Colorectal Cancer Clinic	Hereditary nonpolyposis colon cancer
		Familial adenomatous polyposis; incl FAP and MUTYH
UMCG	Expertise Center Germcell tumors	Testicular germ cell tumor
UMCG	Expertise Center Soft tissue and bone tumors	Osteosarcoma
		Soft tissue sarcomas
UMCG	Pediatric Neuro-oncology Team	Medulloblastoma; and PNET
		Rare nervous system tumor
UMCG	Expertise Center Gyneco-oncology UMCG	Rare vulvovaginal tumor; squamous cell carcinoma of vulva
		Rare cancer of the cervix uteri; incl. squamous cell carcinoma
		Malignant epithelial tumor of ovary; different types adenocarcinoma
UMCG	UMCG/ oesophageal/ gastric cancer tumorgroup	Esophageal adenocarcinoma
UMCG	Small bowel rehabilitation and transplant centre	Chronic intestinal failure
UMCG	Expertise Center pediatric liver disease, pediatric liver surgery and pediatric liver transplantation	Biliary atresia
UMCG	Paediatric centre for Rheumatologic and immunologic diseases	Juvenile idiopathic arthritis
UMCG	Tuberculosis centre Beatrixoord	Tuberculosis
UMCG	Expertise Center for Polycystic Kidney Diseases	Familial cystic renal disease
UMCG	Center for Blistering Diseases	Inherited epidermolysis bullosa

		Autoimmune bullous skin disease; all forms of pemphigus and pemfigoid
UMCG	Groningen Unit for Amyloidosis Research & Development (GUARD)	Primary systemic amyloidosis
		Primary localized amyloidosis
		Secondary amyloidosis
		Familial amyloid polyneuropathy
		Transthyretin-related familial amyloid cardiomyopathy
		Senile systemic amyloidosis
UMCG	Expert centre for systemic vasculitis	Vasculitis
UMCG	Expert centre Sjögren syndrome	Systemic autoimmune disease; Sjögren (including MALT lymphoma)
		Tumor of hematopoietic and lymphoid tissues; MALT lymphoma associated with Sjögren's disease
UMCG	Clinic for Connective tissue disorders	Marfan syndrome
UMCG	The multidisciplinary CHARGE clinic	CHARGE syndrome
UMCG	Clinic for rare chromosome disorders	Autosomal anomaly; wide diversity of chromosomal deletions and duplications (and not the more common trisomies)
UMCG	Expert Centre for Cardiogenetics	Familial isolated arrhythmogenic ventricular dysplasia - biventricular form
UMCG	Expert Center for Children and Adults with rare Congenital Heart Diseases	Congenital heart malformation
UMCG	Dutch expertise centre for lympho-vascular medicine	Primary lymphedema
		Syndromic lymphedema
UMCG	Expert Center for Children and Adults with Pulmonary Hypertension	Pulmonary arterial hypertension
UMCG	Cystic Fibrosis centre Groningen	Cystic fibrosis
UMCG	UMCG Pituitary Center	Rare pituitary disease
UMCG	Pediatric Oncology group UMCG	Tumor of hematopoietic and lymphoid tissues
UMCU	Center Inherited Metabolic Diseases	Disorder of fatty acid oxidation and ketone body metabolism
		Disorder of pyridoxine metabolism
UMCU	Brain Centre Rudolf Magnus, Neuromuscular Diseases	Neuromuscular disease
		Spinal muscular atrophy
UMCU	Center for Refractory Pediatric Epilepsy	Rare epilepsy
		Tuberous sclerosis
		Continuous spikes and waves during sleep and ESES
		Epilepsy syndrome
UMCU	Center of Excellence in Congenital Orofacial and Dental Anomolies	Rare odontologic disease; oligodontia
		22q11.2 deletion syndrome
		Isolated Pierre Robin syndrome
		Hemifaciale microsomie & Microtie
UMCU	Expertise centre for benign hematology, thrombosis and hemostasis, Van Creveld clinic	Rare hemorrhagic disorder due to a coagulation factors defect; hemophilia
		Rare hemorrhagic disorder due to a coagulation factors defect; von Willebrand Disease
		Rare hemorrhagic disorder due to a coagulation factors defect; deficiency of factor II/ V/ VII/X/XI
		Rare hemorrhagic disorder due to a coagulation factors defect; antiplasmin deficiency
		Rare anemia
		Rare constitutional medullar aplasia; Fanconi, Diamond-Blackfan anemia and congenital neutropenia
		Rare hemorrhagic disorder; Congenital and acquired platelet disorders
UMCU	Expertise centre for malignant hematology	Multiple myeloma
		Non Hodgkin lymphoma
		Acute lymphoblastic and myeloid leukemia
UMCU	Centre for rare tumors	Inherited cancer-predisposing syndrome; Men 1
		Multiple endocrine neoplasia type 2A; incl. fam medullary thyroid carcinoma, MEN2B and sporadic medulary thyroid carcinoma
		Von Hippel-Lindau disease

UMCU	Expertise Center Rare GI and hepatic diseases	Progressive familial intrahepatic cholestasis
		Wilson disease
		Intractable diarrhea of infancy; due to genetic defects
UMCU	UMCU ophthalmology uveitisgroup	Anterior uveitis; (non) infectious
		Posterior uveitis; (non) infectious and in syst dis.
