Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	
		Lysosomal Disease	
		Cholesteryl ester storage disease	
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 disease and alfa or beta thalassemia	
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	voor 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	Pancreatic tumor	voor zeldzame cysteuze pancreas tumoren
		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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
AMC	Center for immune-mediated and genetic cholestasis syndromes	Primary biliary cirrhosis	
		Primary sclerosing cholangitis	
		Chronic autoimmune hepatitis	Voorlopig*
		Familial intrahepatic cholestasis	Voorlopig*
		Benign recurrent intrahepatic cholestasis	Voorlopig*
		IgG4-related disease	Voorlopig*
		Crigler-Najjar syndrome	Voorlopig*
		Crigler-Najjar syndrome type 1	
		Crigler-Najjar syndrome type 2	
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	Rare rheumatologic disease; focus on all forms of juvenile idiopathic arthritis, reactive arthritis and all forms of autoinflammatory syndromes	voor JIA
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	
		Hypertrophic cardiomyopathy	
AMC	Centre for Marfan syndrome	Marfan syndrome	
AMC	Amsterdam Expert Center for Developmental Disorders	Cornelia de Lange syndrome	
		Rubinstein-Taybi syndrome	
		Pitt-Hopkins syndrome	
		Marshall-Smith syndrome	
		Hutchinson-Gilford progeria	Voorlopig*
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; Chronic Toxic Encephalopathy	voor rare intoxication
		Manganese poisoning	
		Lead poisoning; and other orphanet rare intoxications incl. different poisonings	
AMC	Center for Idiopathic Nephrotic Syndrome	Primary glomerular disease; Steroid -sensitive and steroid-resistant nephrotic syndrome, incl. congenital nephrotic syndrome, membrano proliferative glomerulonefritis	voor idiopathic nephrotic syndrome
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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
AMC	Amsterdam Multidisciplinary Lyme borreliosis Center	Lyme disease	
AMC	Center for Paediatric oncology	Osteosarcoma, incl. ewing sarcomas and langer hans cell bone lesions	voor osteosarcoma
		soft tissue sarcomas (rhabdomyosarcoma and non rhabdo myosarcoma	voor soft tissue sarcomas
AMC	Expert Centre for congenital anomalies of the urinary tract EKZ-AMC	Non-syndromic renal or urinary tract malformation; CAKUT	
AMC & VUmc	Amsterdam Centre of congenital malformations	Hirschsprung Disease	
		Esophageal atresia	
		Anorectal malformation	
		Sacrococcygeal teratoma	
		Chronic intestinal pseudoobstruction	
AMC	Center for Osteochondral Defects of the Talus	Osteochondritis van tarsaal/metatarsaal bot	
AMC	Center for condylar hyperplasia	Temporomandibular joint anomaly	Voorlopig*
		Condylaire hyperplasie	Voorlopig*
AMC	Amsterdam expert center for bronchopulmonary dysplasia	Bronchopulmonary dysplasia	
AMC & VUmc	Amsterdam Center for ILD and sarcoïdosis	Sarcoidosis	Voorlopig*
AMC & VUmc & LUMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	
		Centrale sereuze chorioretinopathie	
AMC	Center for rare movement disorders	Primary orthostatic tremor	
		Benign adult familial myoclonic epilepsy	
AMC	Center for congenital nevi	Large congenital melanocytic nevus	Voorlopig*
AMC	Center for hematological immune diseases Amsterdam	Autoimmune hemolytic anemia (AIHA)	
AMC	Melioidosis Expertise Center	Melioidosis	Voorlopig*
AMC	LYMMCARE Amsterdam (Lymphoma and Myeloma CAre and REsearch)	Waldenstrom macroglobulinemia	
		B-cell chronic lymphocytic leukemia	
		Lymphoma	
		Multiple myeloma	
AMC	Centre for graft versus host disease Amsterdam	Graft versus host disease (acuut en chronisch)	
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 for PKU, 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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		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)	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	Rare tumor; Hereditary Breast Cancer	
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	voor tumors arising from the epithelium of the nasal, paranasal and skull base regions
		Esthesioneuroblastoma	
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	voor 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	
		Pancreatic tumor (2x)	voor familiair pancreascarc.
