Eligible Diagnoses and Diagnostic Codes for Enhanced Care Coordination Pilot

Group Name	DIAGNOSIS	ICD-10 CODES
Traditional Categories		
Blood Disorders		
	F/S Beta Thalassemia (Major)	D56.1
	Delta-Beta Thalassemia	D56.2
	Beta-Thalassemia (Minor)	D56.3
	Hemoglobin E-Beta Thalassemia	D56.5
	Hemoglobin S/Beta Thalassemia	D57.4
	Hemoglobin C	D57.20
	Hemoglobin SC Disease	D57.2
	Hemophilia Hereditary Factor VIII Deficiency	D66
	Hemophilia Hereditary Factor IX Deficiency	D67
	Other Hemoglobinopathies	D58.2
	Hemoglobin SS Disease with Crisis, Unspecified	D57.00
	Sickle - Cell Disease without Crisis	D57.1
	Thalassemia	D56
	Von Willebrand Disease	D68.0
Cardiac and Circulatory System Disorders		
	Supravalvular Aortic Stenosis	Q25.3
	Aortopulmonary Septal Defect	Q21.4
	Atrioventricular Septal Defect	Q21.2
	Congenital Insufficiency of Aortic Valve	Q23.1
	Coarctation of Aorta	Q25.1
	Congenital Malformation of Heart, Unspecified	Q24.9
	Congenital Heart Block	q24.6
	Hypoplastic Left Heart Syndrome	Q23.4
	Hypoplastic Right Heart Syndrome	Q22.6
	Marfan's Syndrome	Q87.4
	PDA - Patent Ductus Arteriosus	Q25.0
	Congenital Pulmonary Valve Stenosis	Q22.1
	SVT - Superventricular Tachycardia	I47.1
	Total Anomalous Pulmonary Venous Connection	Q26.2
	TOF - Tetralogy of Fallot	Q21.3
	Discordant Ventriculoarterial Connection	Q20.3
	Congenital Tricuspid Atresia	Q22.4
	VSD - Ventriculoseptal Defect	Q21.0
	Atrial Septal Defect	Q21.1

	Pre - excitation Syndrome includes WPW - Wolf-Parkinson-White Syndrome	I45.6
	Congenital Malformations of Cardiac Chambers and Connection	Q20
	Congenital Renal Artery Stenosis	Q27.1
	Congenital Malformations of Circulatory System, Unspecified	Q28.9
	Congenital Subaortic Stenosis	Q24.4
	Congenital Malformation of Heart, Unspecified	Q24.9
	Interruption of Aortic Arch	Q25.21
	Other Atresia of Aorta	Q25.29
	Congenital Malformation of Aorta, Unspecified	Q25.40
	Absence and Aplasia of Aorta	Q25.41
	Congenital Aneurysm of Aorta	Q25.43
	Other Congenital Malformations of Aorta	Q25.49
	Situs Inversus	Q89.3
Craniofacial Disorders		
	Apert Syndrome	Q87.0
	Cleft Palate	Q35
	Cleft Hard Palate	Q35.1
	Cleft Soft Palate	Q35.3
	Cleft Hard Palate with Cleft Soft Palate	Q35.5
	Cleft Uvula	Q35.7
	Cleft Palate, Unspecified	Q35.9
	Cleft Lip	Q36
	Cleft Lip, Bilateral	Q36.0
	Cleft Lip, Median	Q36.1
	Cleft Lip, Unilateral	Q36.9
	Cleft Palate with Cleft Lip	Q37
	Other Congenital Malformations of Skull and Face Bones, Cranial Deformity	Q75
	Craniosynostosis	Q75.0
	Congenital Malformation Syndrome Predominantly Affecting Facial Appearance, Includes Goldenhar Syndrome and Pierre Robin	Q87.0
	Treacher-Collins Syndrome	Q75.4
Endocrine Disorders		
	CAH - Congenital Adrenal Hyperplasia	E25.0
	Adrenogenital Disorder Unspecified	E25.9
	Diabetes Mellitus due to Underlying Condition	E08
	Type I Diabetes Mellitus	E10

