AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
706031	H-11-15	Rare neoplastic diseases	Vulvar squamous cell carcinoma	494448	P	18	NFK: Patiëntenplatform Zeldzame Kankers
706031	H-11-15	Rare neoplastic diseases	Malignant epithelial tumor of ovary	398934	P	88	Stichting Olijf
706031	H-11-15	Rare neoplastic diseases	Rare cancer of cervix uteri	213761	P	88	Stichting Olijf
706031	H-11-15	Rare neoplastic diseases	Rare cancer of corpus uteri	213569	P	88	Stichting Olijf
706031	H-11-15	Rare neoplastic diseases	Rare gynecological tumor	98063	P	88	Stichting Olijf
706075	H-11-43	Rare neoplastic diseases, Rare otorhinolaryngological diseases	Nasopharyngeal carcinoma	150	P	169	Patiëntenvereniging Hoofd - Hals
706138	H-5-4	Rare renal diseases	Genetic renal tubular disease	183592	P	31	Nierpatiëntenvereniging Nederland (NVN)
706138	H-5-4	Rare renal diseases	Nephropathy secondary to a storage or other metabolic disease	93593	P	31	Nierpatiëntenvereniging Nederland (NVN)
706138	H-5-4	Rare renal diseases	Rare renal tubular disease	93603	P	31	Nierpatiëntenvereniging Nederland (NVN)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
706138	H-5-4	Rare renal diseases	Nephropathy secondary to a storage or other metabolic disease	93593	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
706138	H-5-4	Rare renal diseases	Rare renal tubular disease	93603	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
706167	H-14-8	Rare neurological diseases	Rare central nervous system and retinal vascular disease	71281	P	100	Oogvereniging

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
706167	H-14-8	Rare neurological diseases	Acquired aneurysmal subarachnoid hemorrhage	90065	P	315	Stichting Hersenaneurysma Patiënten Platform
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft hard palate	101023	P	17	LaPosa
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip and alveolus	141291	P	17	LaPosa
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip with or without cleft palate	1991	P	17	LaPosa
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip/palate	199306	P	17	LaPosa
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft palate	2014	P	17	LaPosa
706222	H-2-16	Rare surgical maxillo-facial diseases	Submucosal cleft palate	155878	P	17	LaPosa

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft hard palate	101023	P	287	Schisis Nederland
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip and alveolus	141291	P	287	Schisis Nederland
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip with or without cleft palate	1991	P	287	Schisis Nederland
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft lip/palate	199306	P	287	Schisis Nederland
706222	H-2-16	Rare surgical maxillo-facial diseases	Cleft palate	2014	P	287	Schisis Nederland
706222	H-2-16	Rare surgical maxillo-facial diseases	Submucosal cleft palate	155878	P	287	Schisis Nederland
708008	H-12-8	Rare hematologic diseases	Rare hemorrhagic disorder due to a qualitative platelet defect	275736	P	74	Nederlandse Vereniging voor Hemofiliepatiënten (NVHP)
708008	H-12-8	Rare hematologic diseases	Rare hemorrhagic disorder due to a qualitative platelet defect	275736	P	75	Stichting Zeldzame Bloedziekten (SZB)
708027	H-4-1	Rare neurological diseases	Cerebral malformation with epilepsy	166478	P	7	Epilepsie NL

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
708027	H-4-1	Rare neurological diseases	Epilepsy Syndrome	166463	P	7	Epilepsie NL
708027	H-4-1	Rare neurological diseases	Infantile Epilepsy Syndrome	98258	P	7	Epilepsie NL
708027	H-4-1	Rare neurological diseases	Monogenic disease with epilepsy	166472	P	7	Epilepsie NL
708027	H-4-1	Rare neurological diseases	Rare epilepsy	101998	P	7	Epilepsie NL
708027	H-4-1	Rare neurological diseases	Cerebral malformation with epilepsy	166478	P	333	Stichting Dravet Syndroom Nederland / Vlaanderen
708027	H-4-1	Rare neurological diseases	Epilepsy Syndrome	166463	P	333	Stichting Dravet Syndroom Nederland / Vlaanderen
708027	H-4-1	Rare neurological diseases	Infantile Epilepsy Syndrome	98258	P	333	Stichting Dravet Syndroom Nederland / Vlaanderen
708027	H-4-1	Rare neurological diseases	Monogenic disease with epilepsy	166472	P	333	Stichting Dravet Syndroom Nederland / Vlaanderen
708027	H-4-1	Rare neurological diseases	Rare epilepsy	101998	P	333	Stichting Dravet Syndroom Nederland / Vlaanderen

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	HELLP syndrome	244242	P	113	HELLP Stichting
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	Preeclampsia	275555	P	113	HELLP Stichting
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	HELLP syndrome	244242	P	114	Care4Neo
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	Placental insufficiency	439167	P	114	Care4Neo
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	Preeclampsia	275555	P	114	Care4Neo
708047	H-26-5	Rare circulatory system diseases, Rare gynecologic and obstetric diseases	Rare disorders related with pregnancy, childbirth and puerperium	163637	P	114	Care4Neo
708068	H-8-15	Rare respiratory diseases	Pulmonary arterial hypertension	182090	P	158	Stichting Pulmonale Hypertensie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
708068	H-8-15	Rare respiratory diseases	Rare pulmonary hypertension	71198	P	158	Stichting Pulmonale Hypertensie
708087	H-16-8	Rare cardiac diseases	Arrhythmogenic right ventricular dysplasia	247	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Brugada syndrome	130	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Dilated cardiomyopathy	217604	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Familial long QT syndrome	768	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Idiopathic ventricular fibrillation - not Brugada type	228140	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Rare familial disorder with hypertrophic cardiomyopathy	99739	P	58	Harteraad
708087	H-16-8	Rare cardiac diseases	Brugada syndrome	130	P	337	Brugada Community
708145	H-15-7	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	88	Stichting Olijf

