



Medical Necessity Guidelines:

Covered Diagnosis Code Reference Tool for Outpatient Habilitative Services: Physical Therapy, Occupational Therapy, and Speech Therapy

Effective: December 1, 2024

Prior Authorization Required If REQUIRED, submit supporting clinical documentation pertinent to service request to the FAX numbers below	Yes ⊠ No □
Notification Required IF REQUIRED, concurrent review may apply	Yes □ No ⊠
Applies to: Commercial Products	
 □ Harvard Pilgrim Health Care Commercial products; 800-232-0816 ☑ Tufts Health Plan Commercial products; 617-972-9409 CareLinkSM – Refer to CareLink Procedures, Services and Items Requiring Prior Authorization 	
Public Plans Products	
 ☑ Tufts Health Direct – A Massachusetts Qualified Health Plan (QHP) (a commercial product); 888-415-5 ☐ Tufts Health Together – MassHealth MCO Plan and Accountable Care Partnership Plans; 888-415-5 ☐ Tufts Health RITogether – A Rhode Island Medicaid Plan; 857-304-6404 ☐ Tufts Health One Care – A dual-eligible product; 857-304-6304 	
Senior Products	
 □ Harvard Pilgrim Health Care Stride Medicare Advantage; 888-609-0692 □ Tufts Health Plan Senior Care Options (SCO), (a dual-eligible product); 617-673-0965 □ Tufts Medicare Preferred HMO, (a Medicare Advantage product); 617-673-0965 □ Tufts Medicare Preferred PPO, (a Medicare Advantage product); 617-673-0965 	

Note: While you may not be the provider responsible for obtaining prior authorization or notifying Point32Health, as a condition of payment you will need to ensure that any necessary prior authorization has been obtained and/or Point32Health has received proper notification. If notification is required, providers may additionally be required to provide updated clinical information to qualify for continued service.

Overview

Habilitative services are provided for a person to attain, maintain, or prevent deterioration of a skill or function never learned or acquired due to a disabling condition. Habilitative Services are provided per the Member's plan benefit.

Please refer to the Medical Necessity Guidelines for Outpatient Habilitative Services: Physical Therapy, Occupational Therapy and Speech Therapy for additional information regarding coverage.

Codes

The following code(s) require prior authorization:

Table 1: ICD-10-CM Habilitative Codes covered if selection criteria are met

*The following contains diagnosis codes that are considered experimental in nature

Codes	Description
DEVELOPMENT	•
F80.0	Phonological disorder
F80.1	Expressive language disorder
F80.2	Mixed receptive-expressive language disorder
F80.89	Other developmental disorders of speech and language
F80.9	Developmental disorders of speech and language, unspecified
F82	Specific developmental disorder of motor function
F88	Other disorders of psychological development (developmental agnosia)
F89	Unspecified disorder of psychological development
H93.25	Central auditory processing disorder
R62.0	Delayed milestone in childhood
R62.50	Unspecified lack of expected normal physiological development in childhood
R62.52	Short stature (child) (lack of growth, physical retardation)
R62.59	Other lack of expected normal physiological development in childhood
	SEASES/DISORDERS
A50.01	Early congenital syphilitic oculopathy
A50.02	Early congenital syphilitic osteochondropathy
A50.02	Early congenital syphilitic discontinuopatry Early congenital syphilitic pharyngitis
A50.03	Early congenital syphilitic pneumonia
A50.04	Early congenital syphilitic pheumonia Early congenital syphilitic rhinitis
A50.05	Early cutaneous congenital syphilis
A50.07	Early mucocutaneous congenital syphilis
A50.07	Early visceral congenital syphilis
A50.08	Other early congenital syphilis, symptomatic
A50.09	Early congenital syphilis, latent (less than two years after birth)
A50.1	Early congenital syphilis, inspecified
A50.2 A50.30	Late congenital syphilitic oculopathy, unspecified
A50.30	Late congenital syphilitic interstitial keratitis
A50.32	Late congenital syphilitic chorioretinitis
A50.32	Other late congenital syphilitic oculopathy
F80.4	Speech and language development delay due to hearing loss
F80.81	Childhood onset fluency disorder
F80.82	Social pragmatic communication disorder
F84.2	Rett's syndrome
F90.0	Attention-deficit hyperactivity disorder, predominantly inattentive type
F90.0	Attention-deficit hyperactivity disorder, predominantly inattentive type Attention-deficit hyperactivity disorder, predominantly hyperactive type
F90.2	Attention-deficit hyperactivity disorder, predominantly hyperactive type Attention-deficit hyperactivity disorder, combined type
F90.8	Attention-deficit hyperactivity disorder, combined type Attention-deficit hyperactivity disorder, other type
F90.9	Attention-deficit hyperactivity disorder, other type Attention-deficit hyperactivity disorder, unspecified type
F98.2	Other feeding disorders of infancy and childhood
F98.21	Rumination disorder of infancy
F98.29	Other feeding disorders of infancy and early childhood
F98.4 G11.0	Stereotyped movement disorders Congenital nonprogressive ataxia
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G11.10	Early-onset cerebellar ataxia
	Early-onset cerebellar ataxia, unspecified
G11.11	Priedreich ataxia
G11.19	Other early-onset cerebellar ataxia
G11.3	Cerebellar ataxia with defective DNA repair
G11.4	Hereditary spastic paraplegia
G11.8	Other hereditary ataxias