		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	
		Rare hepatic and biliary tract tumor	
Erasmus MC	Immunodeficiency center	Primary immunodeficiency (*)	
Erasmus MC	Center for systemic allergic diseases	Rare immune disease; Systemic mastocytosis	
Erasmus MC	Center of rare skin diseases	Netherton syndrome	
		Systemic disease with skin involvement; Localized scleroderma and Suppurative hidradenitis	voor Supp. 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	Craniofacial Center	Isolated craniosynostosis	
		Syndromic craniosynostosis; craniofrontonasal syndrome	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		Treacher-Collins syndrome; incl. Nager and Burn-McKeown syn	voor Pierre Robin syndrome associated with branchial archs anomalies
		Cleft palate; incl cleft palate only and Pierre Robin sequence (eg Stickler, van der Woude, Wolf Hirschhorn)	
		Rare bone development disorder; involving craniofacial presentation of fibrous dysplasia, McCune Albright, cherubism, cleidocranial dysplasia	
		Acalvaria; including ossification defects of the skull with or without involvement of skin	
		Goldenhar syndrome; incl. hemifacial microsomia	
		Facial cleft	
		Cleft lip with or without cleft palate	
		Cleft palate	
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	voor Syndrome with limb malformations as a major feature
		Multiple congenital anomalies/dysmorphic syndrome without intellectual disability; Radiusdysplasie (Fanconi, TAR etc), Brachydactylie, Holt-Oram/Hart-Hand	
		Syndrome with limb malformations as a major feature; Arthrogryposis, Syndromale polydactylie (o.a. Greig) Transversaal reductie defect	
		Dysostosis with limb anomaly as a major feature	
		Non-syndromic limb malformation	voor Non-syndromic limb malformation
		Non-syndromic limb malformation; Macrodactylie handen voeten geïsoleerd, Hemihypertrofie, Musculaire hyperplasie	
Erasmus MC	Expertise center DSD	Disorder of sex development	
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	voor 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: intervention for diseases of D-2-8, D-6-4, D-6-11, D-6-10, D-6-3; A-5-7	
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	
		Malignant peritoneal mesothelioma	
		Thymoma	
		Small cell lung cancer	
Erasmus MC	Center for Bronchopulmonary Dysplasia	Bronchopulmonary dysplasia	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
Erasmus MC	Erasmus MC Bone Center	Primary bone dysplasia; incl. a.o. list of 6 specific forms of dysplasia	voor primary bone dysplasia
Erasmus MC	Center for Perinatal Psychiatry	Rare disorder related with pregnancy - childbirth and puerperium; Postpartum psychosis	
Erasmus MC	Erasmus MC Leprosy Centre	Leprosy	
Erasmus MC	Academic Center Kidney & Hypertension	rare renal tubular disease	
		familial cystic renal disease	
Erasmus MC	Center for Familian and Hereditary Tumors	Lynch Syndrome	
Erasmus MC	Center for pregnancy induced diseases	Pre-eclampsia (< 34 wks)	
		HELLP-syndrome	
Erasmus MC	Centre of Expertise for Uveitis	Uveitis	Voorlopig*
		Fuchs hetrochromic iridocyclitis	Voorlopig*
		Birdshot chorioretinopathy	Voorlopig*
		Paraneoplastic uveitis (ipv uveitis)	Voorlopig*
		Cancer-associated retinopathy	Voorlopig*
Erasmus MC	Center for Genetic Eye Diseases Rotterdam	Retinal dystrophy	
		Retinitis pigmentosa	
		Developmental defect of the eye	
		Color-vision disease	
		Achromatopsia	
Erasmus MC & LUMC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Myelodysplastic syndrome	
		Inherited acute myeloid leukemia	
		Constitutional neutropenia	
Erasmus MC	Rotterdam Ocular Melanoma Center (ROMC)	Uveal melanoma	
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	
		Primary bone dysplasia with increased bone density	
