Medical Conditions:

Cerebellitis, 2-Hydroxyglutraic aciduria, 3M syndrome, 9p-Syndrome (Monosomy 9p, Alfi's Syndrome), ADHD, Aarskog syndrome, Abdominal Migraine, Abdominal pain, Abnormality to hands/fingers, Absence seizures (Petit Mal), Achondroplasia, Acne (acne vulgaris), Acquired brain injury (ABI), Acrophobia, Acute disseminated encephalomyelitis (ADEM), Acute lymphoblastic leukaemia (ALL), Addison's disease (primary adrenal insufficiency, hypocortisolism), Adenomyosis, Adrenal Crisis (Addisonian Crisis), Adrenal insufficiency, Afebrile seizures, Agenesis of the corpus callosum (ACC), Aicardi syndrome, Alagille syndrome, Albinism, Albinism, Allergic reaction (unknown), Allergic conjunctivitis, Allergy (please specify), Alopecia (hair loss), Alpha 1-antitrypsin deficiency (A1AD), Alpha Thalassemia, Alpha Tryptasemia Syndrome, Alpha-1 Carrier, Amblyopia (lazy eye), Amelogenesis Imperfecta, Amniotic Band Syndrome (ABS), Anaemia, Anaphylactic Food Allergy, Anaphylaxis, Anderson-Fabry disease, Angelman syndrome (AS), Angioedema, Angiomatoid Fibrous Histiocytoma, Anisometropia, Ankle Instability, Ankle-Foot Orthosis (AFO, ankle brace), Ankyloglossia (Tongue-tie), Anorexia nervosa, Anti-thyroid antibodies, Antibody Disorder, Anxiety, Anxiety/Anxiety Disorder, Aortic Stenosis (AS), Aortic insufficiency (Aortic regurgitation, leaky aortic valve), Apert Syndrome, Aphakia, Aphthous stomatitis (Aphthous Ulcers), Aplasia cutis congenita (ACC), Aplastic Anaemia, Apnoea (Apnea), Appendicitis, Arnold-Chiari malformation, Arrhytmogenic right ventricular cardiomyopathy (ARVC), Arteriovenous Malformation (AVM, weakened blood vessels), Arthritis, Arthrogryposis multiplex congenital (AMC, multiple joint contractures), Artificial Limb (Prosthesis), Asperger Syndrome, Aspirin Sensitivity/Intolerance (Salicylate), Asthma, Astigmatism, Astrocytoma (Glioma Brain Tumor), Ataxia, Ataxia-telangiectasia (A-T, Louis Bar syndrome), Atrial Isomerism, Atrial Septal Defect (ASD, Hole in heart), Attachment Disorder, Attention Deficit Hyperactivity Disorder (ADHD), Auditory Processing Disorder (APD), Autism/Austistic Spectrum Disorder, Autistic enterocolitis, Autoimmune lymphoproliferative syndrome (ALPS, Canale-Smith syndrome), Autoinflammatory condition, Autonomic Dysreflexia (Hyperreflexia), Avoidant Restrict Food Intake Disorder (ARFID), Back Pain, Bacterial vaginosis, Balance problem, Balbar Palsy, Baraitser-Winter syndrome, Bardet_Biedl syndrome, Bartter Syndrome, Batten Disease, Beckwith-Wiedemann Syndrome, Behavioural Emotional and Social Difficulties, Behcet's Disease (Behcet's Syndrome), Bell's Palsy, Benign Paroxysmal Positional Vertigo, Benign Reactive Lymphadenopathy/Reactive Lymph nodes, Benign rolandic epilepsy (BCECTS), Beta Thalassemia, Beta thalassemia trait/carrier, Bicuspid aortic valve (BAV), Bilateral Brachial Plexus Palsy, Bilateral Retinoblastoma (Rb), Bilateral hearing loss (BHL), Bipolar Disorder, Bipolar I Disorder, Bipolar II Disorder (BP-II), Birthmark, Bladder condition, Blepharitis, Blindness (Vision loss), Blood Vessel Condition, Body dysmorphic disorder (BDD, Dysmorphia), Body temperature regulation disorder, Bone Defect, Bone Hypertrophy, Bone Marrow Transplant, Bone disease, Bowel condition, Bowel obstruction (intestinal