Excluded pediatric codes

The following pediatric diagnosis codes are excluded from the physical medicine and therapies component of our Physical Medicine program for members aged 17 and younger. Services are subject to benefit limitations.

Code	Description	Code	Description
E75.24	Niemann-Pick disease	G82.51	Quadriplegia, C1-C4 complete
E75.240	Niemann-Pick disease type A	G91.0	Communicating hydrocephalus
E75.241	Niemann-Pick disease type B	G91.1	Obstructive hydrocephalus
E75.242	Niemann-Pick disease type C	G91.3	Post-traumatic hydrocephalus, unspecified
E75.243	Niemann-Pick disease type D	G91.4	Hydrocephalus in diseases classified elsewhere
E75.248	Other Niemann-Pick disease	G91.8	Other hydrocephalus
E75.249	Niemann-Pick disease, unspecified	G91.9	Hydrocephalus, unspecified
E75.3	Sphingolipidosis, unspecified	G93.1	Anoxic brain damage, not elsewhere classified
E75.5	Other lipid storage disorders	G93.40	Encephalopathy, unspecified
E75.6	Lipid storage disorder, unspecified	G93.5	Compression of brain
E76	Disorders of glycosaminoglycan metabolism	G93.6	Cerebral edema
E76.0	Mucopolysaccharidosis, Type I	G93.7	Reye's syndrome
E76.01	Hurler's syndrome	G93.89	Other specified disorders of brain
E76.02	Hurler-Scheie syndrome	G93.9	Disorder of brain, unspecified
E76.03	Scheie's syndrome	G96.9	Disorder of central nervous system, unspecified
P07.30	Preterm newborn, unspecified weeks of gestation	G98.8	Other disorders of nervous system
P07.31	Preterm newborn, gestational age 28 completed weeks	P07.3	Preterm [premature] newborn [other]
P07.32	Preterm newborn, gestational age 29 completed weeks	P83.2	Hydrops fetalis not due to hemolytic disease
P07.33	Preterm newborn, gestational age 30 completed weeks	Q01.0	Feeding problems of newborn
P07.34	Preterm newborn, gestational age 31 completed weeks	Q01.1	Frontal encephalocele
P07.35	Preterm newborn, gestational age 32	Q01.2	Nasofrontal encephalocele

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	completed weeks		
P07.36	Preterm newborn, gestational age 33 completed weeks	Q01.8	Occipital encephalocele
P07.37	Preterm newborn, gestational age 34 completed weeks	Q01.9	Encephalocele of other sites
P07.38	Preterm newborn, gestational age 35 completed weeks	Q02	Encephalocele, unspecified
P07.39	Preterm newborn, gestational age 36 completed weeks	Q03.0	Microcephaly
Q06	Other congenital malformations of spinal cord	Q03.1	Malformations of aqueduct of Sylvius
Q06.0	Amyelia	Q03.8	Atresia of foramina of Magendie and Luschka
Q06.1	Hypoplasia and dysplasia of spinal cord	Q03.9	Other congenital hydrocephalus
Q06.2	Diastematomyelia	Q04.0	Congenital hydrocephalus, unspecified
Q06.3	Other congenital cauda equina malformations	Q04.1	Arhinencephaly
Q06.4	Hydromyelia	Q04.2	Holoprosencephaly
Q06.8	Other specified congenital malformations of spinal cord	Q04.3	Other reduction deformities of brain
Q92.6	Marker chromosomes	Q04.4	Septo-optic dysplasia of brain
Q93	Monosomies and deletions from the autosomes, not elsewhere classified	Q04.5	Megalencephaly
Q93.51	Angelman syndrome	Q04.6	Congenital cerebral cysts
Q93.59	Other deletions of part of a chromosome	Q04.8	Other specified congenital malformations of brain
Q93.8	Other deletions from the autosomes	Q04.9	Congenital malformation of brain, unspecified
Q93.82	Williams syndrome	Q05.0	Cervical spina bifida with hydrocephalus
D82.1	Di George's syndrome	Q05.1	Thoracic spina bifida with hydrocephalus
E75.0	GM2 gangliosidosis	Q05.2	Lumbar spina bifida with hydrocephalus
E75.00	GM2 gangliosidosis, unspecified	Q05.3	Sacral spina bifida with hydrocephalus