		Primary bone dysplasia with disorganized development of skeletal components	
		Chronic recurrent multifocal osteomyelitis	
LUMC	Nerve Centre	Rare neurologic disease, nerve lesion	
LUMC	Neuromuscular Center LUMC	Neuromuscular disease	
		Skeletal muscle disease	voor 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.	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
LUMC	Autonomic Disease Center	Primary orthostatic hypotension, PAF, MSA	voor PAF
LUMC	Gender Clinic Leiden, WAKZ-Curium LUMC	Rare endocrine disease, gender dysphoria	voor gender dysphoria
LUMC	Expertise center for monogenic diabetes mellitus	MODY syndrome	
LUMC	Expertise center for lipodystrophy	Berardinelli-Seip congenital lipodystrophy	voor primary lipodystrophy
		Familial partial lipodystrophy - Dunnigan type	
		Familial partial lipodystrophy associated with PPARG mutations	
		Familial partial lipodystrophy due to AKT2 mutations	
		Familial partial lipodystrophy - bberling type	
		Familial partial lipodystrophy associated with PLIN1 mutations	
		Acquired generalized lipodystrophy, also called Lawrence-Seip	
LUMC	Center for Endocrine Tumors Leiden (CETL)	Endocrine tumor with other location	voor Rare hypothalamic and pituitary disease (behalve MEN 1 en 2)
		Acromegaly	
		Cushing disease	
		Non-functioning pituitary adenoma	
		Pituitary deficiency	
		Hereditary pheochromocytoma-paraganglioma	
		Thyroid tumor	ORPHA:319494: zonder MEN type 2
		Adrenocortical carcinoma	
		Parathyroid carcinoma	
		Acquired chronic primary adrenal insufficiency	
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	Rare bone tumor	
		Multiple osteochondromas	
		Adamantinoma	
		Chondromyxoid fibroma	
		Osteosarcoma	
		Ewing sarcoma	
		Chondrosarcoma	
		Giant cell tumor of bone	
		Rare soft tissue tumor	
LUMC	Leiden Ocular Oncology Center	Uveal melanoma	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	Clinic for Lupus-, Vasculitis- and Complement-mediated systemic diseases	C3 glomerulonephritis	
		Pauci-immune glomerulonephritis	
		Anti-neutrophil cytoplasmic antibody-associated vasculitis	
		Immunoglobulin-mediated membranoproliferative 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 syndr. and FAMMMPC syndr.	
LUMC	Center of expertise Coffin-Siris syndrome	Coffin-Siris syndroom	
LUMC	Familial and hereditary breast cancer center	Hereditary breast and ovarian cancer syndrome	
		Hereditary breast cancer	
LUMC	Center for vestibular schwannoma and NF2	Vestibularis schwannoom	Voorlopig*

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
LUMC & AMC & VUmc	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	
		Centrale sereuze chorioretinopathie	
LUMC & Erasmus MC	Center for Congenital bone marrow failure and leukemia predisposition syndromes	Myelodysplastic syndrome	
		Inherited acute myeloid leukemia	
		Constitutional neutropenia	
LUMC	Center for Polyomavirus-associated skin infections and cancer (Acronym PASIC)	Virus-associated trichodysplasia spinulosa, usually known as trichodysplasia spinulosa	Voorlopig*
LUMC	Expert Center for hematopoietic stem cell transplantation and T-cell immunotherapy	Acute myeloid leukemia	
		Acute lymfoblastic leukemia	
		Plasma cell tumor	
LUMC	Center for primary and secondary immunodeficiencies	Mendelian susceptibility to mycobacterial diseases	
		Combined T and B cell immunodeficiency	
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+	Center for Genodermatoses	Inherited ichthyosis	
		Erythrokeratoderma	
		Lymphedema	
		Birt-Hogg-Du syndrome	
		Other sporadic genodermatoses	voor rare genetic skin disease
		Hereditary palmoplantar keratoderma	
		Basal cell nevus syndrome	
MUMC+	Expertise center Rare syndromes and cognitive disorders	Rare developmental defect during embryogenesis	