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
708145	H-15-7	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	89	Borstkankervereniging (BVN)
708145	H-15-7	Rare genetic diseases	Intestinal polyposis syndrome	104010	P	93	Stichting Lynch Polyposis
708145	H-15-7	Rare genetic diseases	Lynch syndrome	144	P	93	Stichting Lynch Polyposis
708145	H-15-7	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	339	Stichting Erfelijke Kanker Nederland (SEKN)
708169	H-2-10	Rare otorhinolaryngological diseases	Idiopathic Bilateral Vestibulopathy	171684	P	33	Stichting Hoormij
708169	H-2-10	Rare otorhinolaryngological diseases	Idiopathic Bilateral Vestibulopathy	171684	P	282	De 9e van .....
708189	H-11-25	Rare neoplastic diseases	Desmoid tumor	873	P	94	Stichting Patiëntenplatform Sarcomen
708189	H-11-25	Rare neoplastic diseases	Gastrointestinal stromal tumor	44890	P	94	Stichting Patiëntenplatform Sarcomen

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
708208	H-12-7	Rare hematologic diseases, Rare neoplastic diseases	Acute graft versus host disease	99920	P	81	Stichting Hematon
708208	H-12-7	Rare hematologic diseases, Rare neoplastic diseases	B-cell chronic lymphocytic leukemia	67038	P	81	Stichting Hematon
708208	H-12-7	Rare hematologic diseases, Rare neoplastic diseases	Chronic graft versus host disease	99921	P	81	Stichting Hematon
708208	H-12-7	Rare hematologic diseases, Rare neoplastic diseases	Graft versus host disease	39812	P	81	Stichting Hematon
708228	H-14-4	Rare inborn errors of metabolism	Mitochondrial diseases	68380	P	43	Spierziekten Nederland (SN)
708228	H-14-4	Rare inborn errors of metabolism	Mitochondrial diseases	68380	P	199	Stichting Nemo
708228	H-14-4	Rare inborn errors of metabolism	Mitochondrial diseases	68380	P	274	Belangenvereniging LOA/LHON
708228	H-14-4	Rare inborn errors of metabolism	Mitochondrial diseases	68380	P	281	Stichting Cure ADOA Foundation
709008	H-10-11	Rare skin diseases	Merkel Cell carcinoma	79140	P	343	Stichting NETNECkanker
709054	H-11-54	Rare gastroenterologic diseases	Epithelial tumor of anal cancer	424010	P	18	NFK: Patiëntenplatform Zeldzame Kankers

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
709054	H-11-54	Rare gastroenterologic diseases	Rare gastroesophageal tumor	180821	P	63	SPKS - Leven met Maag- of Slokdarmkanker
709091	H-9-19	Rare neurological diseases	Glial tumor	182067	P	82	Vereniging Kinderkanker Nederland
709091	H-9-19	Rare neurological diseases	Glial tumor	182067	P	91	Hersenletsel.nl
709238	H-11-53	Rare neoplastic diseases	Pleural mesothelioma	50251	P	109	Asbestslachtoffers Vereniging Nederland (AVN)
709238	H-11-53	Rare neoplastic diseases	Thymic tumor	100100	P	204	Longkanker Nederland
709290	H-3-19	Rare endocrine diseases	Thyroid carcinoma	100088	P	60	Schildklier Organisatie Nederland (SON)
709290	H-3-19	Rare endocrine diseases	Thyroid tumor	100087	P	60	Schildklier Organisatie Nederland (SON)
709336	H-11-35	Rare neoplastic diseases	Desmoid tumor	873	P	94	Stichting Patiëntenplatform Sarcomen
709386	H-7-19	Rare abdominal surgical diseases	Celiac artery compression syndrome	293208	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
709465	H-9-18	Rare neurological diseases	Genetic cerebral small vessel disease	477754	P	14	HCHWA-D Vereniging
709465	H-9-18	Rare neurological diseases	Rare central nervous system and retinal vascular disease	71281	P	100	Oogvereniging
709465	H-9-18	Rare neurological diseases	Acquired aneurysmal subarachnoid hemorrhage	90065	P	315	Stichting Hersenaneurysma Patiënten Platform
709465	H-9-18	Rare neurological diseases	Cerebral sinovenous thrombosis	329217	P		Zonder patiëntenorganisatie
709517	H-9-17	Rare neurological diseases	Pyridoxine Dependent Epilepsy	3006	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
709738	H-7-18	Rare abdominal surgical diseases, Rare neurological diseases	Early-onset autosomal dominant Alzheimers disease	1020	P	2	Alzheimer Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
709738	H-7-18	Rare abdominal surgical diseases, Rare neurological diseases	Human Prion Disease	56970	P	2	Alzheimer Nederland
709738	H-7-18	Rare abdominal surgical diseases, Rare neurological diseases	Frontotemporal degeneration with dementia	98535	P	9	FTD Lotgenoten
709775	H-5-2	Rare inborn errors of metabolism, Rare renal diseases	Glomerular disease	93548	P	31	Nierpatiëntenvereniging Nederland (NVN)
709775	H-5-2	Rare inborn errors of metabolism, Rare renal diseases	Rare renal tubular disease	93603	P	31	Nierpatiëntenvereniging Nederland (NVN)
709775	H-5-2	Rare inborn errors of metabolism, Rare renal diseases	Thrombotic microangiopathy	93573	P	31	Nierpatiëntenvereniging Nederland (NVN)
709775	H-5-2	Rare inborn errors of metabolism, Rare renal diseases	Rare renal tubular disease	93603	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
709775	H-5-2	Rare inborn errors of metabolism, Rare renal diseases	Thrombotic microangiopathy	93573	P	75	Stichting Zeldzame Bloedziekten (SZB)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
709794	H-22-15	Rare immunological diseases	Primary Sjögren syndrome	289390	P	22	Nationale Vereniging Sjögrenpatiënten (NVSP)
709794	H-22-15	Rare immunological diseases	Vasculitis	52759	P	51	Vasculitis Stichting
709794	H-22-15	Rare immunological diseases	Systemic lupus erythematosus	536	P	128	Nationale vereniging voor lupus, aps, sclerodermie en MCTD (NVLE)
709794	H-22-15	Rare immunological diseases	Systemic sclerosis	801	P	128	Nationale vereniging voor lupus, aps, sclerodermie en MCTD (NVLE)
709794	H-22-15	Rare immunological diseases	Systemic lupus erythematosus	536	P	161	Huid Nederland
709794	H-22-15	Rare immunological diseases	Systemic sclerosis	801	P	161	Huid Nederland
709839	H-11-19	Rare neoplastic diseases	Adenocarcinoma of the small intestine	104075	P	304	Stichting Darmkanker Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
709885	H-1-9	Rare bone diseases	Osteongenesis imperfecta	666	P	53	Vereniging Osteogenesis Imperfecta (VOI)
709929	H-22-14	Rare immunological diseases	Exercise-induced malignant hyperthermia	466650	P		Zonder patiëntenorganisatie
709929	H-22-14	Rare immunological diseases	Malignant hyperthermia of anesthesia	423	P		Zonder patiëntenorganisatie
709929	H-22-14	Rare immunological diseases	Rare disease with malignant hyperthermia	466658	P		Zonder patiëntenorganisatie
710024	H-6-6	Rare genetic diseases	Cerebral visual impairement	447788	P	100	Oogvereniging
710024	H-6-6	Rare genetic diseases	Inherited retinal disorder	71862	P	100	Oogvereniging
710077	H-17-16	Rare surgical thoracic diseases	arterial thoracic outlet syndrome	357107	P		Zonder patiëntenorganisatie
710077	H-17-16	Rare surgical thoracic diseases	neurogenic thoracic outlet syndrome	100073	P		Zonder patiëntenorganisatie
710077	H-17-16	Rare surgical thoracic diseases	Thoracic Outlet Syndrome	97330	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710077	H-17-16	Rare surgical thoracic diseases	venous thoracic outlet syndrome	357131	P		Zonder patiëntenorganisatie
710108	H-10-10	Rare skin diseases	Necrotizing soft tissue infection	440368	P	161	Huid Nederland
710108	H-10-10	Rare skin diseases	Necrotizing soft tissue infection	440368	P	231	Vereniging van Mensen met Brandwonden
710236	H-6-4	Rare ophthalmic disorders	Leber plus disease	99718	P	40	Sarcoïdose Belangenvereniging Nederland
710236	H-6-4	Rare ophthalmic disorders	Uveitis	98715	P	40	Sarcoïdose Belangenvereniging Nederland
710236	H-6-4	Rare ophthalmic disorders	Leber plus disease	99718	P	100	Oogvereniging
710236	H-6-4	Rare ophthalmic disorders	Uveitis	98715	P	100	Oogvereniging
710236	H-6-4	Rare ophthalmic disorders	Leber plus disease	99718	P	125	Macula Vereniging
710236	H-6-4	Rare ophthalmic disorders	Uveitis	98715	P	125	Macula Vereniging
710236	H-6-4	Rare ophthalmic disorders	Leber plus disease	99718	P	240	Stichting Usher Syndroom