Codes	Description
G11.9	Hereditary ataxia, unspecified
G12.0	Infantile spinal muscular atrophy, type I [Werdnig-Hoffman]
G12.0	Other inherited spinal muscular atrophy
G12.15	Progressive spinal muscle atrophy
G24.1	Genetic torsion dystonia
G24.1	Idiopathic nonfamilial dystonia
G24.2 G40.42	Cyclin-Dependent Kinase-Like 5 Deficiency Disorder
G40.833	
G40.834	Dravet syndrome, intractable, with status epilepticus
G40.834 G70.2	Dravet syndrome, intractable, without status epilepticus
G70.2 G71.0	Congenital and developmental myasthenia
G71.00	Muscular dystrophy
	Muscular dystrophy, unspecified
G71.01 G71.02	Duchenne or Becker muscular dystrophy
	Facioscapulohumeral muscular dystrophy Other appointed muscular dystrophics
G71.09 G71.11	Other specified muscular dystrophies
	Myotonic muscular dystrophy
G71.12	Myotonia congenita
G71.13	Myotonic chondrodystrophy
G71.2	Congenital myopathies
G71.20	Congenital myopathy, unspecified
G71.21	Nemaline myopathy
G71.220	X-linked myotubular myopathy
G71.228	Other centronuclear myopathy
G71.29	Other congenital myopathy
G71.3	Mitochondrial myopathy, not elsewhere classified
G80.0	Spastic quadriplegic cerebral palsy
G80.1	Spastic diplegic cerebral palsy
G80.2	Spastic hemiplegic cerebral palsy
G80.3	Athetoid cerebral palsy
G80.4	Ataxic cerebral palsy
G80.8	Other cerebral palsy
G80.9	Cerebral palsy, unspecified
H55.01	Congenital nystagmus
M61.10	Myositis ossificans progressiva, unspecified site
M61.111	Myositis ossificans progressiva, right shoulder
M61.112	Myositis ossificans progressiva, left shoulder
M61.119	Myositis ossificans progressiva, unspecified shoulder
M61.121	Myositis ossificans progressiva, right upper arm
M61.122	Myositis ossificans progressiva, left upper arm
M61.129	Myositis ossificans progressiva, unspecified arm
M61.131	Myositis ossificans progressiva, right forearm
M61.132	Myositis ossificans progressiva, left forearm
M61.139	Myositis ossificans progressiva, unspecified forearm
M61.141	Myositis ossificans progressiva, right hand
M61.142	Myositis ossificans progressiva, left hand
M61.143	Myositis ossificans progressiva, unspecified hand
M61.144	Myositis ossificans progressiva, right finger(s)
M61.145	Myositis ossificans progressiva, left finger(s)
M61.146	Myositis ossificans progressiva, unspecified finger(s)
M61.151	Myositis ossificans progressiva, right thigh
M61.152	Myositis ossificans progressiva, left thigh

Codes	Description
M61.159	Myositis ossificans progressiva, unspecified thigh
M61.161	Myositis ossificans progressiva, unspecified triight Myositis ossificans progressiva, right lower leg
M61.162	Myositis ossificans progressiva, light lower leg
M61.169	Myositis ossificans progressiva, lett lower leg Myositis ossificans progressiva, unspecified lower leg
M61.171	Myositis ossificans progressiva, drispectified lower leg Myositis ossificans progressiva, right ankle
M61.172	Myositis ossificans progressiva, light ankle Myositis ossificans progressiva, left ankle
M61.173	Myositis ossificans progressiva, lett arikle Myositis ossificans progressiva, unspecified ankle
M61.174	Myositis ossificans progressiva, unspecified arrive
M61.175	Myositis ossificans progressiva, light foot
M61.176	Myositis ossificans progressiva, lett loot Myositis ossificans progressiva, unspecified foot
M61.177	Myositis ossificans progressiva, drispectified foot Myositis ossificans progressiva, right toe(s)
M61.178	Myositis ossificans progressiva, light toe(s) Myositis ossificans progressiva, left toe(s)
M61.179	Myositis ossificans progressiva, left toe(s) Myositis ossificans progressiva, unspecified toe(s)
M61.18	Myositis ossificans progressiva, unspecified toe(s) Myositis ossificans progressiva, other site
M61.19	
P91.821	Myositis ossificans progressiva, multiple sites Neonatal cerebral infarction, right side of brain
P91.822	· G
P91.822	Neonatal cerebral infarction, left side of brain
P91.823 P94.1	Neonatal cerebral infarction, bilateral
	Congenital hypertonia
P94.2	Congenital hypotonia
P94.8	Other disorders of muscle tone of newborn
P94.9	Disorder of muscle tone of newborn, unspecified
Q00.0	Anencephaly Craniorachischisis
Q00.1	
Q00.2	Iniencephaly
Q01.0	Frontal encephalocele
Q01.1	Nasofrontal encephalocele
Q01.2	Occipital encephalocele
Q01.8	Encephalocele of other sites
Q01.9	Encephalocele, unspecified
Q02	Microcephaly Malfarranting and a read wat of Cultification
Q03.0	Malformations of aqueduct of Sylvius
Q03.1	Atresia of foramina of Magendie and Luschka
Q03.8	Other congenital hydrocephalus
Q03.9	Congenital hydrocephalus, unspecified
Q04.0	Congenital malformations of corpus callosum
Q04.1	Arhinencephaly
Q04.2	Holoprosencephaly
Q04.3	Other reduction deformities of brain
Q04.4	Septo-optic dysplasia of brain
Q04.5	Megalencephaly Congressite Lorentz Burgets
Q04.6	Congenital cerebral cysts
Q04.8	Other specified congenital malformations of brain
Q04.9	Congenital malformation of brain, unspecified
Q05.0	Cervical spina bifida with hydrocephalus
Q05.1	Thoracic spina bifida with hydrocephalus
Q05.2	Lumbar spina bifida with hydrocephalus
Q05.3	Sacral spina bifida with hydrocephalus
Q05.4	Unspecified spina bifida with hydrocephalus
Q05.5	Cervical spina bifida without hydrocephalus
Q05.6	Thoracic spina bifida without hydrocephalus