		Kabuki syndrome	
		Rett Syndrome	
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+ & Radboudumc	Marfan and related disorders policlinic	Marfan syndrome	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		Loeys-Dietz syndrome	
		Familial thoracic aortic aneurysm and aortic dissection	
MUMC+	Limburg renal registry	EGPA, GPA and MPA	
		Cryoglobulinemic vasculitis	
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	voor biliary tract carcinoom en hepatho-cellular 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	Endocrine tumor with other location; Papillary and follicular thyroid carcinoma, medullary thyroid carcinoma, anaplastic thyroid carcinoma	voor thyroid carcinoma
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	voor 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	
MUMC+	Division of Balance Disorders	Idiopathic Bilateral Vestibulopathy	
MUMC+	MUMC Mediastinal tumors	Thymic tumor	Voorlopig*
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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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; including NF2 patients	excl. NF2
		Tumor of endocrine glands; Hereditary pheochromocytoma-paraganglioma	
Radboudumc	Radboudumc pediatric center for congenital malformations of the intestinal tract	High anorectal malformation	
		Intermediate anorectal malformation	
		Low anorectal malformation	
		Anal fistula	
		VACTERL/VATER association	
		Caudal regression sequence	
		Hirschsprung disease	
		Cloacal exstrophy	
		Intestinal malformation	Voorlopig*
		Esophageal malformation	Voorlopig*
		Gastroschisis	Voorlopig*
		Omphalocele	Voorlopig*
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	Radboudumc Expertise 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)	
		Autoinflammatory syndrome	Voorlopig*
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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		Renal or urinary tract malformation	
		Cystinosis	
		Rare renal disease; Thrombotic microangiopathy (level ontbrak in form voor ref)	
Radboudumc	Centre for genetic neurodevelopmental disorders	Kleefstra syndrome due to a point mutation	
		Multiple congenital anomalies/dysmorphic syndrome-intellectual disability	voor Koolen -de Vries syndrome
		PTEN hamartoma tumor syndrome	voor Cowden syndrome
		KBG syndrome	
		Multiple congenital anomalies/dysmorphic syndrome - variable intellectual disability	voor Noonan syndrome
		Syndromic obesity	voor PWS i.p.v. Syndr. obesity
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	
		Congenital alveolo-capillary dysplasia	
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.	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	
		Tumor of the meninges	voor 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	
Radboudumc	Radboudumc center for congenital disorders of dental development	Oligodontia	
		Schöpf-Schulz-Passarge syndrome	Voorlopig*
Radboudumc	Radboudumc center for facial palsy	Paralytic facial malformation	Voorlopig*
Radboudumc	Radboud University Medical Centre CF centre	Cystic fibrosis	
Radboudumc	Radboudumc Center for male infertility	Male infertility due to obstructive azoospermia	Voorlopig*
		Rare idiopathic male infertility	Voorlopig*
		Male infertility due to gonadal dysgenesis or sperm disorder	Voorlopig*
Radboudumc	Radboud Center for Infectious diseases	Lyme disease	Voorlopig*
		Q fever	Voorlopig*
		Arbovirus fever	Voorlopig*
		Malaria	Voorlopig*
		Idiopathic recurrent and disabling cutaneous herpes	Voorlopig*
		Whooping cough (kinkhoest)	Voorlopig*
		Viral hemorrhagic fevers	Voorlopig*
		Fulminant virale hepatitis	Voorlopig*
Radboudumc	Radboudumc center for systemic autoimmune diseases	Systemic sclerosis	