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710236	H-6-4	Rare ophthalmic disorders	Uveitis	98715	P	240	Stichting Usher Syndroom
710236	H-6-4	Rare ophthalmic disorders	Leber plus disease	99718	P	274	Belangenvereniging LOA/LHON
710286	H-6-5	Rare ophthalmic disorders	IgG4-related ophthalmic disease	449563	P	51	Vasculitis Stichting
710286	H-6-5	Rare ophthalmic disorders	Necrobiotic xanthogranuloma	158011	P	60	Schildklier Organisatie Nederland (SON)
710286	H-6-5	Rare ophthalmic disorders	IgG4-related ophthalmic disease	449563	P	60	Schildklier Organisatie Nederland (SON)
710372	H-9-14	Rare neurological diseases	Benign peripheral nerve sheath tumor	252131	P	30	Neurofibromatosevereniging Nederland (NFVN)
710372	H-9-14	Rare neurological diseases	Tumor of meninges	252025	P	33	Stichting Hoormij
710372	H-9-14	Rare neurological diseases	Benign peripheral nerve sheath tumor	252131	P	33	Stichting Hoormij
710372	H-9-14	Rare neurological diseases	Glial tumor	182067	P	33	Stichting Hoormij
710372	H-9-14	Rare neurological diseases	Glial tumor	182067	P	82	Vereniging Kinderkanker Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710372	H-9-14	Rare neurological diseases	Glial tumor	182067	P	91	Hersenletsel.nl
710455	H-17-11	Rare genetic diseases	Down syndrome	870	P	48	Stichting Rubinstein-Taybi
710455	H-17-11	Rare genetic diseases	16p11.2p12.2 microdeletion syndrome	261211	P	48	Stichting Rubinstein-Taybi
710455	H-17-11	Rare genetic diseases	Distal 16p11.2 microdeletion syndrome	261222	P	48	Stichting Rubinstein-Taybi
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microdeletion syndrome	261197	P	48	Stichting Rubinstein-Taybi
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microduplication syndrome	370079	P	48	Stichting Rubinstein-Taybi
710455	H-17-11	Rare genetic diseases	Down syndrome	870	P	71	Stichting Down Syndroom (SDS)
710455	H-17-11	Rare genetic diseases	Down syndrome	870	P	145	Stichting Pitt Hopkins
710455	H-17-11	Rare genetic diseases	16p11.2p12.2 microdeletion syndrome	261211	P	145	Stichting Pitt Hopkins

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710455	H-17-11	Rare genetic diseases	Distal 16p11.2 microdeletion syndrome	261222	P	145	Stichting Pitt Hopkins
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microdeletion syndrome	261197	P	145	Stichting Pitt Hopkins
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microduplication syndrome	370079	P	145	Stichting Pitt Hopkins
710455	H-17-11	Rare genetic diseases	Down syndrome	870	P	146	Marshall Smith Syndroom Research Foundation
710455	H-17-11	Rare genetic diseases	16p11.2p12.2 microdeletion syndrome	261211	P	146	Marshall Smith Syndroom Research Foundation
710455	H-17-11	Rare genetic diseases	Distal 16p11.2 microdeletion syndrome	261222	P	146	Marshall Smith Syndroom Research Foundation
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microdeletion syndrome	261197	P	146	Marshall Smith Syndroom Research Foundation