OS.7 Lumbar spina bifida without hydrocephalus OS.8 Sacral spina bifida, without hydrocephalus Spina bifida, unspecified OS.0 Amyelia OS.1 Hypoplasia and dysplasia of spinal cord OS.2 Diastematomyelia OS.3 Other congenital cauda equina malformations OS.4 Hydromyelia OS.5 Other specified congenital malformations of spinal cord OS.6 Other specified congenital malformations of spinal cord OS.7 Other specified congenital malformations of phydrocephalus OS.7 Other specified congenital malformations of nervous system OS.7 Other specified congenital malformations of spelid OT.0 Congenital entropion OT.0 Congenital entropion OT.0 Other congenital malformations of eyelid OT.0 Congenital entropion OT.0 Other congenital malformations of actimal apparatus OT.0 Congenital stenosis and stricture of lacrimal duct OT.0 Congenital stenosis and stricture of lacrimal apparatus OT.0 Congenital stenosis and stricture of lacrimal apparatus OT.1 Congenital malformations of actimal apparatus OT.1 Congenital malformations of actimal apparatus OT.1 Congenital malformation of orbit OT.1 Congenital malformation of orbit OT.1 Congenital malformations OT.2 Congenital malformations OT.3 Congenital malformations OT.3 Congenital pahakia OT.2 Congenital malformation of orbit OT.3 Congenital malformations OT.3 Congenital malformation of orbit OT.3 Congenital malformation of orbit OT.3 Congenital malformation	Codes	Description
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Q14.9 Congenital malformation of posterior segment of eye, unspecified		

Codes	Description
Q15.8	Other specified congenital malformations of eye
Q15.8 Q15.9	Congenital malformation of eye, unspecified
Q16.0	Congenital absence of (ear) auricle
Q16.0	
	Congenital absence, atresia and stricture of auditory canal (external) Absence of eustachian tube
Q16.2	
Q16.3	Congenital malformation of ear ossicles
Q16.4	Other congenital malformations of middle ear
Q16.5	Congenital malformation of inner ear
Q16.9	Congenital malformation of ear causing impairment of hearing, unspecified
Q17.0	Accessory auricle
Q17.1	Macrotia
Q17.2	Microtia
Q17.3	Other misshapen ear
Q17.4	Misplaced ear
Q17.5	Prominent ear
Q17.8	Other specified congenital malformations of ear
Q17.9	Congenital malformation of ear, unspecified
Q18.0	Sinus, fistula and cyst of branchial cleft
Q18.1	Preauricular sinus and cyst
Q18.2	Other branchial cleft malformations
Q18.3	Webbing of neck
Q18.4	Macrostomia
Q18.5	Microstomia
Q18.6	Macrocheilia
Q18.7	Microcheilia
Q18.8	Other specified congenital malformations of face and neck
Q18.9	Congenital malformation of face and neck, unspecified
Q20.0	Common arterial trunk
Q20.1	Double outlet right ventricle
Q20.2	Double outlet left ventricle
Q20.3	Discordant ventriculoarterial connection
Q20.4	Double inlet ventricle
Q20.5	Discordant atrioventricular connection
Q20.6	Isomerism of atrial appendages
Q20.8	Other congenital malformations of cardiac chambers and connections
Q20.9	Congenital malformation of cardiac chambers and connections, unspecified
Q21.0	Ventricular septal defect
Q21.10	Atrial septal defect, unspecified
Q21.11	Secundum atrial septal defect
Q21.12	Patent foramen ovale
Q21.13	Coronary sinus atrial septal defect
Q21.14	Superior sinus venosus atrial septal defect
Q21.15	Inferior sinus venosus atrial septal defect
Q21.16	Sinus venosus atrial septal defect, unspecifie
Q21.19	Other specified atrial septal defect
Q21.20	Atrioventricular septal defect, unspecified as to partial or complete
Q21.21	Partial atrioventricular septal defect
Q21.22	Transitional atrioventricular septal defect
Q21.23	Complete atrioventricular septal defect
Q21.3	Tetralogy of Fallot
Q21.4	Aortopulmonary septal defect
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Codes	Description
Q21.8	Other congenital malformations of cardiac septa
Q21.8 Q21.9	Congenital malformations of cardiac septum, unspecified
Q21.9 Q22.0	Pulmonary valve atresia
Q22.0 Q22.1	,
	Congenital pulmonary valve stenosis
Q22.2	Congenital pulmonary valve insufficiency
Q22.3	Other congenital malformations of pulmonary valve
Q22.4	Congenital tricuspid stenosis
Q22.5	Ebstein's anomaly
Q22.6	Hypoplastic right heart syndrome
Q22.8	Other congenital malformations of tricuspid valve
Q22.9	Congenital malformation of tricuspid valve, unspecified
Q23.0	Congenital stenosis of aortic valve
Q23.1	Congenital insufficiency of aortic valve
Q23.2	Congenital mitral stenosis
Q23.3	Congenital mitral insufficiency
Q23.4	Hypoplastic left heart syndrome
Q23.8	Other congenital malformations of aortic and mitral valves
Q23.9	Congenital malformation of aortic and mitral valves, unspecified
Q24.0	Dextrocardia
Q24.1	Levocardia
Q24.2	Cor triatriatum
Q24.3	Pulmonary infundibular stenosis
Q24.4	Congenital subaortic stenosis
Q24.5	Malformation of coronary vessels
Q24.6	Congenital heart block
Q24.8	Other specified congenital malformations of heart
Q24.9	Congenital malformation of heart, unspecified
Q25.0	Patent ductus arteriosus
Q25.1	Coarctation of aorta
Q25.2	Atresia of aorta
Q25.21	Interruption of aortic arch
Q25.29	Other atresia of aorta
Q25.3	Supravalvular aortic stenosis
Q25.4	Other congenital malformations of aorta
Q25.40	Congenital malformation of aorta unspecified
Q25.41	Absence and aplasia of aorta
Q25.42	Hypoplasia of aorta
Q25.43	Congenital aneurysm of aorta
Q25.44	Congenital dilation of aorta
Q25.45	Double aortic arch
Q25.46	Tortuous aortic arch
Q25.47	Right aortic arch
Q25.48	Anomalous origin of subclavian artery
Q25.49	Other congenital malformations of aorta
Q25.5	Atresia of pulmonary artery
Q25.6	Stenosis of pulmonary artery
Q25.71	Coarctation of pulmonary artery
Q25.72	Congenital pulmonary arteriovenous malformation
Q25.79	Other congenital malformations of pulmonary artery
Q25.8	Other congenital malformations of other great arteries
Q25.9	Congenital malformation of great arteries, unspecified
	0