	Type II Diabetes Mellitus	E11.8
	Type II Diabetes Mellitus with Unspecified Complications	E11.8
	Congenital Hypothyroidism with Diffuse Goiter	E03.0
	Congenital Hypothyroidism without Goiter	E03.1
ENT Disorders		
	Hearing Loss - Conductive and Sensorineural	H90
	Hearing Loss - Other and Unspecified	H91
	Congenital Malformations of Ear Causing impairment of Hearing	Q16
	Congenital Absence of (Ear) Auricle Causing Hearing Loss	Q16.0
	Congenital Malformation of Ear Causing Impairment of Hearing, Unspecified	Q16.9
	Conductive Hearing Loss, Unspecified	H90.2
Eye Disorders		
	Other Specified Congenital Malformations of the Eye	Q15.8
	Aphakia (Aquired)	H27.0
	Aphakia (Congenital)	Q12.3
	Aphakia, Unspecified	H27.00
	Aphakia, Right Eye	H27.01
	Aphakia, Left Eye	H27.02
	Aphakia, Bilateral	H27.03
	Infantile and Juvenile Cataracts	H26.0
	Congenital Cataracts	Q12.0
	Eye Surgery - Enophitalmos due to Trauma or Surgery, Right Eye	H05.421
	Eye Surgery - Enophitalmos due to Trauma or Surgery, Left Eye	H05.422
	Unspecified Glaucoma	H40.9
	Legal Blindness, as Defined in USA	H54.8
	Retinopathy of Prematurity, Unspecified	H35.1
	Retinopathy of Prematurity, Right Eye	H35.101
	Retinopathy of Prematurity, Left Eye	H35.102
	Retinopathy of Prematurity, Bilateral	H35.103
	Retinopathy of Prematurity, Unspecified, Unspecified Eye	H35.109
	Presence of Artificial Eye	Z97.0
Congenital and/or Genetic Disorders		
	2-Methylbutyryl-CoA Dehydrogenase Deficiency Long Chain / Very Long Chain	E71.310
	2-Methylbutyryl-CoA Dehydrogenase Deficiency Medium Chain	E71.311

	2-Methylbutyryl-CoA Dehydrogenase Deficiency Short Chain	E71.312
	Adrenoleukodystrophy	E71.511
	Biotindase Deficiency	D81.810
	CAH - Congenital Adrenal Hyperplasia with Enzyme Deficiency	E25.0
	Adrenogenital Disorder Unspecified	E25.9
	Chromosomal Anomaly	O28.5
	Cystic Fibrosis, Unspecified	E84.9
	Down Syndrome, Unspecified	E90.9
	Galactosemia	E74.21
	Hurler's Syndrome	E76.01
	Hypomelanosis of Ito (Other Disorder of Diminished melanin Formation)	L81.6
	Congenital Ichthyosis, Unspecified	Q80.9
	Marfan Syndrome	Q87.4
	MCAD	E71.311
	MSUD Maple Syrup Urine Disease	E71.0
	Other Unspecified Metabolic Disorders	E88
	PKU - Phenylketonuria, Classical	E70.0
	PPA - Propionic Acidemia	E71.121
	Congenital Malformation Syndrome Predominantly Involving Limbs Includes Rubinstein - Taybi Syndrome	Q87.2
	Sanfilippo Syndrome	E76.22
	Turner's Syndrome	Q96
	TYR I Tyrosinemia Type I	E70.21
	Fetal Alcohol Syndrome	Q86.0
	Congenital Cytomegalovirus Infection	P35.1
	Congenital Toxoplasmosis	P37.1
	Prune Belly Syndrome	Q79.4
	Ehlers Danlos, Unspecified	Q79.60
	Sanfilippo Syndrome	E76.22
	Hurler's Syndrome	E76.01
	Pompe Disease	E74.02
	Other Glycogen Storage Disease	E74.09
	Mitochondrial Metabolism Disorder, Unspecified	E88.40
	Trisomy 18, Unspecified	Q91.3
	Trisomy 13, Unspecified	Q91.7
	Trisomy and Partial Trisomy of Autosomes, Unspecified	Q92.9
	Phakomatoses, Not Elsewhere Classified	Q85.8
	Phakomatoses, Unspecified	Q85.9