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microduplication syndrome	370079	P	146	Marshall Smith Syndroom Research Foundation
710455	H-17-11	Rare genetic diseases	16p11.2p12.2 microdeletion syndrome	261211	P	290	Zeldsamen
710455	H-17-11	Rare genetic diseases	Distal 16p11.2 microdeletion syndrome	261222	P	290	Zeldsamen
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microdeletion syndrome	261197	P	290	Zeldsamen
710455	H-17-11	Rare genetic diseases	Proximal 16p11.2 microduplication syndrome	370079	P	290	Zeldsamen
710477	H-12-13	Rare hematologic diseases	Chronic Myeloid Leukemia	521	P	81	Stichting Hematon
710505	H-7-3	Rare gastroenterologic diseases	Omphalocele	660	P	114	Care4Neo
710505	H-7-3	Rare gastroenterologic diseases	Congenital Chronic Diarrhea with protein-losing enteropathy	329242	P	120	Vereniging Anusatresie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710505	H-7-3	Rare gastroenterologic diseases	Congenital tufting enteropathy	92050	P	120	Vereniging Anusatresie
710505	H-7-3	Rare gastroenterologic diseases	Duodenum atresia	1203	P	120	Vereniging Anusatresie
710505	H-7-3	Rare gastroenterologic diseases	Microvillus inclusion disease	2290	P	120	Vereniging Anusatresie
710505	H-7-3	Rare gastroenterologic diseases	Rare diseases involving intestinal motility	104009	P	120	Vereniging Anusatresie
710505	H-7-3	Rare gastroenterologic diseases	Omphalocele	660	P	120	Vereniging Anusatresie
710505	H-7-3	Rare gastroenterologic diseases	Rare diseases involving intestinal motility	104009	P	123	Vereniging Ziekte van Hirschsprung
710545	H-7-2	Rare gastroenterologic diseases	Eosinophilic esophagitis	73247	P	124	Vereniging Ouderen en Kinderen met Slokdarmafsluiting (VOKS)
710545	H-7-2	Rare gastroenterologic diseases	Idiopathic achalasia	930	P	124	Vereniging Ouderen en Kinderen met Slokdarmafsluiting (VOKS)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
710545	H-7-2	Rare gastroenterologic diseases	Eosinophilic esophagitis	73247	P	182	Vereniging voor Allergiepatiënten
711854	H-8-26	Rare respiratory diseases	Bronchopulmonary Dysplasia	70589	P	114	Care4Neo
712016	H-8-25	Rare respiratory diseases	Cystic Fibrosis	586	P	159	Nederlandse Cystic Fibrosis Stichting (NCFS)
712061	H-7-16	Rare abdominal surgical diseases, Rare gastroenterologic diseases	Gastroschisis	2368	P	114	Care4Neo
715007	H-26-7	Rare infectious diseases	Leishmaniasis	507	P	161	Huid Nederland
715007	H-26-7	Rare infectious diseases	Arbovirus fever	344	P		Zonder patiëntenorganisatie
715007	H-26-7	Rare infectious diseases	Malaria	673	P		Zonder patiëntenorganisatie
715007	H-26-7	Rare infectious diseases	Rare parasitic disease	163588	P		Zonder patiëntenorganisatie
715007	H-26-7	Rare infectious diseases	Schistosomiasis	1247	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715026	H-11-19	Rare neoplastic diseases	Desmoid tumor	873	P	94	Stichting Patiëntenplatform Sarcomen
715026	H-11-19	Rare neoplastic diseases	Gastrointestinal stromal tumor	44890	P	94	Stichting Patiëntenplatform Sarcomen
715026	H-11-19	Rare neoplastic diseases	Soft tissue sarcoma	3394	P	94	Stichting Patiëntenplatform Sarcomen
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Dermatofibrosarcoma protuberans	31112	P	94	Stichting Patiëntenplatform Sarcomen
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Pyogenic arthritis-pyoderma gangrenosum-acne syndrome	69126	P	135	Stichting KAISZ
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Large congenital melanocytic nevus	626	P	141	Nevus Netwerk
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Neurocutaneous melanocytosis	2481	P	141	Nevus Netwerk
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Neurocutaneous melanocytosis	2481	P	161	Huid Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Pyoderma gangrenosum - acne - hidradenitis suppurativa	289478	P	161	Huid Nederland
715061	H-10-4	Rare neoplastic diseases, Rare skin diseases	Cutaneous neuroendocrine carcinoma	79140	P	343	Stichting NETNECkanker
715080	H-21-6	Rare systemic and rheumatological diseases	Mixed connective tissue disease	809	P	22	Nationale Vereniging Sjögren Patiënten
715080	H-21-6	Rare systemic and rheumatological diseases	Mixed connective tissue disease	809	P	128	Nationale vereniging voor lupus, aps, sclerodermie en MTCD (NVLE)
715100	H-26-6	Rare gynecologic and obstetric diseases	Retinopathy of prematurity	90050	P	100	Oogvereniging
715100	H-26-6	Rare gynecologic and obstetric diseases	Retinopathy of prematurity	90050	P	114	Care4Neo
715100	H-26-6	Rare gynecologic and obstetric diseases	Selective Intrauterine Growth Restriction	617301	P	317	Stichting Taps Support