Codes	Description
Q26.0	Congenital stenosis of vena cava
Q26.1	Persistent left superior vena cava
Q26.2	Total anomalous pulmonary venous connection
Q26.3	Partial anomalous pulmonary venous connection
Q26.4	Anomalous pulmonary venous connection, unspecified
Q26.5	Anomalous portal venous connection
Q26.6	Portal vein-hepatic artery fistula
Q26.8	Other congenital malformations of great veins
Q26.9	Congenital malformation of great vein, unspecified
Q27.0	Congenital absence and hypoplasia of umbilical artery
Q27.1	Congenital renal artery stenosis
Q27.2	Other congenital malformations of renal artery
Q27.30	Arteriovenous malformation, site unspecified
Q27.31	Arteriovenous malformation, site unspecified Arteriovenous malformation of vessel of upper limb
Q27.32	Arteriovenous malformation of vessel of dyper limb
Q27.33	Arteriovenous malformation of vessel of lower limb Arteriovenous malformation of digestive system vessel
Q27.34	Arteriovenous malformation of digestive system vessel Arteriovenous malformation of renal vessel
Q27.39	Arteriovenous malformation, other site
Q27.4	Congenital phlebectasia
Q27.8	Other specified congenital malformations of peripheral vascular system
Q27.9	Congenital malformation of peripheral vascular system, unspecified
Q28.0	Arteriovenous malformation of precerebral vessels
Q28.1	Other malformations of precerebral vessels
Q28.2	Arteriovenous malformation of cerebral vessels
Q28.3	Other malformations of cerebral vessels
Q28.8	Other specified congenital malformations of circulatory system
Q28.9	Congenital malformation of circulatory system, unspecified
Q30.0	Choanal atresia
Q30.1	Agenesis and underdevelopment of nose
Q30.2	Fissured, notched and cleft nose
Q30.3	Congenital perforated nasal septum
Q30.8	Other congenital malformations of nose
Q30.9	Congenital malformation of nose, unspecified
Q31.0	Web of larynx
Q31.1	Congenital subglottic stenosis
Q31.2	Laryngeal hypoplasia
Q31.3	Laryngocele
Q31.5	Congenital laryngomalacia
Q31.8	Other congenital malformations of larynx
Q31.9	Congenital malformation of larynx, unspecified
Q32.0	Congenital tracheomalacia
Q32.1	Other congenital malformations of trachea
Q32.2	Congenital bronchomalacia
Q32.3	Congenital stenosis of bronchus
Q32.4	Other congenital malformations of bronchus
Q33.0	Congenital cystic lung
Q33.1	Accessory lobe of lung
Q33.2	Sequestration of lung
Q33.3	Agenesis of lung
Q33.4	Congenital bronchiectasis
Q33.5	Ectopic tissue in lung