	Other Liped Storage Disorder	E75.5
	Other GM2 Gangliosidosis	E75.09
	Other Congenital Malformation Syndrome Predominantly Associated with Short Stature	Q87.19
	VATER Syndrome	C24.1
	Velocardiofacial Syndrome / Digeorge Syndrome	D82.1
	Ichthyosis Vulgaris	Q80.0
	Fragile X	Q99.2
	Hereditary Vitamin D - Dependent Rickets (Type I) (Type II)	E83.32
	Other Specified Chromosome Abnormalities	Q99.8
	Williams Syndrome	Q93.82
	Prader - Willi Syndrome	Q87.11
	Klinefelter's Syndrome, Unspecified	Q98.4
	Marfan's Syndrome, Unspecified	Q87.40
	Polyostic Fibrous Dysplasia - McCune Albright Stemberg Syndrome	Q78.1
	Multiple Congenital Anomalies, Not Elsewhere Classified	Q89.7
	Other Specified Congenital Anomalies	Q89.8
	Mitochondrial Disorder, Unspecified	E88.40
	Metabolic Disorder, Unspecified	E88.9
	Juvenile Dermatoymyositis with Myopathy	M33.02
	Juvenile Dermatoymyositis without Myopathy	M33.03
Rheumatoid Disorders		
	Juvenile Arthritis, Unspecified, Unspecified Site	M08.09
	Rheumatiod Arthritis, Unspecified	M06.9
Genitourinary System		
	Indeterminate Sex, Unspecified - Ambiguous Genitalia	Q56.4
	Renal Agenesis, Unilateral	Q60.0
	Renal Agenesis, Bilateral	Q60.1
	Renal Agenesis, Unspecified	Q60.2
	Renal Hypoplasia, Unilateral	Q60.3
	Renal Hypoplasia, Bilateral	Q60.4
	Renal Hypoplasia, Unspecified	Q60.5
	Potter's Syndrome	Q60.6
	Other Unspecified Congenital Malformations of the Bladder and Urethra	Q64.7
	Hermaphroditism, Not Elsewhere Classified	Q56.0
	Congenital Malformation of Male Genitalia Organ, Unspecified	Q55.9

	Congenital Malformation of Female Genitalia, Unspecified	Q52.9
	Congenital Malformation of Ovaries, Fallopian Tubes and Broad Ligaments, Unilateral	Q50.01
	Congenital Malformation of Ovaries, Fallopian Tubes and Broad Ligaments, Bilateral	Q50.02
	Developmental Ovarian Cyst	Q50.1
	Congenital Torision of Ovary	Q50.2
	Accessory Ovary	Q50.31
	Ovarian Streak	Q50.32
	Other Congenital Malformation of Ovary	Q50.39
	Embryonic Cyst of Fallopian Tube	Q50.4
	Embryonic Cyst of Broad Ligament	Q50.5
	Other Congenital Malformations of Fallopian Tube and Broad Ligament	Q50.6
	Congenital Malformation of Uterus and Cervix, Unspecified	Q51.9
	Undescended Testicle, Unspecified	Q53.9
	Ectopic Testicle, Unspecified	Q53.00
	Hypospadias, Unspecified	Q54.9
	Indeterminate Sex and Pseudohermaphroditism	Q56.4
	Other Specified Congenital Malformation of Intestines	Q43.8
	Neuromuscular Dysfunction of Bladder, Unspecified - Neuogenic Bladder	N31.9
	Renal Agenesis, Unspecified	Q60.2
	Renal Dysplasia / Hypoplasia	Q61.4
	Congenital Hydronephrosis	Q62.0
	Agenesis of Ureter	Q62.4
	Hypospadias, Unspecified	Q54.9
	Polycystic Kidney Infantile Type	Q61.1
Malformation of Other Organ Systems		
	Laryngeal Hypoplasia	Q31.2
	Hirschsprung's Disease	Q43.1
	Congenital Malformation of Intestines, Unspecified	Q43.9
	Neurogenic Bowel, Not Elsewhere Classified	K59.2
	Congenital Malformations of Musculoskeletal System	Q79.8
	Congenital Tracheomalacia	Q32.0
	Other Specified Congenital Malformations of Digestive System	Q45.8
	Gastroschisis	Q79.3