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715100	H-26-6	Rare gynecologic and obstetric diseases	Twin Anemia Polycythemia Sequence	617294	P	317	Stichting Taps Support
715100	H-26-6	Rare gynecologic and obstetric diseases	Twin Reversed Arterial Perfusion sequence	617297	P	317	Stichting Taps Support
715100	H-26-6	Rare gynecologic and obstetric diseases	Retinopathy of prematurity	90050	P	322	CHI Support
715100	H-26-6	Rare gynecologic and obstetric diseases	Selective Intrauterine Growth Restriction	617301	P	322	CHI Support
715100	H-26-6	Rare gynecologic and obstetric diseases	Twin Anemia Polycythemia Sequence	617294	P	322	CHI Support
715100	H-26-6	Rare gynecologic and obstetric diseases	Twin Reversed Arterial Perfusion sequence	617297	P	322	CHI Support
715121	H-0-4	Rare neurological diseases	Idiopathic hypersomnia	33208	P	210	Narcolepsie Vereniging Nederland
715121	H-0-4	Rare neurological diseases	Narcolepsy type 1	2073	P	210	Narcolepsie Vereniging Nederland
715121	H-0-4	Rare neurological diseases	Narcolepsy type 2	83465	P	210	Narcolepsie Vereniging Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715121	H-0-4	Rare neurological diseases	Rare sleep disorder	68354	P	210	Narcolepsie Vereniging Nederland
715121	H-0-4	Rare neurological diseases	Kleine-Levin syndrome	33543	P		Zonder patiëntenorganisatie
715145	H-9-11	Rare neurological diseases	Pure autonomic failure	441	P	242	Stichting Orthostatische Tremor
715145	H-9-11	Rare neurological diseases	Rare autonomic nervous system disorder	423662	P	329	Lotgenoten Lage Bloeddruk Hartpatiënten Nederland
715145	H-9-11	Rare neurological diseases	Pure autonomic failure	441	P	329	Lotgenoten Lage Bloeddruk Hartpatiënten Nederland
715165	H-25-11	Rare infectious diseases	Arboviral fever	344	P	118	Nederlandse Meningitis Stichting
715165	H-25-11	Rare infectious diseases	Malaria	673	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Middle East respiratory syndrome	576074	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715165	H-25-11	Rare infectious diseases	Rabies	770	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Rare parasitic disease	163588	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Schistosomiasis	1247	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Severe acute respiratory syndrome	140896	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Strongyloidiasis	76	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Viral hemorrhagic fever	341	P		Zonder patiëntenorganisatie
715165	H-25-11	Rare infectious diseases	Yellow fever	99829	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715185	H-17-14	Rare developmental anomalies during embryogenesis	Coffin-Siris syndrome	1465	P	290	Zeldsamen
715185	H-17-14	Rare developmental anomalies during embryogenesis	Intellectual disability-severe speech delay-mild dysmorphism syndrome	391372	P	290	Zeldsamen
715185	H-17-14	Rare developmental anomalies during embryogenesis	Rare intellectual disability	87277	P	290	Zeldsamen
715185	H-17-14	Rare developmental anomalies during embryogenesis	Intellectual disability-severe speech delay-mild dysmorphism syndrome	391372	P	342	Stichting Kind Beter
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Non-recovering obstetric brachial plexus lesion	439202	P	8	Erbse Parese Vereniging Nederland
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Acquired periheral neuropathy	182086	P	43	Spierziekten Nederland (SN)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Acquired periheral neuropathy	182086	P	288	SOS Nederland
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Anterior cutaneous nerve entrapment syndrome	51890	P	295	Stichting ACNES
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Anterior cutaneous nerve entrapment syndrome	51890	P	296	ACNES Foundation
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Neurogenic thoracic outlet syndrome	100073	P		Zonder patiëntenorganisatie
715205	H-14-11	Rare neoplastic diseases, Rare neurological diseases	Pudendal neuralgia	60039	P		Zonder patiëntenorganisatie
715227	H-3-17	Rare endocrine diseases	Meningioma	2495	P	34	Osteoporose Vereniging
715227	H-3-17	Rare endocrine diseases	Acquired premature ovarian failure	95709	P	34	Osteoporose Vereniging
715227	H-3-17	Rare endocrine diseases	Meningioma	2495	P	64	NVACP

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715227	H-3-17	Rare endocrine diseases	Meningioma	2495	P	65	Nederlandse Hypofyse Stichting (NHS)
715227	H-3-17	Rare endocrine diseases	Meningioma	2495	P	91	Hersenletsel.nl
715227	H-3-17	Rare endocrine diseases	Acquired premature ovarian failure	95709	P	312	Patiëntenvereniging POI-POF
715254	H-20-8	Rare hepatic diseases	Autoimmune hepatitis	2137	P	19	Nederlandse Leverpatiëntenvereniging (NLV)
715254	H-20-8	Rare hepatic diseases	IgG4-related sclerosing cholangitis	447764	P	19	Nederlandse Leverpatiëntenvereniging (NLV)
715254	H-20-8	Rare hepatic diseases	Primary biliary cholangitis	186	P	19	Nederlandse Leverpatiëntenvereniging (NLV)
715254	H-20-8	Rare hepatic diseases	Primary sclerosing cholangitis	171	P	19	Nederlandse Leverpatiëntenvereniging (NLV)
715274	H-8-23	Rare hepatic diseases, Rare respiratory diseases	Alpha-1-antitrypsin deficiency	60	P	121	Longfonds

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715304	H-2-15	Rare otorhinolaryngological diseases	Cranio-cervical dystonia with laryngeal and upper-limb involvement	420485	P	59	Dystonievereniging
715304	H-2-15	Rare otorhinolaryngological diseases	Congenital laryngeal cyst	141124	P	59	Dystonievereniging
715304	H-2-15	Rare otorhinolaryngological diseases	Laryngeal abductor paralysis	2808	P	59	Dystonievereniging
715304	H-2-15	Rare otorhinolaryngological diseases	Larynx anomaly	156249	P	59	Dystonievereniging
715304	H-2-15	Rare otorhinolaryngological diseases	Congenital subglottic stenosis	141121	P	59	Dystonievereniging
715304	H-2-15	Rare otorhinolaryngological diseases	Congenital subglottic stenosis	141121	P	114	Care4Neo
715332	H-16-12	Rare cardiac diseases	Arrhythmogenic right ventricular cardiomyopathy	247	P	58	Harteraad