Congenital hypoplasia and dysplasia of lung Casa.8 Other congenital malformations of lung Casa.8 Other congenital malformations of lung Casa.8 Other congenital malformations of lung Casa.4 Congenital cyst of mediastinum Casa.4 Congenital cyst of mediastinum Casa.4 Congenital cyst of mediastinum Casa.4 Congenital malformation of respiratory system Casa.4 Congenital malformation of respiratory system Casa.5 Cleft soft palate Casa.5 Cleft hard palate Casa.5 Cleft hard palate Casa.5 Cleft hard palate with cleft soft palate Casa.5 Cleft palate, unspecified Casa.6 Cleft lip, bitarial Casa.9 Cleft palate, unspecified Casa.6 Cleft lip, median Casa.9 Cleft lip, median Casa.9 Cleft lip, median Casa.9 Cleft lip, unilateral Casa.9 Cleft lip, unilateral Casa.9 Cleft lip, unilateral Casa.9 Cleft lip, median Casa.9 Cleft soft palate with bilateral cleft lip Casa.1 Cleft hard palate with bilateral cleft lip Casa.2 Cleft soft palate with bilateral cleft lip Casa.2 Cleft soft palate with bilateral cleft lip Casa.3 Cleft soft palate with bilateral cleft lip Casa.4 Cleft hard and soft palate with bilateral cleft lip Casa.5 Cleft hard and soft palate with bilateral cleft lip Casa.6 Cleft hard and soft palate with bilateral cleft lip Casa.6 Cleft hard and soft palate with bilateral cleft lip Casa.6 Congenital malformations of lips, not elsewhere classified Casa.6 Ankyloglossia Casa.6 Macroglossia Casa.7 Congenital malformations of palate, not elsewhere classified Casa.6 Charcongenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.6 Cher congenital malformations of palate, not elsewhere classified Casa.7 Congenital palatery cherophic palate, not elsewhere classified	Codes	Description
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Q40.9 Congenital malformation of upper alimentary tract, unspecified Q41.0 Congenital absence, atresia and stenosis of duodenum Q41.1 Congenital absence, atresia and stenosis of jejunum Q41.2 Congenital absence, atresia and stenosis of ileum	Q40.8	
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Q41.2 Congenital absence, atresia and stenosis of ileum		
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Codes	Description
Q41.9	Congenital absence, atresia and stenosis of small intestine, part unspecified
Q42.0	Congenital absence, atresia and stenosis of rectum with fistula
Q42.0 Q42.1	Congenital absence, atresia and stenosis of rectum with listula Congenital absence, atresia and stenosis of rectum without fistula
Q42.1 Q42.2	Congenital absence, atresia and stenosis of rectum without ristula Congenital absence, atresia and stenosis of anus with fistula
Q42.2 Q42.3	
	Congenital absence, atresia and stenosis of anus without fistula
Q42.8	Congenital absence, atresia and stenosis of other parts of large intestine
Q42.9	Congenital absence, atresia and stenosis of large intestine, part unspecified
Q43.0	Meckel's diverticulum (displaced) (hypertrophic)
Q43.1	Hirschsprung's disease
Q43.2	Other congenital functional disorders of colon
Q43.3	Congenital malformations of intestinal fixation
Q43.4	Duplication of intestine
Q43.5	Ectopic anus
Q43.6	Congenital fistula of rectum and anus
Q43.7	Persistent cloaca
Q43.8	Other specified congenital malformations of intestine
Q43.9	Congenital malformation of intestine, unspecified
Q44.0	Agenesis, aplasia and hypoplasia of gallbladder
Q44.1	Other congenital malformations of gallbladder
Q44.2	Atresia of bile ducts
Q44.3	Congenital stenosis and stricture of bile ducts
Q44.4	Choledochal cyst
Q44.5	Other congenital malformations of bile ducts
Q44.6	Cystic disease of liver
Q44.7	Other congenital malformations of liver
Q45.0	Agenesis, aplasia and hypoplasia of pancreas
Q45.1	Annular pancreas
Q45.2	Congenital pancreatic cyst
Q45.3	Other congenital malformations of pancreas and pancreatic duct
Q45.8	Other specified congenital malformations of digestive system
Q45.9	Congenital malformation of digestive system, unspecified
Q50.01	Congenital absence of ovary, unilateral
Q50.02	Congenital absence of ovary, bilateral
Q50.1	Developmental ovarian cyst
Q50.2	Congenital torsion of ovary
Q50.31	Accessory ovary
Q50.32	Ovarian streak
Q50.39	Other congenital malformation of ovary
Q50.4	Embryonic cyst of fallopian tube
Q50.5	Embryonic cyst of broad ligament
Q50.6	Other congenital malformations of fallopian tube and broad ligament
Q51.0	Agenesis and aplasia of uterus
Q51.10	Doubling of uterus with doubling of cervix and vagina without obstruction
Q51.11	Doubling of uterus with doubling of cervix and vagina with obstruction
Q51.2	Other doubling of uterus
Q51.21	Other complete doubling of uterus
Q51.22	Other partial doubling of uterus
Q51.28	Other doubling of uterus, other specified
Q51.3	Bicornate uterus
Q51.4	Unicornate uterus
Q51.5	Agenesis and aplasia of cervix
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Codes	Description
Q51.6	Embryonic cyst of cervix
Q51.7	Congenital fistulae between uterus and digestive and urinary tracts
Q51.810	Arcuate uterus
Q51.811	Hypoplasia of uterus
Q51.818	Other congenital malformations of uterus
Q51.820	Cervical duplication
Q51.821	Hypoplasia of cervix
Q51.828	Other congenital malformations of cervix
Q51.9	Congenital malformation of uterus and cervix, unspecified
Q52.0	Congenital absence of vagina
Q52.10	Doubling of vagina, unspecified
Q52.11	Transverse vaginal septum
Q52.12	Longitudinal vaginal septum
Q52.120	Longitudinal vaginal septum, nonobstructing
Q52.121	Longitudinal vaginal septum, obstructing, right side
Q52.122	Longitudinal vaginal septum, obstructing, left side
Q52.123	Longitudinal vaginal septum, microperforate, right side
Q52.123	Longitudinal vaginal septum, microperforate, left side
Q52.124 Q52.129	Other and unspecified longitudinal vaginal septum
Q52.129	Congenital rectovaginal fistula
Q52.2 Q52.3	Imperforate hymen
Q52.4	Other congenital malformations of vagina
Q52.5	Fusion of labia
Q52.6	Congenital malformation of clitoris
Q52.70	Unspecified congenital malformations of vulva
Q52.71	Congenital absence of vulva
Q52.71	Other congenital malformations of vulva
Q52.79	Other specified congenital malformations of female genitalia
Q52.9	Congenital malformation of female genitalia, unspecified
Q53.00	Ectopic testis, unspecified
Q53.00	Ectopic testis, unispecified Ectopic testis, unilateral
Q53.01	Ectopic testis, unitateral Ectopic testes, bilateral
Q53.02 Q53.10	Unspecified undescended testicle, unilateral
Q53.10	Unilateral intraabdominal testis
Q53.111 Q53.112	Unilateral inguinal testis
Q53.112 Q53.12	Ectopic perineal testis, unilateral
Q53.12 Q53.13	Unilateral high scrotal testis
Q53.13	Undescended testicle, unspecified, bilateral
Q53.20 Q53.211	Bilateral intraabdominal testes
Q53.211 Q53.212	Bilateral inguinal testes
Q53.212 Q53.22	Ectopic perineal testis, bilateral
Q53.22 Q53.23	Bilateral high scrotal testes
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Q53.9	Undescended testicle, unspecified
Q54.0	Hypospadias, balanic
Q54.1	Hypospadias, penile
Q54.2	Hypospadias, peripad
Q54.3	Hypospadias, perineal
Q54.4	Congenital chordee
Q54.8	Other hypospadias
Q54.9	Hypospadias, unspecified
Q55.0	Absence and aplasia of testis