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E75.01	Sandhoff disease	Q05.4	Unspecified spina bifida with hydrocephalus
E75.02	Tay-Sachs disease	Q05.5	Cervical spina bifida without hydrocephalus
E75.09	Other GM2 gangliosidosis	Q05.6	Thoracic spina bifida without hydrocephalus
E75.1	Other and unspecified gangliosidosis	Q05.7	Lumbar spina bifida without hydrocephalus
E75.10	Unspecified gangliosidosis	Q05.8	Sacral spina bifida without hydrocephalus
E75.11	Mucolipidosis IV	Q05.9	Spina bifida, unspecified
E75.19	Other gangliosidosis	Q06.9	Congenital malformation of spinal cord, unspecified
E75.2	Other sphingolipidosis	Q07.00	Arnold-Chiari syndrome without spina bifida or hydrocephalus
E75.21	Fabry (-Anderson) disease	Q07.01	Arnold-Chiari syndrome with spina bifida
E75.22	Gaucher disease	Q07.02	Arnold-Chiari syndrome with hydrocephalus
E75.23	Krabbe disease	Q07.03	Arnold-Chiari syndrome with spina bifida and hydrocephalus
E75.25	Metachromatic leukodystrophy	Q07.8	Other specified congenital malformation of nervous system
E75.26	Sulfatase deficiency	Q07.9	Congenital malformation of nervous system, unspecified
E75.29	Other sphingolipidosis	Q90.0	Trisomy 21, nonmosaicism (meiotic nondisjunction)
E75.4	Neuronal ceroid lipofuscinosis	Q90.1	Trisomy 21, mosaicism (mitotic nondisjunction)
E78.71	Barth syndrome	Q90.2	Trisomy 21, translocation
E78.72	Smith-Lemli-Opitz syndrome	Q90.9	Down syndrome, unspecified
F70	Mild intellectual disabilities	Q91.0	Trisomy 18, nonmosaicism (meiotic nondisjunction)
F71	Moderate intellectual disabilities	Q91.1	Trisomy 18, mosaicism (mitotic nondisjunction)
F72	Severe intellectual disabilities	Q91.2	Trisomy 18, translocation
F73	Profound intellectual disabilities	Q91.3	Trisomy 18, unspecified
F78	Other intellectual disabilities	Q91.4	Trisomy 13, nonmosaicism (meiotic nondisjunction)
F79	Unspecified intellectual disabilities	Q91.5	Trisomy 13, mosaicism (mitotic nondisjunction)
F82	Specific developmental disorder of motor	Q91.6	Trisomy 13, translocation

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	function		
F84	Pervasive development disorders	Q91.7	Trisomy 13, unspecified
F84.0	Autistic disorder	Q92.0	Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)
F84.2	Rett's syndrome	Q92.1	Whole chromosome trisomy, mosaicism (mitotic nondisjunction)
F84.3	Other childhood disintegrative disorder	Q92.2	Partial trisomy
F84.5	Asperger's syndrome	Q92.5	Duplications with other complex rearrangements
F84.8	Other pervasive developmental disorders	Q92.61	Marker chromosomes in normal individual
F84.9	Pervasive developmental disorder, unspecified	Q92.62	Marker chromosomes in abnormal individual
F88	Other disorders of psychological development	Q92.7	Triploidy and polyploidy
F89	Unspecified disorder of psychological development	Q92.8	Other specified trisomies and partial trisomies of autosomes
F90	Attention-deficit hyperactivity disorders	Q92.9	Trisomy and partial trisomy of autosomes, unspecified
F98.2	Other feeding disorders of infancy and childhood	Q93.0	Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)
F98.9	Unspecified behavioral and emotional disorders with onset usually occurring in childhood and adolescence	Q93.1	Whole chromosome monosomy, mosaicism (mitotic nondisjunction)
G11.1	Early-onset cerebellar ataxia	Q93.2	Chromosome replaced with ring, dicentric or isochromosome
G12.0	Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]	Q93.3	Deletion of short arm of chromosome 4
G12.1	Other inherited spinal muscular atrophy	Q93.4	Deletion of short arm of chromosome 5
G31.84	Mild cognitive impairment, so stated	Q93.5	Other deletions of part of a chromosome
G71.0	Muscular Dystrophy	Q93.7	Deletions with other complex rearrangements
G71.00	Muscular dystrophy, unspecified	Q93.81	Velo-cardio-facial syndrome

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G71.01	Duchenne or Becker muscular dystrophy	Q93.88	Other microdeletions
G71.02	Facioscapulohumeral muscular dystrophy	Q93.89	Other deletions from the autosomes
G71.09	Other specified muscular dystrophies	Q93.9	Deletion from autosomes, unspecified
G71.11	Myotonic muscular dystrophy	Q95.2	Balanced autosomal rearrangement in abnormal individual
G71.12	Myotonia congenita	Q95.3	Balanced sex/autosomal rearrangement in abnormal individual
G71.13	Myotonic chondrodystrophy	Q99.2	Fragile X chromosome
G71.14	Drug induced myotonia	Q99.8	Other specified chromosome abnormalities
G71.19	Other specified myotonic disorders	Q99.9	Chromosomal abnormality, unspecified
G71.2	Congenital myopathies	R27.9	Unspecified lack of coordination
G80.0	Spastic quadriplegic cerebral palsy	R62.0	Delayed milestone in childhood
G80.1	Spastic diplegic cerebral palsy	R62.50	Unspecified lack of expected normal physiological development in childhood
G80.2	Spastic hemiplegic cerebral palsy	R62.51	Failure to thrive (child)
G80.3	Athetoid cerebral palsy	R62.59	Other lack of expected normal physiological development in childhood
G80.4	Ataxic cerebral palsy	R63.3	Feeding difficulties
G80.8	Other cerebral palsy	T74.4XXA	Shaken infant syndrome, initial encounter
G80.9	Cerebral palsy, unspecified	T74.4XXD	Shaken infant syndrome, subsequent encounter
		T74.4XXS	Shaken infant syndrome, sequela