		Localized scleroderma	
		Eosinophilic fasciitis	
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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		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; pediatric CNS tumours	voor 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	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	Granulomatosis with polyangiitis	voor vasculitis
		Microscopic polyangiitis	
		Eosinophilic granulomatosis with polyangiitis	
		Takayasu arteritis	
		Giant cell arteritis	
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	
		Cardiomyopathy	
		Genetic cardiac rhythm disease	
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 hypothalamic or pituitary disease	voor rare pituitary disease
UMCG	Pediatric Oncology group UMCG	Tumor of hematopoietic and lymphoid tissues	
UMCG	Center for rare inherited inborn errors of metabolism	Molybdenum cofactor deficientie type A	
UMCG	Paediatric Colorectal Expertise Center Groningen	Hirschsprung disease	Voorlopig*
		Anorectal malformation	Voorlopig*
UMCG	Centre of expertise for choledochal malformations	Choledochal cyst	
UMCG	Centre for Necrotizing Enterocolitis	Necrotiserende Enterocolitis	
UMCU	Center Inherited Metabolic Diseases	Disorder of fatty acid oxidation and ketone body metabolism	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		Disorder of pyridoxine metabolism	
UMCU	Brain Centre Rudolf Magnus, Neuromuscular Diseases	Neuromuscular disease; ALS, PLS, Progressive Spinal Muscular Atrophy and polyneuropathy	voor Neuro-muscular disease
			voor Spinal muscular atrophy
UMCU	Center for Refractory Pediatric Epilepsy	Rare epilepsy	
		Tuberous sclerosis	
		Continuous spikes and waves during sleep and ESES	
		Genetic causes of refractory pediatric epilepsy	voor 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	
		Cleft/lip palate	
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	Myeloproliferative neoplasm ; Polycytemia vera (PV) and Essential thrombocythemia (ET)	voor Multiple myeloma
		Myeloid hemopathy; AML/RAEB-T, RAEB, Myelodysplasia and remaining diseases	
		Myeloproliferative neoplasm; CML	
		Myeloid hemopathy, ao CMMol / Myelofibrosis/ eosinophilic disorders	voor Non Hodgkin lymphoma
		Plasma cell tumor; MM/ Amyloidosis and Monoclonal gammopathy(MGUS)	
		Lymphoma; NHL, Hodgkin, CLL, Waldenstrom, hairycell and other lymfoproliferative diseases	
		Primary central nervous system lymphoma	voor acute lymphoblastic and myeloid leukemia
		Acute lymphoblastic leukemia	
		Systemic mastocytosis	
		Myeloid hemopathy; HES and Langerhanscel histiocytose	
UMCU	Centre for rare tumors	Inherited cancer-predisposing syndrome	voor Men 1
		Multiple endocrine neoplasia type 2A; incl. fam medullary thyroid carcinoma, MEN2B and sporadic medulary thyroid carcinoma	
		Von Hippel-Lindau disease	
		Thyroid tumor	
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.	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
		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	
		Vasculitis	
		Systemic sclerosis	
		Rare coagulation disorder; Antiphospholipid syndrome	
		Systemic autoimmune disease; extraglandular manifestations in Sjogren	
UMCU	WKZ center for congenital malformations	Esophageal atresia	
		Anorectal malformation	
		Hirschsprung Disease	
UMCU	Multidisciplinary Center for Limb Reduction Defects	Rare bone disease; m.n. primaire skeletdysplasieen en dysostoses, ook osteogenesis imperfecta en achondroplasie	
		Non-syndromic limb reduction defects	
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	PHACE syndrome	voor vascular anomaly
		Kaposiform hemangioendothelioma	
		Diffuse neonatal hemangiomatosis	