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715332	H-16-12	Rare cardiac diseases	Dilated cardiomyopathy	217604	P	58	Harteraad
715332	H-16-12	Rare cardiac diseases	Rare cardiomyopathy	167848	P	58	Harteraad
715332	H-16-12	Rare cardiac diseases	Rare hypertrophic cardiomyopathy	217569	P	58	Harteraad
715332	H-16-12	Rare cardiac diseases	Arrhythmogenic right ventricular cardiomyopathy	247	P	206	Stichting ICD Dragers Nederland (STIN)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715332	H-16-12	Rare cardiac diseases	Dilated cardiomyopathy	217604	P	206	Stichting ICD Dragers Nederland (STIN)
715332	H-16-12	Rare cardiac diseases	Rare cardiomyopathy	167848	P	206	Stichting ICD Dragers Nederland (STIN)
715332	H-16-12	Rare cardiac diseases	Rare hypertrophic cardiomyopathy	217569	P	206	Stichting ICD Dragers Nederland (STIN)
715332	H-16-12	Rare cardiac diseases	Arrhythmogenic right ventricular cardiomyopathy	247	P	338	Stichting LMNA Cardiac

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715332	H-16-12	Rare cardiac diseases	Dilated cardiomyopathy	217604	P	338	Stichting LMNA Cardiac
715332	H-16-12	Rare cardiac diseases	Rare cardiomyopathy	167848	P	338	Stichting LMNA Cardiac
715332	H-16-12	Rare cardiac diseases	Rare hypertrophic cardiomyopathy	217569	P	338	Stichting LMNA Cardiac
715366	H-11-52	Rare neoplastic diseases	Rare vulvovaginal tumor	180312	P	18	NFK: Patiëntenplatform Zeldzame Kankers
715366	H-11-52	Rare neoplastic diseases	Rare hepatic and biliary tract tumor	101943	P	19	Nederlandse Leverpatiëntenvereniging (NLV)
715366	H-11-52	Rare neoplastic diseases	Benign peripheral nerve sheath tumor	252131	P	30	Neurofibromatosevereniging Nederland (NFVN)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715366	H-11-52	Rare neoplastic diseases	Malignant peripheral nerve sheath tumor	3148	P	30	Neurofibromatosevereniging Nederland (NFVN)
715366	H-11-52	Rare neoplastic diseases	Rare gastroesophageal tumor	180821	P	63	SPKS - Leven met Maag- en Slokdarmkanker
715366	H-11-52	Rare neoplastic diseases	Glial Tumors	182067	P	82	Vereniging Kinderkanker Nederland
715366	H-11-52	Rare neoplastic diseases	Rare carcinoma of pancreas	217074	P	87	Alvleeskliervereniging
715366	H-11-52	Rare neoplastic diseases	Rare tumor of pancreas	180824	P	87	Alvleeskliervereniging
715366	H-11-52	Rare neoplastic diseases	Rare gynaecological tumor	98063	P	88	Stichting Olijf
715366	H-11-52	Rare neoplastic diseases	Rare uterine cancer	213564	P	88	Stichting Olijf

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715366	H-11-52	Rare neoplastic diseases	Rare vulvovaginal tumor	180312	P	88	Stichting Olijf
715366	H-11-52	Rare neoplastic diseases	Rare ovarian cancer	213500	P	88	Stichting Olijf
715366	H-11-52	Rare neoplastic diseases	Glial Tumors	182067	P	91	Hersenletsel.nl
715366	H-11-52	Rare neoplastic diseases	Malignant peripheral nerve sheath tumor	3148	P	94	Stichting Patiëntenplatform Sarcomen
715366	H-11-52	Rare neoplastic diseases	Malignant melanoma of the mucosa	168999	P	96	Stichting Melanoom
715366	H-11-52	Rare neoplastic diseases	Rare carcinoma of pancreas	217074	P	260	Living with Hope
715366	H-11-52	Rare neoplastic diseases	Rare tumor of pancreas	180824	P	260	Living with Hope
715386	H-12-18	Rare hematologic diseases	Systemic mastocytosis	2467	P	83	Mastocytose Vereniging Nederland

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
715430	H-2-8	Rare developmental anomalies during embryogenesis, Rare neurological diseases Rare otorhinolaryngological diseases Rare surgical maxillo-facial diseases	Paralytic facial malformation	156224	P	17	LaPosa
715450	H-3-18	Rare endocrine diseases, Rare gynecologic and obstetric diseases	Non-acquired premature ovarian failure	95710	P	312	Patiëntenvereniging POI-POF
715486	H-10-9	Rare skin diseases	Large congenital melanocytic nevus	626	P	141	Nevus Netwerk
716037	H-18-1	Rare inborn errors of metabolism	Rare dyslipidemia	101953	P	43	Spierziekten Nederland (SN)
716037	H-18-1	Rare inborn errors of metabolism	Rare dyslipidemia	101953	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
716058	H-2-2	Rare surgical maxillo-facial diseases	Oculo-auriculo-vertebral spectrum	141132	P	17	LaPosa