obstruction), Brachydactyly, Bradycardia (Low Heart Rate), Brain Aneurysm, Brain Development Condition, Brain Injury, Brain Tumor, Brain/spinal cord cyst, Branchiootorenal (BOR) Syndrome, Breath-holding spells (Episodic apnea), Breathing difficulties, Broccoli intolerance, Broken Bone, Bronchiectasis, Bronchioltis, Bronchitis, Broncholithiasis, Bronchomalacia, Bronchospasm, Brown's syndrome, Brugada Syndrome (BrS), Bulimia nervosa, Burns, Börjeson-Forssman-Lehmann syndrome (BFLS), CHARGE Syndrome, Cancer, Cardiac arrhythmia (Irregular heartbeat, heart rhythm disorders), Cardiomyopathy, Carnitine Palmitoyltransferase II deficiency (CPT-11, CPT2), Cataplexy, Cataract, Caudal regression syndrome (sacral agenesis, hypoplasia of the sacrum), Cavernous hemangioma (Cavernoma of the brain/spinal cord), Cellulitis, Cerebral Palsy, Cerebral Venous Sinus Thrombosis (CVST), Cerebral edema/oedema, Cerebrocostomandibular Syndrome, Cervical Intraepithelial Neoplasia, Cervical Kyphosis (Military/straight neck), Channelopathy, Charcot-Marie-Tooth disease (CMT), Chest Infection, Chest/Lung Condition, Chicken Pox, Chilblains, Chondromalacia patellae (CMP), Chromosome 2q37 deletion syndrome, Chromosome 6 Abnormality, Chromosome 9p deletion syndrome, Chromosome 9p duplication, Chromosome 12q deletion, Chromosome 14 Trisomy Mosaic, Chromosome 17 trisomy, Chromosome 19p13.2 microdeletion, Chromosome 22 Trisomy, Chromosome Deletion No.9, Chromosome abnormality (anomaly, aberration or mutation), Chronic Constipation, Chronic Elbow Instability (Recurrent dislocations), Chronic Fatigue Syndrome (CFS, ME, PVFS), Chronic Inflammatory Demyelinating Polyneuropathy (CIDP), Chronic Kidney Disease (CKD, Chronic Renal Failure), Chronic Lung Disease (CLD), Chronic Motor Tic Disorder, Chronic granulomatous disease (CGD), Chronic idiopathic constipation, Chronic obstructive pulmonary disease (COPD), Chronic pain, Chronic pulmonary aspiration (CPA), Cibophobia (Food phobia), Circulation condition/issues, Citrullinemia, Claustrophobia, Cleft Lip (CL), Cleft Palate (CP), Cleidocranial dysostosis (CCD, cleidocranial dysplasia), Clinical Obesity, Club Foot/Positional Talipes (PTEV), Club Foot/Structural Talipes (CTEV), Cobalamin F Defect, Cochlear Implant due to Deafness, Cochlear Implantation after kidney transplantation, Cockayne Syndrome (CS), Coeliac Disease (Gluten free), Coffin-Lowry syndrome, Coffin-Siris syndrome, Cohen syndrome (Pepper syndrome), Cold Sores, Colitis, Collapsed Lung, Coloboma, Colostomy Bag Fitted, Colour Blindness, Comorbid Disorders/Diseases, Conduct disorder (CD), Conductive hearing loss, Congenital Cataract, Congenital Diaphragmatic Hernia, Congenital Heart Disease/Defect, Congenital Hip Dislocation (Dysplasia), Congenital Hypoplasia, Congenital Neutropenia, Congenital adrenal hyperplasia (CAH), Congenital hepatic fibrosis (CHF), Congenital melanocytic nevus, Congenital myasthenic syndrome (CMS), Congenital

