Diagnoses that qualify for CoachArt include everything below

If your child has a physical diagnosis that isn't listed, please contact info@coachart.org to inquire about eligibility. Please note: children with behavioral, mental, and brain development disorders (such as ADHD or autism) do not qualify for CoachArt unless they also have an underlying chronic, physical illness.

chronic, physical liness.
Achondroplasia
Agenesis of the Corpus Callosum
ALCL Lymphoma
Alopecia
ALL- Acute Lymphoblastic Leukemia
Amniotic Bands Syndrome
Aplastic Anemia
Arthrogryposis
Arthritis
Asthma
Astrocytoma Brain Tumor & Glioma
Bilateral Cleft Lip & Palate
Birt-Hogg Dube Syndrome
Bone Marrow Transplant
Brachial Plexus Palsy
Brain Hemmorhage, L Paralysis
Brain Tumor
Brain Tumor Anaplastic Astrocytoma
Burkitt's Lymphoma
Cancer
Cerebral Palsy
Cleft Lip and Palate
Cleft Palate
Cockayne Syndrome
Colitis
Common Variable Immunodeficiency
Complex Partial Seizures
Congenital Adrenal Hyperplasia
Craneofacial Abnormalities
Craniosynostosis
Crohn's Disease
Cryptogenic Generalized Seizure Disorder
Cystic Fibrosis
Dermatomyositis
Diabetes
Diffuse Intrinsic Pontine Glioma
Dystonia
Eczema
Ehlers Danlos
Ellis Van Creveld Syndrome
Embryonic Testicular Rhabdomyosarcoma
End Stage Renal Disease
Ependymoma-Brain Tumor
Epilepsy
Ewings Sarcoma

Fanconi Anemia
Fetal Alcohol Syndrome
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Fibromyalgia GERD
Germ Cell Tumor
Glaucoma
Graves Disease
Heart Disease or Defects
Hemifacial Microsomia and Anotia
Hemoglobin H Constant Spring
Hemolytic Anemia, Thrombotic Thrombocytopenic Purpura Hemophilia
Hepatoblastoma
High Risk GI Reflux
High Risk T-Cell ALL
-
Hirschsprung's Disease Histiocytosis
HIV
Hodgkins Lymphoma
Holocarboxylase Synthetase Deficiency
Hunter Syndrome
Hydrocephalus
Hypoplastic Left Heart Syndrome
Hypopituitarism
Hypothyroidism
Juvenile Ankylosing Spondylitis
Juvenile Rheumatoid Arthritis
Kawasaki Disease
Kidney Disease
Kidney Failure/Transplant
Kostmann's Syndrome
Left Hemifacial Microsomia
Leukemia
Low Grade Astrocytoma
Lung Disease
Lupus
Lymphoblastoma Inflammatory
Lymphoma
Lymphomablastic Leukemia
Mast Cell Activation Syndrome
Medulloblastoma
Menkes Syndrome
Metabolic Acidosis
Metabolism Disorder
Mitochondrial Disease
Mixed Connective Tissue Disease
Mixed Oligoastrocytoma
Moebius Syndrome
Morquio Syndrome
Mucopolysaccharidoses (MPS)
Multiple Congenital Anomolies

Museules Dustee but
Muscular Dystrophy
Neuroblastoma
Neurofibromatosis
Neuromelitis Optic
Niemann Pick Disease
Non-Hodgkins Lymphoma
Optic Glioma
Orafacial Type 1
Orbital Rhabdomysarcoma
Osteomyelitis
Osteosarcoma
Ovarian Tumor
Overlap Syndrome JRA
Pancreatitis
PANDAS
Paralysis
Pierre Robin Sequence/ Marshall Syndrome
Pitt Hopkins Syndrome
PNET Brain Tumor
Polyarticular Juvenile Rheumatoid Arthritis
Pompe Disease
Prader-Willi
Progressive Neuromuscular Degenerative Disorder Psoriatic Arthritis
Renal Failure
Retinoblastoma
Rhabdomyosarcoma
Right Hemifacial Microsomia
Sacral Agenesis- Malformation of the Spine
Sarcoidosis
Sarcoma
SCD-SS Pulmonary Arterial Hypertension
Scleroderma Arthritis
Scoliosis
Severe Aplastic Anemia
Severe Asthma
Severe Hemophilia A
Sickle Cell Anemia
Skin Disease
Spastic Cerebral Palsy
Spina Bifida
Systemic Lupus Erythematosus
Thalassemia
Thrombocytopenia
Treacher Collins Syndrome
Tuberous Sclerosis
Tumor
Turner Syndrome
VACTERL
Von Willebrand Disease
Wiedemann-Steiner Syndrome

Wilm's Tumor Hereditary motor and sensory neuropathy.