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
716078	H-24-2	Rare circulatory system diseases	Congenital chylothorax	264688	P	114	Care4Neo
716078	H-24-2	Rare circulatory system diseases	Congenital chylothorax	264688	P	200	LGD Alliance
716098	H-24-6	Rare circulatory system diseases	Aortic malformation	98718	P	147	Contactgroep Marfan Nederland
716098	H-24-6	Rare circulatory system diseases	Ascending aorta anomaly	98725	P	147	Contactgroep Marfan Nederland
716098	H-24-6	Rare circulatory system diseases	Rare disease with thoracic aortic aneurysm and aortic dissection	285014	P	147	Contactgroep Marfan Nederland
716098	H-24-6	Rare circulatory system diseases	Aortic malformation	98718	P	148	Vereniging van Ehlers Danlos Patiënten (VED)
716098	H-24-6	Rare circulatory system diseases	Ascending aorta anomaly	98725	P	148	Vereniging van Ehlers Danlos Patiënten (VED)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
716098	H-24-6	Rare circulatory system diseases	Rare disease with thoracic aortic aneurysm and aortic dissection	285014	P	148	Vereniging van Ehlers Danlos Patiënten (VED)
716098	H-24-6	Rare circulatory system diseases	Aortic malformation	98718	P	309	EDS Fonds
716098	H-24-6	Rare circulatory system diseases	Ascending aorta anomaly	98725	P	309	EDS Fonds
716098	H-24-6	Rare circulatory system diseases	Rare disease with thoracic aortic aneurysm and aortic dissection	285014	P	309	EDS Fonds
716098	H-24-6	Rare circulatory system diseases	Rare disease with thoracic aortic aneurysm and aortic dissection	285014	P	315	Stichting Hersenaneurysma Patiënten Platform
716120	H-11-16	Rare endocrine diseases	Non-functioning paraganglioma	94080	P	29	Nederlandse Vereniging voor Mensen met Paragangliomen (NVPG)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
716120	H-11-16	Rare endocrine diseases	Non-functioning paraganglioma	94080	P	64	Bijniervereniging NVACP
716145	H-12-3	Rare neoplastic diseases	Myeloproliferative neoplasm	98274	P	78	MPN Stichting
716145	H-12-3	Rare neoplastic diseases	Primary myelofibrosis	824	P	78	MPN Stichting
716145	H-12-3	Rare neoplastic diseases	Acute myeloid leukemia	519	P	81	Stichting Hematon
716145	H-12-3	Rare neoplastic diseases	Chronic myeloid leukemia	521	P	81	Stichting Hematon
716145	H-12-3	Rare neoplastic diseases	Chronic myelomonocytic leukemia	98823	P	81	Stichting Hematon
716145	H-12-3	Rare neoplastic diseases	Myelodysplastic syndrome	52688	P	81	Stichting Hematon

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
716145	H-12-3	Rare neoplastic diseases	Myeloid hemopathy	171895	P	81	Stichting Hematon
716145	H-12-3	Rare neoplastic diseases	Myeloproliferative neoplasm	98274	P	81	Stichting Hematon
716145	H-12-3	Rare neoplastic diseases	Primary myelofibrosis	824	P	81	Stichting Hematon
716165	H-25-2	Rare infectious diseases	Rare mycoses	163591	P	72	Stichting voor Afweerstoornissen (SAS)
716165	H-25-2	Rare infectious diseases	Rare mycoses	163591	P	161	Huid Nederland
717024	H-11-51	Rare neoplastic diseases	Germ cell tumor	3399	P	226	Stichting Zaadbalkanker
719220	H-7-15	Rare gastroenterologic diseases, Rare transplant-related disorders, Rare abdominal surgical	Chronic intestinal failure	294422	P	123	Vereniging Ziekte van Hirschsprung
719220	H-7-15	Rare gastroenterologic diseases	Chronic intestinal pseudoobstruction	2978	P	123	Vereniging Ziekte van Hirschsprung

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
719220	H-7-15	Rare gastroenterologic diseases	Secondary short bowel syndrome	95427	P	123	Vereniging Ziekte van Hirschsprung
719220	H-7-15	Rare gastroenterologic diseases	Parenteral nutrition-associated cholestasis	567983	P		zonder patiëntenorganisatie
719220	H-7-15	Rare gastroenterologic diseases	Microvillus inclusion disease	2290	P		Zonder patiëntenorganisatie
720077	H-16-13	Rare cardiac diseases	Aortic malformation	98718	P	127	Patiëntenvereniging voor Aangeboren Hartafwijkingen (PAH)
720077	H-16-13	Rare cardiac diseases	Conotruncal heart malformations	2445	P	127	Patiëntenvereniging voor Aangeboren Hartafwijkingen (PAH)
720077	H-16-13	Rare cardiac diseases	Rare congenital non-syndromic heart malformation	88991	P	127	Patiëntenvereniging voor Aangeboren Hartafwijkingen (PAH)
720077	H-16-13	Rare cardiac diseases	Transposition of the great arteries	216675	P	127	Patiëntenvereniging voor Aangeboren Hartafwijkingen (PAH)
720077	H-16-13	Rare cardiac diseases	Univentricular cardiopathy	95483	P	127	Patiëntenvereniging voor Aangeboren Hartafwijkingen (PAH)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
720256	H-12-11	Rare hematologic diseases	Dyskeratosis Congenita	1775	P	161	Huid Nederland
724127	H-2-12	Rare surgical maxillo-facial diseases	22q11.2 deletion syndrome	567	P	17	LaPosa
724127	H-2-12	Rare surgical maxillo-facial diseases	22q11.2 duplication syndrome	1727	P	17	LaPosa
724127	H-2-12	Rare surgical maxillo-facial diseases	Isolated Pierre Robin syndrome	718	P	17	LaPosa
724127	H-2-12	Rare surgical maxillo-facial diseases	Rare disease with Pierre Robin syndrome	138044	P	17	LaPosa
724127	H-2-12	Rare surgical maxillo-facial diseases	22q11.2 deletion syndrome	567	P	271	Stichting Steun 22q11
724127	H-2-12	Rare surgical maxillo-facial diseases	22q11.2 duplication syndrome	1727	P	271	Stichting Steun 22q11

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724127	H-2-12	Rare surgical maxillo-facial diseases	Isolated Pierre Robin syndrome	718	P	279	Stichting Pierre Robin Europe
724127	H-2-12	Rare surgical maxillo-facial diseases	Rare disease with Pierre Robin syndrome	138044	P	279	Stichting Pierre Robin Europe
724127	H-2-12	Rare surgical maxillo-facial diseases	Isolated Pierre Robin syndrome	718	P	287	Schisis Nederland
724127	H-2-12	Rare surgical maxillo-facial diseases	Rare disease with Pierre Robin syndrome	138044	P	287	Schisis Nederland
724182	H-11-32	Rare neoplastic diseases	Vulvar carcinoma	494418	P	88	Stichting Olijf
724182	H-11-32	Rare neoplastic diseases	Malignant graniulosacel tumor of ovary	99915	P	88	Stichting Olijf
724182	H-11-32	Rare neoplastic diseases	Rare cancer of the cervix uteri	213761	P	88	Stichting Olijf
724182	H-11-32	Rare neoplastic diseases	Rare gynecological tumor	98063	P	88	Stichting Olijf
724182	H-11-32	Rare neoplastic diseases	Ovarian cancer	213500	P	88	Stichting Olijf