		Rapidly involuting congenital hemangioma; RICH and NICH	
		PELVIS syndrome	
		Laryngotracheal angioma	
		Mucocutaneous venous malformations	
		Macrocystic lymphatic malformation	
		Vascular malformation	
		Microcystic lymphatic malformation	
UMCU	Sylvia Toth Center for Multi-disciplinary follow up of Lysosomal Storage Disorders, University Medical Center Utrecht	Hurler disease	voor Hurler disease
UMCU	Mobility Clinic	Osteochondritits Dissecans	
UMCU	Center for inherited cardiovascular disease	Cardiomyopathy	Voorlopig*
		Genetic cardiac rhytm disease	Voorlopig*
		Rare genetic vascular disease	Voorlopig*
UMCU	Centre of expertise for extracranial carotid artery aneurysms (ECAA)	Extracraniele carotis aneurysma	Voorlopig*

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
UMCU	Dutch National Expertise Center for Pseudoxanthoma elasticum (DNECP)	Pseudoxanthoma elasticum-	
UMCU	Utrecht Center for Keratoconus and Corneal Dystrophies	Keratoconus	Voorlopig*
UMCU	Center for Neonatal Neurology	Hypoxic ischemic brain injury	
		Pediatric arterial ischemic stroke	
		Periventricular leukomalacia	
UMCU	Center for Intracranial Vasculopathies	Sneddon syndrome	Voorlopig*
		Primary central nervous system vasculitis	Voorlopig*
		Moya Moya angiopathy:	Voorlopig*
UMCU	Head and Neck Working Group Utrecht	Squamous cell carcinoma of head and neck	
		Squamous cell carcinoma of the oral tongue	
		Rare tumors of salivary glands	
		Nasopharyngeal carcinoma	
UMCU	UMCU Neuro-Oncology Center	Glial tumor	
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	Rare odontologic disease, Robin sequence	voor Robin seq.
VUmc	Expertise Center Head and Neck tumors	Nasopharyngeal carcinoma	
		Squamous cell carcinoma of the oral cavity and lip	
		Rare head and neck tumor	
		Hereditary pheochromocytoma-paraganglioma	
VUmc	Celiac disease center	Refractory celiac disease	
		Autoimmune hepatitis	
		Short Bowel Syndrome	
		Enteropathy Associated T-cell Lymphoma	
VUmc	Center for rare haematologic cancers	Acute myeloid leukemia	voor Multiple myeloma
		Multiple myeloma; incl Amyloidosis, Castlemans disease and POEMS	
		Myelofibrosis with myeloid metaplasia	
		Chronic myeloid leukemia	voor Non Hodgkin lymphoma
		Myelodysplastic syndromes	
		Acute lymphoblastic leukemia	
		Hodgkin lymphoma	voor acute lymphoblastic and myeloid leukemia
		Enteropathy-associated T-cell lymphoma	
		Aggressive B-cell non-Hodgkin lymphoma	
		Langerhans cell histiocytosis	
		Myelodysplastic syndromes (al erkend onder cluster NHL)	
		Chronic Myeloid Leukemia (al erkend onder cluster NHL)	
		Primary Myelofibrosis	
		Hodgkin Lymphoma (al erkend onder cluster AL&ML)	
		Myeloid hemopathy	
		Lymphoid hemopathy	
VUmc	Dutch Center for Oral Medicine and Oral Pathology	Squamous cell carcinoma of head and neck; Ameloblastomas and Keratocystic odontogenic tumors, Gorlin-Goltz syndrome	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
VUmc	Brain Tumor Center Amsterdam	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	
		X linked osteoporosis with fractures	Voorlopig*
		Fibrous dysplasia of bone (Head)	Voorlopig*
VUmc	VUmc multidisciplinary (genetic) breast cancer team	Rare genetic tumor; Hereditary breast cancer, CHEK2*1100delC related	voor 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	neuromusculaire aandoeningen, in het bijzonder Foetale Akinesie	voor FADS
VUmc & AMC	Amsterdam Centre of congenital malformations	Hirschsprung Disease	
		Esophageal atresia	
		Anorectal malformation	
		Sacrococcygeal teratoma	
		Chronic intestinal pseudoobstruction	
VUmc & AMC	Amsterdam Center for ILD and sarcoïdosis	Sarcoidosis	Voorlopig*
VUmc	Center on Atypical Sex or Gender development	Gender dysphoria	
		Disorder of sex development	