toxoplasmosis, Conjunctivitis, Cornelia de Lange Syndrome (CdLS), Coronary Artery Fistula, Corpus Callosum Dysgenesis, Costochondritis (chest wall pain), Coulrophobia, Cowden syndrome, Cradle cap, Craniosynostosis, Crepitus, Cri-du-chat (cat's cry) syndrome (5p-syndrome), Chron's Disease (Regional Enteritis), Croup (Laryngotracheobronchitis), Cutis Laxa, Cutis Marmorata (CMTC) Cyanosis, Cyclical Vomiting Syndrome (CVS), Cyclothymia, Cystic Fibrosis, Cystic hygroma, Cytochrome C Oxidase Deficiency, Dandy-Walker Syndrome (DWS), De Grouchy Syndrome, Deaf, Deep vein thrombosis (DVT), Deficits in attention, motor control and perception (DAMP), Degenerative disease, Delayed Swallowing, Dental condition (Tooth pathology), Dentinogenesis imperfecta, Depression, Dermoid cyst, Detrusor Instability, Developmental Delay, Dextrocardia, DiGeorge syndrome (22q11.2 deletion syndrome), Diabetes, Diamond Blackfan Anemia (DBA), Diarrhea condition, Digestive Disorder, Dilated cardiomyopathy (DCM), Diplegia, Disabled Hand, Disciform keratitis, Disorder of Sexual Development, Disruptive Behaviour Disorder, Dizziness, Double Incontinence (DI), Down Syndrome, Dravet syndrome (severe myoclonic epilepsy of infancy, SMEI), Drooling, Duane syndrome, Dubin-Johnson syndrome (DJS), Durchenne muscular dystrophy (DMD), Duplex kidneys, Dysautonomia (autonomic dysfunction), Dyscalculia, Dyscalculia, Dysfibrinogenemia, Dysgraphia, Dyslexia, Dysphagia (swallowing problems), Dysphasia (Communications Difficulties), Dysplasia (Development disorder), Dyspraxia (Development Coordination Disorder), Dystonia (Muscle disorder), ENT (Ear Nose Throat) problems, Ear Infection, Ear Considition, Eating Problem, Ectopic Atrial Tachycardia, Eczema/Dermatitis, Edema (oedema), Edwards syndrome (trisomy 18), Ehler-Danlos syndrome (EDS), Elevated gonadotropins, Emetophobia (vomiting fear), Emotional Detachment Disorder, Encephalitis, Encopresis, Endometriosis, Enlarged adenoids (Adenoid hypertrophy), Enterovirus Myocarditis, Enuresis/Urinary Incontinence (weak bladder), Eosinophilic Colitis, Eosinophilic Oesophagitis, Eosinophilic gastroenteritis (EG), Epidermolysis bullosa simplex (EBS), Epilepsy, Epileptic spasms (West syndrome, infantile spasms), Epstein-Barr Virus (EBV), Erb's-palsy (Erb-Duchenne palsy), Erythema Nodosum (EN), Exocrine pancreatic insufficiency (EPI), Exophoria, Exophthalmos (exophthalmia, proptosis, exorbitism), Exostoses, Exotropia (Divergent Squint), Eye condition, Facial Palsy, Factor V Leiden, Factor V deficiency (Owren's disease, parahemophilia), Factor VII deficiency, Factor X deficiency, Factor XII deficiency (Hageman factor deficiency, Fainting, Faints at sight / mentions of blood, Familial Mediterranean Fever, Familial exudative vitreoretinopathy (FEVR) Familial hypercholesterolemia (FH), Fanconi Syndrome, Fanconi Anaemia (FA), Fat intolerance, Favism (G6PD deficiency, Glucose-6-phosphate dehydrogenase deficiency, Febrile Seizures/Convulsions, Fecal Impaction, Femoral Anteversion (Pigeon Toe), Fetal Valproate syndrome, Fetal alcohol syndrome (FAS), Fibrodysplasia ossificans progressiva (FOP), Fibromyalgia, Fissure, Flat Feet (Dropped/Fallen Arches, Plano-valgus foot), Foetal Anticonvulsant Syndrome (FACS), Follicular hyperplasia, Food intolerance, Foot development problems, Fowler's Syndrome, FoxG1 Syndrome, Fragile X syndrome (FXS), Fragrance/Perfume Sensitivity, Frey's syndrome, Fructose malabsorption (dietary fructose intolerance), Fucosidosis, Fundoplication in place, GLUT1 deficiency syndrome (Glucose transporter type 1 deficiency), Gag Reflex Condition, Gallstones (Choleithiasis), Ganglioneuroblastoma, Gastritis, Gastroesophageal Reflux (Gastric Reflux, GERD), Gastrointestinal Disorder, Gastrostomy, General Infection, General blood disorder, Genetic Disorder), Genu recurvatum (knee hyperextension), Geographic tongue (benign migratory glossitis, erythema migrans lingualis), Gilbert syndrome (Liver Disorder), Glandular Fever (Infectious mononucleosis), Glaucoma, Glenoid hypoplasia, Glomerulosclerosis (Renal/Kidney scarring), Glucose-Galactose Malabsorption, Glue Ear (Otitis Media), Glutaric aciduria type 1 (GA1), Gluten Intolerance, Glycerol Kinase Deficiency (GKD), Glycogen Storage Disease (GSD), Glycogen storage disease type IX (GSD IX), Goldenhar syndrome (Oculo-Aurícula-Vertebral (OAV) syndrome, Gomez Lopez syndrome, Gordon Syndrome, Gorlin syndrome, Granuloma annulare, Graves Disease (Toxic diffuse goiter), Grommets in ear, Guanidinoacetate methyltransferase (GAMT) deficiency, Guillain-Barre syndrome (GBS), Guttate Psoriasis, Gynaecological condition, Gynaecomastia, Haemangiomas, Haemoglobin D carrier, Haemoglobin E carrier, Haemophilia, Hair Pulling Disorder (Trichotillomania, TTM), Hameodialysis Catheter (Haemocath), Hand or Elbow condition, Hayfever, Hb Koln (blood disorder), Head Injury (cannot do contact sports), Hearing loss, Hearing problem, Heart Block, Heart Murmur, Heart Palpitations, Heart condition, Heartburn (Pyrosis), Hemihypertrophy (Hemihyperplasia), Hemiparesis (weakness on one side of body), Hemiplegia, Hemiplegic migraines, Hemoglobin H disease, Henoch-Schontein Purpura (HSP, Anaphylactoid Purpura), Hepatitis, Hepatitis B, Hereditary Angioedema (HAE), Hereditary motor and sensory neuropathy (HMSN), Hereditary motor and sensory neuropathy type 1A (HMSN 1A), Hereditary multiple exostoses (HME, MHE, diaphyseal aclasis), Hereditary neuropathy with pressure palsies, Hereditary spastic paraplegia (HSP), Hernia, Herpes virus, Heterochromia, Hickman Line, High pain threshold, Hip Dislocation (Dysplasia), Hip problem, Hirschsprung's disease (HD), Hirschsprung-associated enterocolitis (HAEC), Hodgkin Lymphoma, Holoprosencephaly, Holt-Oram Syndrome, Homocystinuria (CBS deficiency), Hormone injections, (Horner's syndrome (Bernard-Horner syndrome), Hunter syndrome (mucopolysaccharidosis type II, MPS II), Hurler syndrome (mucopolysaccharidosis type IH, MPS IH), Hurler-Scheie syndrome (mucopolysaccharidosis type I, H-S), Hutchinson-Gilford Progeria, Hyaluronidase deficiency (mucopolysaccharidosis type IX, MPS IX), Hydrocephalus (water on the brain), Hydromyelia, Hyperacusis, Hypercholesterolemia (High Cholesterol), Hyperglycaemia (High blood-sugar level),

Hyperhidrosis (Excessive sweating), Hyperinsulinism, Hyperplasia, Hypersensitivity, Hypertension (High blood pressure), Hyperthyroidism (overactive thyroid), Hypertonia, Hypertrophic cardiomyopathy (HCM), Hypocalcemia (low calcium), Hypoglycaemia (Low blood sugar level), Hypoparathyroidism (HypoPARA), Hypopituitarism, Hypoplastic left heart syndrome (HLHS), Hypospadias, Hypotension (low blood pressure), Hypothyroidism (underactive thyroid), Ichthyosis (skin disorder), Idiopathic intracranial hypertension, Idiopathic thrombocytopenic purpura (ITP), IgA Nephropathy (Berger's Disease), Immune system disorder, Immunodeficiency (Immune