		Systemic diseases with panuveitis
		Rare inflammatory eye disease
		Intermediate uveitis; (non) infectious
UMCU	Expertise centre for primary immunodeficiences	Immunodeficiency predominantly affecting antibody production; incl. CVID, XLA, other types of complete agammaglobulinemia
		Immunodeficiency predominantly affecting antibody production; incl. SADNI, IgG subclass- and IgA def.
		Primary immunodeficiency due to a defect in adaptive immunity; B and T cell immunodeficiencies: SCID, CID
		Primary immunodeficiency due to a defect in innate immunity
		Primary hemophagocytic lymphohistiocytosis
		Graft versus host disease
UMCU	Expertise centre systemic autoimmune diseases	Juvenile idiopathic arthritis
		Periodic fever syndrome, incl. CAPS, FMF, behcet, Traps, Pfapa
		Juvenile dermatomyositis
		Pediatric systemic lupus erythematosus
		Vasculitis
		Systemic sclerosis
		Rare coagulation disorder; Antiphospholipid syndrome
		Systemic autoimmune disease; extraglandular manifestations in Sjogren
UMCU	WKZ center for congenital malformations	Esophageal atresia
UMCU	Centre for skeletal malformations	Rare bone disease; m.n. primaire skeletdysplasieen en dysostoses, ook osteogenesis imperfecta en achondroplasie
UMCU	Children's Heartcenter WKZ	Congenital heart malformation
UMCU	Clinic for Tuberous Sclerosis Complex	Rare genetic neurological disorder; Tubererous Sclerosis Complex
UMCU	Cystic Fibrosis Clinic	Cystic fibrosis
UMCU	Expert Centre Hereditary and congenital nephrologic and urologic disorders	Rare renal disease; congenital or inherited renal or urinary tract disease
		Familial cystic renal disease; nephronophthisis, as feature of i.e. Joubert and Meckel or isolated or part of other ciliopathies.
		Non-syndromic renal or urinary tract malformation; CAKUT
UMCU	Center for Rare Ear and Hearing Diseases	Middle ear anomaly
UMCU & NKI-AvL	Expert Center of Neuroendocriene carcinomas	Gastroenteropancreatic endocrine tumor
UMCU	Center of vascular anomalies Utrecht	Vascular anomaly
VUmc	Centre for Genetic Metabolic Diseases Amsterdam (CGMA) - VUmc specific	Disorder of creatine biosynthesis
VUmc	Center for Childhood White Matter Disorders	Rare neurologic disease
VUmc	Expertise Center for Osteogenesis Imperfecta	Primary bone dysplasia with decreased bone density
VUmc	Center for obstetric bracial plexus lesion	Rare neurologic disease; obstetric brachial plexus lesions
VUmc	Center for pediatric rehabilitation medicine	Spastic diplegia - infantile type
VUmc	Center for pediatric oral and maxillofacial surgery	Robin sequence
VUmc	Expertise Center Head and Neck tumors	Nasopharyngeal carcinoma
VUmc	Celiac disease center	Refractory celiac disease
VUmc	Center for rare haematologic cancers	Multiple myeloma
		Non Hodgkin lymphoma
		Acute lymphoblastic and myeloid leukemia
VUmc	Dutch Center for Oral Medicine and Oral Pathology	Squamous cell carcinoma of head and neck; malignant disorders of oral mucosa
		Squamous cell carcinoma of head and neck; Ameloblastomas and Keratocystic odontogenic tumors, Gorlin-Goltz syndrome

VUmc	Brain Tumor Center	Rare tumor; Brain Tumors
VUmc	Dutch Retinoblastoma Center	Retinoblastoma
VUmc	Expertise Center Neurophthalmology	Neuromyelitis optica: autoimmune optic neuropathies incl. NMO, CRION, RION, ION, MSON
		Paraneoplastic neurologic syndrome; optic neuropathies relevant to DD of CRION, RION and NMO-SD
		Adult-onset myasthenia gravis; ocular MG
		Idiopathic intracranial hypertension; loss of vision due to IIH
		Acute zonal occult outer retinopathy
VUmc	Expertise Center Preeclampsia from origin to healhy aging	Preeclampsia
VUmc	Centre for Systemic Sclerosis and Systemic Lupus Erythematosus, embedded in Amsterdam Rheumatology and immunology Centre	Systemic sclerosis
VUmc	Pediatric nephrology centre	Idiopathic nephrotic syndrome
		Non-syndromic renal or urinary tract malformation; CAKUT
VUmc	Cleft Lip and Palate Team	Rare developmental defect during embryogenesis; cleft lip and palate
VUmc	Birt-Hogg-Dubé task force	Inherited renal cell cancer-predisposing syndrome; Birt-Hogg-Dubé syndrome
VUmc	Expertcenter for Fybrodysplasia Ossificans Progressiva	Fibrodysplasia ossificans progressiva
VUmc	VUmc multidisciplinary (genetic) breast cancer team	Hereditary breast cancer
VUmc	Down Center the Netherlands, location West	Down syndrome
VUmc	VUMC PH Centre	Rare pulmonary hypertension; many subforms of PH; associated: collagen vascular disease, Hereditary PH, IPAH, type III WHO associated with emfysema.