Codes	Description
Q55.1	Hypoplasia of testis and scrotum
Q55.20	Unspecified congenital malformations of testis and scrotum
Q55.21	Polyorchism
Q55.22	Retractile testis
Q55.23	Scrotal transposition
Q55.29	Other congenital malformations of testis and scrotum
Q55.3	Atresia of vas deferens
Q55.4	Other congenital malformations of vas deferens, epididymis, seminal vesicles and
Q55.5	prostate Congenital absence and aplasia of penis
Q55.61	Curvature of penis (lateral)
Q55.62	Hypoplasia of penis
Q55.63	Congenital torsion of penis
Q55.64	Hidden penis
Q55.69	· ·
Q55.7	Other congenital malformation of penis Congenital vasocutaneous fistula
Q55.8	Other specified congenital malformations of male genital organs
Q55.9	Congenital malformation of male genital organ, unspecified
Q56.0	Hermaphroditism, not elsewhere classified
Q56.1	Male pseudohermaphroditism, not elsewhere classified
Q56.2	Female pseudohermaphroditism, not elsewhere classified
Q56.3	Pseudohermaphroditism, unspecified
Q56.4	Indeterminate sex, unspecified
Q60.0	Renal agenesis, unilateral
Q60.1	Renal agenesis, bilateral
Q60.2	Renal agenesis, unspecified
Q60.3	Renal hypoplasia, unilateral
Q60.4	Renal hypoplasia, bilateral
Q60.5	Renal hypoplasia, unspecified
Q60.6	Potter's syndrome
Q61.00	Congenital renal cyst, unspecified
Q61.01	Congenital single renal cyst
Q61.02	Congenital multiple renal cysts
Q61.11	Cystic dilatation of collecting ducts
Q61.19	Other polycystic kidney, infantile type
Q61.2	Polycystic kidney, adult type
Q61.3	Polycystic kidney, unspecified
Q61.4	Renal dysplasia
Q61.5	Medullary cystic kidney
Q61.8	Other cystic kidney diseases
Q61.9	Cystic kidney disease, unspecified
Q62.0	Congenital hydronephrosis
Q62.10	Congenital occlusion of ureter, unspecified
Q62.11	Congenital occlusion of ureteropelvic junction
Q62.12	Congenital occlusion of ureterovesical orifice
Q62.2	Congenital megaureter
Q62.31	Congenital ureterocele, orthotopic
Q62.32	Cecoureterocele
Q62.39	Other obstructive defects of renal pelvis and ureter
Q62.4	Agenesis of ureter
Q62.5	Duplication of ureter
Q62.60	Malposition of ureter, unspecified