deficiency), Immunosuppression, Impetigo, Impulsive Aggression, In-toe walking (Gait, turned-in-feet), Inappropriate sinus tachycardia (IST), Incontinentia Pigmenti - genetic disorder, Indeterminate Colitis (IC), Inflammatory Bowel Disease (IBD), Inhaler (as and when required), Insomnia, Insulin Injection, Intestinal Atresia, Intestinal malrotation, Iron Deficiency, Irritable Bowel Syndrome (IBS), Irritable Hip, Ischemic cardiomyopathy, Ischiopubic synchondrosis asymmetry, Jacobsen Syndrome (11q/11q24 deletion), Jaundice, Job syndrome (AD-HIES, Autosomal dominant hyper-IgE syndrome), Joint hypermobility, Joint disorder, Joubert syndrome, KBG syndrome, Kabuki syndrome (KBS), Keratitis (inflammation of the cornea), Keratosis pilaris (KP, follicular keratosis, lichen pilaris, chicken skin), Ketotic hypoglycemia, Kidney (Renal) problems, Kidney stone, Kidney transplant, Klinefelter syndrome (KS, 47, XXY), Klippel-Feil syndrome (KFS, Cervical vertebral fusion), Klippel-Trenaunary-Weber syndrome (KTS), Knee cartilage damage, Knee inflammation, Knee problem, Kniest dysplasia (swiss cheese cartilage syndrome, Metatropic Dwarfism Type I), Kyphosis (Hunchback), Kohler disease, LCHAD deficiency (Long-chain 3-hydroxyacyl-coenzyme A dehydrogenase deficiency), Lactose Intolerance, Larsen Syndrome (LS), Laryngo-tracheo-esophageal cleft, Laryngomalacia (soft larynx), Latent coeliac disease, Laurence-Moon syndrome (LMS), Lazy bowel syndrome, Leaky gut syndrome, Left Hemiplegia, Legs and foot problem, Leigh syndrome, Lemierre's syndrome (Lemierre's disease), Lennox-Gastaut syndrome (LGS), Leukaemia, Leukodystrophy, Ligamentous laxity (weak ligaments), Limb amputation, Limited movement in limb, Linear IgA Dermatosis, Lissencephaly, Liver condition, Long QT syndrome (LQTS), Loss of sense of smell (Anosmia), Low energy levels, Low mood, Low muscle tone (hypotonia), Low Platelet count (thrombocytopenia), Lumbar Lordosis, Lumbar spondylitis, Lumbar puncture (LP, Spinal Tap), Lung malformation, Lupus Pernio, Luxating patella (floating knee-caps), Lymph node condition, Lymphangioma, Lymphatic malformations, Lymphoedema, MCADD deficiency (Medium-chain acyl-CoA dehydrogenase deficiency), MCR4 Deficiency, MED13L haploinsufficiency syndrome, Macrocephaly, Maculopapular rash, Malignant Hyperthermia under anesthesia, Malignant hyperthermia (MH) or malignant hyperpyrexia, Maple syrup urine disease, Marcus Gunn Jaw-winking syndrome, Marfan syndrome, Maroteaux-Lamy syndrome (mucopolysaccharidosis type VI, MPS VI), Mastocytoma, Meckel's diverticulum, Medulloblastom (brain tumor), Memory problems, Meningitis, Menstrual condition, Mental health issues (mental illness, psychological disorder), Mesenteric adenitis, Metabolic bone disease, Metabolic disorder, Metatarsus Adductus (Metatarsus Varus), Microcephaly, Micrognathia, Microphthalmia, Microtia, Migraine with aura, Migraines/Headaches, Miliaria (heat rash, sweat rash), Milk protein intolerance, Miller-Dieker syndrome (MDS), Mitochondrial cytopathy, Mitral valve prolapse (MVP), Mobility condition, Molluscum contagiosum (MC) Mongolian spot (Mongolian blue spot), Monocular vision, Mood swings, Morbid obesity, Morning glory disc anomaly (MGDA), Morphea (localised scleroderma), Morquio syndrome (mucopolysaccharidosis