		Chronic thromboembolic pulmonary hypertension
Vumc & AMC	Cystic Fibrosis Centre Amsterdam	Cystic Fibrosis
VUmc	PCD-center Vumc	Primary ciliary dyskinesia
VUmc	Fetal Akinesia Deformation Sequence Centre	Fetal Akinesia Deformation Sequence
NKI-AvL	Sarcoma Expertise Centre Amsterdam	Soft tissue sarcoma
		Rare soft tissue tumor; Gastrointestinal stromal tumor (GIST)
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
		Familial gastric cancer
NKI-AvL& UMCU	Expert Center of Neuroendocriene carcinomas	Gastroenteropancreatic endocrine tumor
NKI-AvL	Expert Centre of rare thoracic tumours	Mesothelioom
NKI-AvL	Expert centre for rare urological diseases	Testicular cancer
		Penile cancer
NKI-AvL	Center of rare head and neck tumours	Rare otorhinolaryngologic disease; Tumours of the head and neck
		Rare otorhinolaryngologic disease;Tumours of the salivary glands
NKI-AvL	Expert Center for Hereditary Cancer	Li-Fraumeni syndrome
		Hereditary breast and ovarian cancer syndrome
Bartiméus	Bartiméus Diagnostisch Centrum	Genetic vitreous-retinal disease
		Congenital stationary night blindness
		Oculocutaneous or ocular albinism
		Cerebral visual impairment in children
Kempenhaeghe	Center of Sleep Medicine Kempenhaeghe	Rare Sleep disorders
The Rotterdam Eye Hospital	Rare Eye Disease Center Rotterdam	Uveal melanoma
		Central serous chorioretinopathy
		Rare acquired eye disease; Herpes simplex virus keratitis, stromal, neurtrophic and endotheliitis
		Uveitis

		Rare genetic eye disease; Retinal Dystrophies
STZ-Albert Schweitzer hospital, Dordrecht	Centre of expertise Retroperitoneal Fibrosis	Retroperitoneal fibrosis
STZ-OLVG	Center for HME-MO (Hereditaire Multipele exostosen-Multipele Osteochondromen)	Multiple osteochondromas; Hereditary Multiple Exostoses - Multiple Osteochondromas
STZ-OLVG	EC for interstitial lungdiseases OLVG	Idiopathic pulmonary fibrosis en Idiopathic interstitial pneumonia
		Exposure-related interstitial lung disease; in its broadest sense
STZ-St. Antonius Ziekenhuis	Interstitial Lung Diseases Center of Excellence	Interstitial lung disease
		idiopathic pulmonary fibrosis (IPF)
		hypersensitivity pneumonitis (PH)
		sarcoidosis
STZ-St. Antonius Ziekenhuis	Center for Pulmonary vascular diseases	Hereditary hemorrhagic telangiectasia
		Chronic thromboembolic pulmonary hypertension
		Idiopathic pulmonary arterial hypertension
STZ - St. Antonius Ziekenhuis	St. Antonius Oesofagus Centrum	Esophageal carcinoma
STZ-CWZ Nijmegen	Malignant Hyperthermia investigation unit Nijmegen	Malignant hyperthermia
STZ-Maxima Medisch Centrum	SolviMáx, Center of Excellence for Abdominal Wall and Groin Pain	Anterior cutaneous nerve syndrome (ACNES)
STZ-Maxima Medisch Centrum	Center for Adrenal Tumors	Adrenocortical carcinoma
SZT-St. Elisabeth Ziekenhuis	Neurovascular Center Tilburg	Neurovascular malformation
SZT-St. Elisabeth Ziekenhuis	Neurovascular Center Tilburg	Acquired aneurysmal subarachnoid hemorrhage
STZ- St Elisabeth Hospital Tilburg	Neuro-oncology Center Tilburg	Glial tumor
STZ-Jeroen Bosch Hospital	Center for Primary immunodeficiencies	Immuno-deficiency predominantly affecting antibody production, mainly the various types of unclassified antibody deficiency (voorlopig erkenning t/m sep 2017)
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
STZ-Medisch Centrum Haaglanden-Bronovo-Nebo	Center for Neuro-oncology The Hague	Gliomen