Codes	Description
Q62.61	Deviation of ureter
Q62.62	Displacement of ureter
Q62.63	Anomalous implantation of ureter
Q62.69	Other malposition of ureter
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Q62.7	Congenital vesico-uretero-renal reflux
Q62.8	Other congenital malformations of ureter
Q63.0	Accessory kidney
Q63.1	Lobulated, fused and horseshoe kidney
Q63.2	Ectopic kidney
Q63.3	Hyperplastic and giant kidney
Q63.8	Other specified congenital malformations of kidney
Q63.9	Congenital malformation of kidney, unspecified
Q64.0	Epispadias
Q64.10	Exstrophy of urinary bladder, unspecified
Q64.11	Supravesical fissure of urinary bladder
Q64.12	Cloacal exstrophy of urinary bladder
Q64.19	Other exstrophy of urinary bladder
Q64.2	Congenital posterior urethral valves
Q64.31	Congenital bladder neck obstruction
Q64.32	Congenital stricture of urethra
Q64.33	Congenital stricture of urinary meatus
Q64.39	Other atresia and stenosis of urethra and bladder neck
Q64.4	Malformation of urachus
Q64.5	Congenital absence of bladder and urethra
Q64.6	Congenital diverticulum of bladder
Q64.70	Unspecified congenital malformation of bladder and urethra
Q64.71	Congenital prolapse of urethra
Q64.72	Congenital prolapse of urinary meatus
Q64.73	Congenital urethrorectal fistula
Q64.74	Double urethra
Q64.75	Double urinary meatus
Q64.79	Other congenital malformations of bladder and urethra
Q64.8	Other specified congenital malformations of urinary system
Q64.9	Congenital malformation of urinary system, unspecified
Q65.01	Congenital dislocation of right hip, unilateral
Q65.02	Congenital dislocation of left hip, unilateral
Q65.1	Congenital dislocation of hip, bilateral
Q65.2	Congenital dislocation of hip, unspecified
Q65.31	Congenital partial dislocation of right hip, unilateral
Q65.32	Congenital partial dislocation of left hip, unilateral
Q65.4	Congenital partial dislocation of hip, bilateral
Q65.5	Congenital partial dislocation of hip, unspecified
Q65.6	Congenital unstable hip
Q65.81	Congenital coxa valga
Q65.82	Congenital coxa vara
Q65.89	Congenital coxa vara Other specified congenital deformities of hip
Q65.9	Congenital deformity of hip, unspecified
Q66.01	Congenital talipes equinovarus, right foot
Q66.02	Congenital talipes equinovarus, left foot
Q66.11	Congenital talipes calcaneovarus, right foot
Q66.12	Congenital talipes calcaneovarus, left foot
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Codes	Description
Q66.211	Congenital metatarsus primus varus, right foot
Q66.212	Congenital metatarsus primus varus, left foot
Q66.221	Congenital metatarsus adductus, right foot
Q66.222	Congenital metatarsus adductus, left foot
Q66.31	Other congenital varus deformities of feet, right foot
Q66.32	Other congenital varus deformities of feet, left foot
Q66.41	Congenital talipes calcaneovalgus, right foot
Q66.42	Congenital talipes calcaneovalgus, left foot
Q66.51	Congenital pes planus, right foot
Q66.52	Congenital pes plants, right foot  Congenital pes plants, left foot
Q66.6	Other congenital valgus deformities of feet
Q66.71	Congenital pes cavus, right foot
Q66.71	
Q66.81	Congenital pes cavus, left foot  Congenital vertical talus deformity, right foot
Q66.82	Congenital vertical talus deformity, right foot  Congenital vertical talus deformity, left foot
Q66.89	Other specified congenital deformities of feet
	Congenital deformity of feet, unspecified, right foot
Q66.91	
Q66.92	Congenital deformity of feet, unspecified, left foot
Q67.0	Congenital facial asymmetry
Q67.1	Congenital compression facies
Q67.2	Dolichocephaly
Q67.3	Plagiocephaly  Other parametric deformation of alvell force and investigations.
Q67.4	Other congenital deformities of skull, face and jaw
Q67.5	Congenital deformity of spine
Q67.6	Pectus excavatum
Q67.7	Pectus carinatum
Q67.8	Other congenital deformities of chest
Q68.0	Congenital deformity of sternocleidomastoid muscle
Q68.1	Congenital deformity of finger(s) and hand
Q68.2	Congenital deformity of knee
Q68.3	Congenital bowing of femur
Q68.4	Congenital bowing of tibia and fibula
Q68.5	Congenital bowing of long bones of leg, unspecified
Q68.6	Discoid meniscus
Q68.8	Other specified congenital musculoskeletal deformities
Q69.0	Accessory finger(s)
Q69.1	Accessory thumb(s)
Q69.2	Accessory toe(s)
Q69.9	Polydactyly, unspecified
Q70.01	Fused fingers, right hand
Q70.02	Fused fingers, left hand
Q70.03	Fused fingers, bilateral
Q70.11	Webbed fingers, right hand
Q70.12	Webbed fingers, left hand
Q70.13	Webbed fingers, bilateral
Q70.21	Fused toes, right foot
Q70.22	Fused toes, left foot
Q70.23	Fused toes, bilateral
Q70.31	Webbed toes, right foot
Q70.32	Webbed toes, left foot
Q70.33	Webbed toes, bilateral
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O70.4 Polysyndactyly, unspecified O70.9 Syndactyly, unspecified O71.01 Congenital complete absence of right upper limb O71.02 Congenital complete absence of left upper limb O71.03 Congenital complete absence of upper limb, bilateral O71.10 Congenital absence of right upper arm and forearm with hand present O71.11 Congenital absence of left upper arm and forearm with hand present O71.12 Congenital absence of upper arm and forearm with hand present O71.13 Congenital absence of both forearm and hand, right upper limb O71.21 Congenital absence of both forearm and hand, right upper limb O71.22 Congenital absence of both forearm and hand, left upper limb O71.23 Congenital absence of both forearm and hand, left upper limb O71.31 Congenital absence of both forearm and hand, left upper limb O71.32 Congenital absence of left hand and finger O71.33 Congenital absence of left hand and finger O71.34 Longitudinal reduction defect of right radius O71.45 Longitudinal reduction defect of right radius O71.46 Longitudinal reduction defect of right radius O71.47 Longitudinal reduction defect of right ulna O71.51 Longitudinal reduction defect of right ulna O71.53 Longitudinal reduction defect of light ulna O71.54 Longitudinal reduction defect of light ulna O71.55 Longitudinal reduction defect of light ulna O71.61 Lobster-claw light hand O71.62 Lobster-claw left hand O71.63 Lobster-claw left hand O71.64 Lobster-claw left hand O71.65 Lobster-claw left hand O71.66 Lobster-claw left hand O71.67 Lopenital shortening of right upper limb O71.810 Congenital shortening of right upper limb O71.811 Congenital shortening of left upper limb O71.820 Congenital shortening of left upper limb O71.831 Congenital shortening of left upper limb O71.892 Other reduction defects of right upper limb O71.893 Other reduction defects of right upper limb O71.893 Other reduction defects of right upper limb O72.01 Congenital absence of left upper limb O73.02 Congenital absence of left upper limb O74.93 Unspecified reduction defect of right upper limb O75.93 Cong	Codes	Description