IV, MPS, IV), Mosaic Turner syndrome (TS with mosaicism), Motor disorders and stereotypies, Mowat-Wilson syndrome (MWS), Moyamoya disease, Mucocele, Mucopolysaccharidosis type I (MPS I), Muenke syndrome, Multicystic dysplastic kidney (MCDK), Multiple Pituitary Hormone Deficiency (MPHD), Multiple Sclerosis, Multiple endocrine neoplasia (MEN), Multiple endocrine neoplasia type 1 (MEN1, Wermer syndrome), Multiple endocrine neoplasia type 2 (MEN2), Multiple endocrine neoplasia type 2A (MEN 2A, Sipple syndrome), Multiple endocrine neoplasia type 2B (MEN 2B, Wagenmnn-Froboese syndrome), Multiple epiphyseal dysplasia (MED, Fairbank's disease), Mumps, Muscle problem/disorder, Muscle spasms, Muscle Atrophy, Muscular Dystrophy (MD), Musculoskeletal disorder, Myasthenia gravis (MG, muscle fatigue), Myoclonic epilepsy, Myopathy, Myopia (short-sightedness), Myotonic dystrophy, Myotubular Myopathy, Mobius syndrome (Moebius), Nail-patella syndrome (NPS, No kneecaps), Narcolepsy, Nasal septum deviation, Nasogastric feeding tube, Neck problem, Nemaline myopathy, Nephrectomy (Kidney removal), Nephritis (inflammation of the kidneys), Nephrotic syndrome, Neuroblastoma (NB), Neurofibromatosis, Neurofibromatosis type I (NF-1), Neurofibromatosis type II (NF-2), Neurogenic bladder/bowel, Neurological condition/disorder, Neuromuscular condition, Neuronal ceroid lipofuscinoses (NCL), Neuropathic Bladder/Bowel, Neuropathy, Neuropenia, Nissen fundoplication, No blood transfusions, No immunisations/vaccinations given, No oxygen therapy, Non epileptic attack disorder (NEAD), Non-Hodgkin's Lymphoma, Noonan syndrome, Nose disorder/injury, Nosebleeds (Epistaxis), Nystagmus, OPHN1 mutation, Oat intolerance, Obsessive Compulsive Disorder, Occasional rigors/shivers, Oculocutaneous Albinism (OCA), Oculofaciocardiodental syndrome, Oculomotor Apraxia, Oesophageal atresia, Ohdo Blepharophimosis syndrome, Ollier disease, Omenn syndrome, Oppositional Defiant disorder, Optic nerve glioma, Optic nerve hypoplasia (ONH) Oral allergy syndrome, Oral Aversion, Oral Crohn's Disease, Oral Dyspraxia, Oral/Tongue/Mouth condition, Orbital Cellulitis, Orofacial Granulomatosis, Orthostatic hypotension (Postural hypotension), Orthotic Insoles, Osgood-Schlatter Disease (OSD), Osteochondromas, Osteodystrophy, Osteogenesis imperfecta (Brittle bones), Osteomyelitis (OM),

Osteoporosis, Other, PFAPA syndrome (periodic fever, aphthous stomatitis, pharyngitis, adenitis), Pacemaker fitted (heart condition), Pallister-Killian syndrome, Pancreatitis, Panic attacks, Paralysis, Paranoia, Paris-Trousseau syndrome (PTS), Paroxysmal Kinesigenic Dyskinesia (PKD), Paroxysmal dystonia, Paroxysmal Kinesigenic Choreoathetosis/dyskinesia (PKC, PKD), Partially sighted (vision impaired), Patellar dislocation (Dislocated kneecap), Patent Ductus Arteriosus (PDA), Pathological demand avoidance (PDA, Newson's syndrome), Pectus excavatum (funnel chest), Pediatric autoimmune neuropsychiatric disorder associated with streptococcal infections (PANDAS), Pelizaeus-Merzbacher Disease (PMD), Pendred syndrome, Percutaneous endoscopic gastrostomy (PEG) feeding tube, Perforated eardrum, Peri-oral reactions, Period pain (dysmenorrhoea), Peripheral Venous Catheter, Periventricular Heterotopia, Periventricular leukomalacia (PVL), Pernicious anemia (Vitamin B12 deficiency anaemia), Perthes Disease (Hip), Pervasive Refusal syndrome, Peutz Jeghers syndrome (PJS), Pfeiffer syndrome, Phenylketonuria (PKU), Phobia/Fear, Phocomelia, Photophobia (light sensitivity), Photosensitive Epilepsy, Physical Disability, Pica, Pierre Robin Syndrome, Pitt Hopkins syndrome, Pityriasis lichenoides (PL), Plagiocephaly (flat head syndrome), Plantar fasciitis (Jogger's heel), Plexiform neurofibroma, Pneumonia, Poland syndrome, Polyarthiritis, Polycystic kidney disease (PCKD), Polycystic ovary syndrome (PCOS), Polycythemia Vera, Polymicrogyria (PMG), Polymorphous light eruption (PLE, polymorphic light eruption), Polysplenia, Porphyria, Post Brain Tumour, Post Meningitis, Post-op Achilles Tendon Repair, Post-traumatic stress disorder, Postural Orthostatic tachycardia syndrome (POTS), Potocki-Lupski syndrome (PTLS), Prader-Willis syndrome (PWS), Pragmatic language impairment (PLI)/Social communication disorder (SCD), Precocious puberty (early puberty), Premature baby, Premature adrenarche, Priapism, Primary Immune Thrombocytopenia, Primary ciliary dyskinesia (PCD), Primordial Dwarfism, Proctalgia Fugux, Progressive supranuclear palsy (PSP), Proteus syndrome, Prune belly syndrome, Psoriasis, Ptosis (drooping eyelid), Pulled Elbow (nursemaid's elbow, radial head subluxation), Pulmonary Atresia, Pulmonary Haemosiderosis, Pulmonary Vessel under chest bone, Pulmonic stenosis (Pulmonary valve stenosis), Punctate epithelial erosions (punctate erosive keratophy, superficial punctate keratitis), Pyloric stenosis, Pyruvate dehydrogenase deficiency (PDCD), Radioulnar synostosis, Rasmussen's encephalitis (chronic focal encephalitis, CFE), Raynaud's Disease, Reactive airway disease (RAD), Rectus Divarication, Reduced lung volume, Reflex Anoxic Seizures, Reflex sympathetic dystrophy (RSD, CRPS, Complex Regional pain syndrome), Renal agenesis (only one kidney), Renal pelvis dilation (hydronephrosis), Respiratory Disease (respirator condition), Restless leg syndrome, Restrictive cardiomyopathy (RCM), Retinitis Pigmentosa, Retinoblastoma (Rb), Retinopathy, Rett syndrome (RTT), Rheumatic heart disease (RHD), Rheumatoid Arthritis (RA), Rhinitis, Rhombencephalosynapsis, Ricketts, Right Bundle Branch Block (RBBB), Right Hemiplegia, Risk of choking, Rubinstein-Taybi syndrome (RTS, broad thumb-hallux syndrome), Sagittal Synostosis, Salivary gland stones (Sialolithiasis), Salt intolerance, Sandifer syndrome, Sanfilippo syndrome (MPS-III), Scarlet fever, Scheie syndrome (mucopolysaccharidosis I-S, MPS I-S), Scheuermann's disease, Scimitar syndrome, Scleroderma, Scoliosis, Scotopic sensitivity syndrome (SSS, Irlen syndrome), Seasonal Affective Disorder, Seizures (fits), Selective immunoglobulin A (IgA) deficiency (SIgAD), Sensitive skin, Sensory based motor disorder, Sensory discrimination disorder, Sensory Processing Disorder (SPD), Septo-optic dysplasia (SOD, de Morsier syndrome), Sever's disease (Calcaneal apophystis), Severe Factor V11 deficiency, Severe Primary IGF-1 Deficiency, Short bowel syndrome (SBS, short gut), Shoulder condition/injury, Shprintzen syndrome, Shunt, Sickle Cell Anaemia (HbSS), Sickle Cell Disease, Sickel cell trait (Sicklemia), Side effects of regular medication, Silent