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724234	H-15-11	Rare neoplastic diseases	Benign schwannoma	252164	P	30	Neurofibromatosevereniging Nederland (NFVN)
724234	H-15-11	Rare neoplastic diseases	Vestibular schwannoom	252175	P	30	Neurofibromatosevereniging Nederland (NFVN)
724234	H-15-11	Rare neoplastic diseases	Benign schwannoma	252164	P	33	Stichting Hoormij
724234	H-15-11	Rare neoplastic diseases	Vestibular schwannoom	252175	P	33	Stichting Hoormij
724234	H-15-11	Rare neoplastic diseases	Glial tumor	182067	P	33	Stichting Hoormij
724234	H-15-11	Rare neoplastic diseases	Rare tumor of neuroepithelial tissue	251558	P	33	Stichting Hoormij
724234	H-15-11	Rare neoplastic diseases	Meningioma	2495	P	33	Stichting Hoormij

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724234	H-15-11	Rare neoplastic diseases	Glial tumor	182067	P	82	Vereniging Kinderkanker Nederland
724234	H-15-11	Rare neoplastic diseases	Rare tumor of neuroepithelial tissue	251558	P	82	Vereniging Kinderkanker Nederland
724234	H-15-11	Rare neoplastic diseases	Glial tumor	182067	P	91	Hersenletsel.nl
724234	H-15-11	Rare neoplastic diseases	Meningioma	2495	P	91	Hersenletsel.nl
724296	H-9-13	Rare neurological diseases	Moyamoya angiopathy	477768	P		Zonder patiëntenorganisatie
724296	H-9-13	Rare neurological diseases	Moyamoya disease	2573	P		Zonder patiëntenorganisatie
724346	H-15-9	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	88	Stichting Olijf
724346	H-15-9	Rare genetic diseases	Hereditary site-specific ovarian cancer syndrome	213524	P	88	Stichting Olijf

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724346	H-15-9	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	89	Borstkankervereniging (BVN)
724346	H-15-9	Rare genetic diseases	Hereditary breast cancer	227535	P	89	Borstkankervereniging (BVN)
724346	H-15-9	Rare genetic diseases	Hereditary breast and ovarian cancer syndrome	145	P	339	Stichting Erfelijke Kanker Nederland (SEKN)
724406	H-9-12	Rare genetic diseases	Bilateral striopallidodentate calcinosis	1980	P	2	Alzheimer Nederland
724406	H-9-12	Rare genetic diseases	Bilateral striopallidodentate calcinosis	1980	P	346	Fahr Patiënten Vereniging Nederland
724467	H-11-55	Rare neurological diseases	Neonatal hypoxic and ischemic brain injury	137577	P	114	Care4Neo
724467	H-11-55	Rare neurological diseases	Neonatal hypoxic and ischemic brain injury	137577	P	284	CP Nederland
724467	H-11-55	Rare neurological diseases	Cerebral sinovenous thrombosis	329217	P		Zonder patiëntenorganisatie

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724467	H-11-55	Rare neurological diseases	Pediatric arterial ischemic stroke	439175	P		Zonder patiëntenorganisatie
724526	H-11-50	Rare neoplastic diseases	Nasopharyngeal carcinoma	150	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Rare tumor of salivary glands	276142	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of oral cavity and lip	502369	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of the hypopharynx	494547	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of the larynx	494550	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of the nasal cavity and paranasal sinuses	500464	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of the oral cavity	502363	P	169	Patiëntenvereniging Hoofd - Hals

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724526	H-11-50	Rare neoplastic diseases	Squamous cell carcinoma of the oropharynx	500478	P	169	Patiëntenvereniging Hoofd - Hals
724526	H-11-50	Rare neoplastic diseases	Rare tumor of salivary glands	276142	P	255	Patiëntenvereniging Speekselklierkanker
724576	H-11-41	Rare neoplastic diseases	Desmoid tumor	873	P	94	Stichting Patiëntenplatform Sarcomen
724576	H-11-41	Rare neoplastic diseases	Soft tissue	3394	P	94	Stichting Patiëntenplatform Sarcomen
724628	H-11-31	Rare genetic diseases	Gastro-intestinale stromale tumor	44890	P	94	Stichting Patiëntenplatform Sarcomen
724687	H-18-11	Rare inborn errors of metabolism	Classic organic aciduria	79163	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	CLN3 disease	228346	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724687	H-18-11	Rare inborn errors of metabolism	Hurler syndrome	93473	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Juvenile neuronal ceroid lipofuscinosis	79264	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Lysosomal disease	68366	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Methylmalonic acidemia with homocystinuria	26	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Methylmalonic acidemia without homocystinuria	293355	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Neonatal epileptic encephalopathy due to glutaminase deficiency	557064	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)

AIMS ID:	Assessment Number:	Expertisegebied:	Aandoening:		P	nr	Patiëntenorganisatie:
724687	H-18-11	Rare inborn errors of metabolism	Neuronal ceroid lipofuscinosis	216	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Propionic acidemia	35	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	Rare inborn errors of metabolism	68367	P	56	Volwassenen Kinderen en Stofwisselingsziekten (VKS)
724687	H-18-11	Rare inborn errors of metabolism	CLN3 disease	228346	P	310	Beat Batten
724687	H-18-11	Rare inborn errors of metabolism	Juvenile neuronal ceroid lipofuscinosis	79264	P	310	Beat Batten