VUmc & LUMC & AMC	Center for Hereditary Retinal Diseases Leiden Amsterdam	Retinal dystrophy	
		Centrale sereuze chorioretinopathie	
NKI-AvL	Sarcoma Expertise Centre Amsterdam	Soft tissue sarcoma	
		Rare soft tissue tumor; Gastrointestinal stromal tumor (GIST)	

Instelling	Naam Expertise Centrum	Cluster van / Specifieke aandoening	Toelichting erkenning
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	Rare respiratory tumor; Rare pleural malignancies	voor mesothelioom
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	voor testicular cancer
		Squamous cell carcinoma of penis	voor 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	
NKI-AvL	Rare Skin Cancer Center	Merkel Cell carcinoma	
NKI-AvL	Centre for rare nervous system tumor	Glial tumor	
NKI-AvL	Center of rare GI tumours	Epithelial tumor of anal cancer	
		Rare gastroesophageal tumor	Voorlopig*
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	Sleep disorders	voor rare sleep disorder
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	voor alle genoemde aandoeningen
		Uveitis	
		Rare genetic eye disease; Retinal Dystrophies	
STZ-Albert Schweitzer hospital, Dordrecht	Centre of expertise Retroperitoneal Fibrosis	Retroperitoneal fibrosis	
SZT-Medisch Spectrum Twente	Gastrointestinal ischemia Centre	Celiac trunk compression syndrome	
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; incl. idiopathic interstitial pneumonia	voor IPF en IIP
		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	

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STZ-St. Antonius Ziekenhuis	Center for Pulmonary vascular diseases	Hereditary hemorrhagic telangiectasia	
		Chronic thromboembolic pulmonary hypertension	
		Idiopathic pulmonary arterial hypertension	
STZ - St. Antonius Ziekenhuis, Nieuwegein	St. Antonius Oesofagus Centrum	Esophageal carcinoma	
STZ-CWZ Nijmegen	Malignant Hyperthermia investigation unit Nijmegen	Malignant hyperthermia	
STZ-CWZ Nijmegen	Center for Cerebrotendinous xanthomatosis	Cerebrotendinous xanthomatosis	
STZ-Maxima Medisch Centrum	SolviMáx, Center of Excellence for Abdominal Wall and Groin Pain	Acquired peripheral neuropathy; Anterior cutaneous nerve syndrome (ACNES)	alleen voor ACNES
STZ-Maxima Medisch Centrum	Center for Adrenal Tumors	Adrenocortical carcinoma; also 'Catecholamine-producing tumor'and the 'Adrenal incidentaloma'	alleen voor 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	Transient hypogammaglobulinemia of infancy	Voorlopig t/m sep 2017, voor Immuno-deficiency predominantly affecting antibody production, mainly the various types of unclassified antibody deficiency
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	Rare nervous system tumor	voor gliomen
PMC	Princess Máxima Center for pediatric solid tumors	(Pediatric) germ cell tumor	Voorlopig*
		(Pediatric) rare soft tissue tumor	Voorlopig*
		(Pediatric) rare renal tumor	Voorlopig*
		Pediatric) rare digestive tumor	Voorlopig*
		Neuroblastoma	Voorlopig*
PMC	Princess Máxima Center for pediatric hematological malignancies and stem cell transplants	(Pediatric) tumor of hematopoietic and lymphoid tissues	Voorlopig*
		(Pediatric) myeloid hemopathy	Voorlopig*
		(Pediatric) lymphoid hemopathy	Voorlopig*
		(Pediatric) acute myeloid leukemia	Voorlopig*
		(Pediatric) acute lymphoblastic leukemia	Voorlopig*
		(Pediatric) lymphoma	Voorlopig*
		(Pediatric) myelodysplastic syndrome	Voorlopig*

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PMC	Princess Máxima Center for pediatric brain tumors	(Pediatric) rare nervous system tumor	Voorlopig*
		(Pediatric) glial tumor	Voorlopig*
		(Pediatric) astrocytoma	Voorlopig*
		(Pediatric) medulloblastoma	Voorlopig*
			*voor deze aandoeningen geldt een voorlopige erkenning voor 1 jaar