Aspiration, Silver-Russell dwarfism (russell-Silver syndrome, RSS, SRS), Simpson-Golabi-Behmel syndrome (SGBS, Bulldog syndrome), Sinoatrial Node, Sinusitis, Situs Inversus, Skeletal Dysplasia (Dwarfism), Skin condition, Skin cyst, Skin rash, Skull condition, Sleep apnea, Sleeping disorder, Slipped disc (spinal disc herniation), Sly syndrome (mucopolysaccharidosis, type VII, MPS 7), Small intestine bacterial overgrowth (SIBO, SBBOS), Smith-Lemli-Opitz syndrom, Smith-Magenis syndrome (SMS), Soft spot on head (Fontanelle), Solitary kidney, Sotos syndrome, Spherocytosis, Spina bifida, Spinal injury/disorder (back problem), Spinal muscular atrophy (SMA), Spleen problem, Splenomegaly (enlarged spleen), Stammering, Stap skin infection (staphyloccus infection), Stargardt disease (visual impairment), Stickler syndrome, Stoma, Stomach migraines, Stomach ulcer (Peptic ulcer disease, PUD), Strabismus (squint), Stress cardiomyopathy (Takosubo cardiomyopathy), Stoke, Sturge-Weber syndrome (SWS), Subglottic Stenosis, Suboptimal Cortisol Response, Sucrose intolerance (sucrase-isomaltase deficiency, CSID, GSID), Sudden shock syndrome, Sulphite/Sulfite sensitivity, Supraventricular tachycardia (SVT), Surgery, Susceptible to fractures, Symbrachydactyly, Symphysis pubis dysfunction (SPD), Syndactyly (Webbed Fingers or Toes), Synthetic Fabric / Textiles / Polyester Allergy, T Cell Deficiency, Tall stature, Tardive Dyskinesia, Tear duct problems, Tendonitis, Testicular regression syndrome/vanishing testes (TRS), Tetanus, Tetralogy of Fallot (TOF), Tetrasomy 9p, Thyroid condition, Tic disorder, Tietze syndrom (chondropathia tuberosa), Tight tendons, Tinnitus, Tonsillitis, Torticollis (Wry neck, Loxia), Tourette's syndrome, Tracheal compression, Tracheobronchomalacia (TBM), Tracheoesophageal fistula (TEF, TOF), Tracheomalacia, Tracheostomy, Transcobalamin II deficiency, Travel sickness, Treacher Collins syndrome (TCS), Tremors, Trichothiodystrophy (TTD), Trigger finger/trigger thumb, Trimethylaminuria (MAU, fish odour syndrome), Trismus (Lockjaw), Tuberculosis, Tuberous sclerosis

(TSC), Tumor, Turner syndrome (TS), Ulcerative colitis (UC), Undiagnosed condition, Undifferentiated connective tissue disease (UCTD), Upper lip tie, Urethritis, Urinary Tract Infection (bladder infection), Urinary catheter, Urticaria (hives), Uveitis (Iridocyclitis), VACTERL association, Valvular heart disease, Varices, Vasculitis (inflammation of blood vessels), Vasomotor Instability, Vasovagal response (Vasovagal Syncope), Velocardofacial syndrome (VCFS), Ventricular Tachycardia (VT), Ventricular septal defect (VSD, Hole in the heart), Ventriculomegaly, Vernal keratoconjunctivitis (VKC, spring catarrh), Vertigo, Vesicoureteral Reflux (Kidney reflux), Viral induced wheeze, Vision problems, Vitamin C deficiency, Vitamin D deficiency, Vitiligo, Vocal cord paresis, Volvulus (twisted bowel), Von Hippel-Lindau Syndrome, Von Willebrand disease (VWD), Von Willebrand disease (VWD) Type 1, Wears glasses, Wears hearing aids, Wears an eyepatch, Wears contact lenses, Wheelchair user, Williams syndrome (WS), Wolf-Parkinson-White syndrome (WPW), Wolf-Hirschhorn syndrome (WHS), Worster-Drought syndrome (WDS), x-linked Retinoschisis (XLRS), x-linked adrenal hypoplasia congenita, XYY syndrome