	Congenital Diaphragmatic Hernia	Q79.0
	Congenital Malformations of the Gastointestional Tract, Unspecified	Q40.9
	Other Congenital Malformation Syndromes Due to Known Exogenous Causes	Q86.8
	Congenital Malformation of Respiratory System, Unspecified	Q34.9
	Atresia of Esophagus with Tracheo - Esophageal Fistula	Q39.1
	Congenital Tracheo - Esophageal Fistula without Atresia	Q39.2
Neurological Disorders		
	Brachial Plexus Disorders	G54.0
	Cerebral Palsy, Unspecified	G80.9
	Demyelinating Disorder - Multiple Sclerosis	G35
	Neurofibromatosis	Q85.00
	Other Demyelinating Disease of Central Nervous System, Unspecified	G37.9
	Encephalocele, Unspecified	Q01.9
	Erbs Palsy, Unspecified	G80.9
	Fetal Alcohol Syndrome (Dysmorphic)	Q86.0
	Tuberous Sclerosis	Q85.1
	Hydrocephalus, Unspecified	G91.9
	Phakomatosis, Unspecified - Neurocutaneous Syndrome	Q85.9
	Neurofibromatosis, Unspecified	Q85.00
	Seizures (Epilepsy)	G40
	Spina Bifida Occulta	Q76.0
	Spina Bifida, Unspecified	Q05.9
	Congenital Malformation, Unspecified	Q89.9
	Encephalpathy, Unspecified	G93.40
	Wernick's Encephalpathy	E51.2
	Spastic Diplegic Cerebral Palsy	G80.1
	Spastic Hemiplegia Affecting Unspecified Side	G81.10
	Spastic Quadriplegia Cerebral Palsy	G80.0
	Other Spondylosis with Myelopathy, Lumbar Region	M47.16
	Microcephaly	Q02
	Congenital Hydrocephalus, Unspecified	Q03.9
	Congenital Malformation of Brain, Unspecified	Q04.9
	Other Specified Congenital Malformations of Spinal Cord - Tethered Cord	Q06.8
Orthopedic Disorders / Musculoskeletal		

	Achondroplasia	Q77.4
	Congenital Malformation of Limb(s)	Q74.9
	Unspecified Acquired Deformity of Limb and Hand	M21.9
	Unspecified Acquired Deformity of Unspecified Limb	M21.90
	Other Congenital Malformation of Musculoskeletal System - Amniotic Band Syndrome	Q79.8
	Aquired Absense of Limb, Unspecified	Z89.9
	Congenital Absence of Unspecified Limbs	Q73.0
	Arthrogryposis Multiplex Congenita	Q74.3
	Unspecified Congenital malformation of Limb(s)	Q74.9
	Benign Neoplasm of Bone and Articular Cartilage, Unspecified	D16.9
	Juvenile Osteochondrosis of Tibia and Fibula, Unspecified leg - Blount Disease	M92.50
	Juvenile Osteochondrosis, Unspecified	M92.9
	Solitary Bone Cyst, Unspecified Site	M85.40
	Aneurysmal Bone Cyst	M85.5
	Congenital Deformity of feet, Unspecified, Unspecified Foot - Club Foot / Feet	q66.90
	Congenital Kyphosis, Unspecified Region	Q76.419
	Unspecified Congenital Malformation of Limb(s)	Q74.9
	Syndactyly, Unspecified	Q70.9
	Contracture, Unspecified Joint	M24.50
	Congenital Dislocation of Hip, Unspecified	Q65.2
	Benign Lipomatus Neoplasm of skin and Subcutaneous Tissue of Right Leg	D17.23
	Benign Lipomatus Neoplasm of skin and Subcutaneous Tissue of Left Leg	D17.24
	Unspecified Reduction Defect of Unspecified Lower Limb - Limb Length	Q72.90
	Juvenile Osteochondrosis Legg Calvé Perthes Disease Right Leg	M91.11
	Juvenile Osteochondrosis Legg Calvé Perthes Disease Left Leg	M91.12
	Spinal Osteochondroma, Unspecified	M42.9
	Unspecified Congenital Malformation of Limb(s)	Q74.9
	Osteogenesis Imperfecta	Q78.0
	Rickets, Active	E55.0
	Scoliosis, Unspecified	M41.9

	Congenital Deformity of Spine	Q67.5
	Skeletal Fluorasis, Unspecified Site	M85.10
	Other Osteochondrodysplasia with Defects of Growth Tubular Bones and Spines	Q77.8
	Osteochondrodysplasias, Unspecified	Q78.9
	Osteogenesis Imperfecta	Q78.0
	Pes Planus, Unspecified Foot	Q66.50
	Polysyndactyly, Unspecified	Q70.4
Skin / Subcutaneous / Vascular Tissue Disorders Causing Disrupted Function		
	Congenital Ichthyosis, Unspecified	Q80.9
	Hypomelanosis of Ito (Other Disorder of Diminished melanin Formation)	L81.6
	Hemangiomas, Unspecified Site	D18.00
	Ectodermal Dysplasia (Anhidrotic)	Q82.4
	Congenital Ichthyosis, Unspecified	Q80.9
Developmental / Intellectual Impairment		
	Borderline Intellectual Functioning	R41.83
	Mild Intellectual Disability	F70
	Moderate Intellectual Disability	F71
	Severe Intellectual Disability	F72
	Profound Intellectual Disability	F73
	Intellectual Disability, Unspecified	F79
	Pervasive Developmental Disorder	F84
	Pervasive Developmental Disorder, Unspecified	F84.9
	Autistic Disorder (Autism Spectrum Disorder - ASD, Infantile Autism, Infantile Psychosis, Kanner's Syndrome)	F84.0
	Asperger's Syndrome	F84.5
	Rett's Syndrome	F84.2
	Other Childhood Disintegrative Disorders	F84.3
	Specific Reading Disorder	F81.0
	Mathematics Disorder	F81.2
	Disorder of Written Expression	F81.81
	Other Developmental Disorders of Scholastic Skills	F81.89
	Developmental Disorder of Scholastic Skills, Unspecified	F81.9
	Specific Developmental Disorder of Motor Function	F82
	Other Pervasive Developmenta Disorders	F84.8
	Other Disorders of Psychological Development	F88

	Unspecified Disorder of Psychological Development	F89
	Attention - Deficit Hyperactivity Disorders	F90
	Attention - Deficit Hyperactivity Disorder, Predominantly Inattentive Type	F90.0
	Attention - Deficit Hyperactivity Disorder, Predominantly Hyperactive Type	F90.1
	Attention - Deficit Hyperactivity Disorder, Combination Type	F90.2
	Attention - Deficit Hyperactivity Disorder, Other Type	F90.8
	Attention-Deficit Hyperactivity Disorder, Unspecified Type	F90.9
Behavioral / Mental Health		
	Behavioral Insomnia, Unspecified Type	Z73.819
	Behavioral Insomnia of Childhood, Sleep-Onset Associated Type	Z73.810
	Mental Disorder, Not Otherwise Specified	F99
	Insomnia, Unspecified	G47.00
	Other Insomnia	G47.09
	Hypersomnia, Unspecified	G47.10
	Insomnia Due to Other Mental Disorder	F51.05
	Hypersomnia, Due to other Mental Disorder	F51.13
	Separation Anxiety Disorder of Childhood	F93.0
	Social Phobia, Social Anxiety Disorder	F40.1
	Bipolar Disorder, Unspecified	F31.9
	Schizoaffective Disorder, Unspecified	F25.9
	Schizophrenia, Unspecified	F20.9
Expanded List of Chronic Disorders		
	Unspecified Sever Protein - Calorie Malnutrition	E43
	Obesity, Unspecified	E66.9
	Prediabetes	R73.03
	Type 2 Diabetes Mellitus with Unspecified Complications	E11.8
	Other Asthma	J45.998
	Congenital Zika Virus Disease	P35.4
Prematurity / Low Birthweight		
	Preterm Newborn, Unspecified Weeks of Gestation	P07.30
	Birthweight 2499 Grams or Less	P07.10
	Birthweight - Extreme 999 Grams or less	P07.00
	Birthweight - 499 Grams or less	P07.01
	Birthweight - 500 -750 Grams	P07.02
	Birthweight - 750 - 999 Grams	P07.03

	Birthweight - 1000 -1249 Grams	P07.14
	Birthweight - 1250 - 1499 Grams	P07.15
	Birthweight - 1500 - 1749 Grams	P07.16
	Birthweight - 1750 -1999 Grams	P07.17
	Birthweight - 2000 - 2499 Grams	P07.18
Viral or Congenital / Acquired Infectious Diseases		
	COVID 19	U07.1
	Congenital Zika Virus Disease	P35.4
	Toxoplasmosis, Unspecified	B58.9
	Rubella Encephalitis	B06.01
	Cytomegaloviral Pancreatitis - CMV	B25.2
	Herpesviral Infection, Unspecified	B00.9
	Congenital Syphilis, Unspecified	A50.9
	Human Immunodeficiency Virus Disease	B20
	Congenital Viral Diseases, Unspecified	P35.9
Exposures or Substance Abuse		
	Abnormal Lead Level in Blood	R78.71
	Fetal Hydantoin Syndrome	Q86.1
	Neonatal Withdrawal Symptoms from Maternal Use of Drugs of Addiction (Includes NAS - Neonatal Abstinence Syndrome)	P96.1