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Q71.12         Congenital absence of left upper arm and forearm with hand present.           Q71.13         Congenital absence of upper arm and forearm with hand present, bilateral           Q71.21         Congenital absence of both forearm and hand, left upper limb           Q71.22         Congenital absence of both forearm and hand, left upper limb           Q71.23         Congenital absence of right hand and finger           Q71.32         Congenital absence of right hand and finger           Q71.33         Congenital absence of hand and finger, bilateral           Q71.41         Longitudinal reduction defect of right tradius           Q71.42         Longitudinal reduction defect of left radius           Q71.43         Longitudinal reduction defect of right ulna           Q71.51         Longitudinal reduction defect of right ulna           Q71.52         Longitudinal reduction defect of left ulna           Q71.53         Longitudinal reduction defect of left ulna           Q71.52         Longitudinal reduction defect of left ulna           Q71.53         Longitudinal reduction defect of left ulna           Q71.54         Lobster-claw left hand           Q71.55         Lobster-claw left hand           Q71.61         Lobster-claw left hand           Q71.62         Lobster-claw left hand           Q71.81         Congenital shor		
Q71.13         Congenital absence of upper arm and forearm with hand present, bilateral           Q71.21         Congenital absence of both forearm and hand, right upper limb           Q71.22         Congenital absence of both forearm and hand, left upper limb           Q71.33         Congenital absence of right hand and finger           Q71.31         Congenital absence of left hand and finger           Q71.32         Congenital absence of left hand and finger           Q71.33         Congenital absence of left hand and finger           Q71.41         Longitudinal reduction defect of left radius           Q71.42         Longitudinal reduction defect of left radius           Q71.43         Longitudinal reduction defect of left unda           Q71.51         Longitudinal reduction defect of left unda           Q71.52         Longitudinal reduction defect of left unda           Q71.53         Longitudinal reduction defect of left unda           Q71.61         Lobster-claw left hand           Q71.62         Lobster-claw left hand           Q71.63         Lobster-claw left hand           Q71.811         Congenital shortening of right upper limb           Q71.812         Congenital shortening of right upper limb           Q71.813         Congenital shortening of left upper limb           Q71.814         Congenital shortening of left		
Q71.21         Congenital absence of both forearm and hand, right upper limb           Q71.22         Congenital absence of both forearm and hand, left upper limb           Q71.31         Congenital absence of both forearm and hand, bilateral           Q71.32         Congenital absence of left hand and finger           Q71.33         Congenital absence of left hand and finger           Q71.41         Longitudinal reduction defect of right radius           Q71.42         Longitudinal reduction defect of left radius           Q71.43         Longitudinal reduction defect of right ulna           Q71.51         Longitudinal reduction defect of left una           Q71.52         Longitudinal reduction defect of left una           Q71.53         Longitudinal reduction defect of left una           Q71.53         Longitudinal reduction defect of left una           Q71.53         Longitudinal reduction defect of una, bilateral           Q71.64         Lobster-claw right hand           Q71.65         Lobster-claw hand, bilateral           Q71.61         Lobster-claw hand, bilateral           Q71.62         Lobster-claw hand, bilateral           Q71.81         Congenital shortening of right upper limb           Q71.81         Congenital shortening of left upper limb           Q71.81         Congenital shortening of upper limb, bilateral <td></td> <td></td>		
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Q71.42         Longitudinal reduction defect of left radius, bilateral           Q71.43         Longitudinal reduction defect of right ulna           Q71.51         Longitudinal reduction defect of left ulna           Q71.52         Longitudinal reduction defect of left ulna           Q71.53         Longitudinal reduction defect of ulna, bilateral           Q71.61         Lobster-claw left hand           Q71.62         Lobster-claw left hand           Q71.83         Lobster-claw hand, bilateral           Q71.811         Congenital shortening of right upper limb           Q71.812         Congenital shortening of left upper limb           Q71.813         Congenital shortening of upper limb, bilateral           Q71.891         Other reduction defects of right upper limb           Q71.892         Other reduction defects of left upper limb           Q71.893         Other reduction defects of left upper limb, bilateral           Q71.92         Unspecified reduction defect of left upper limb           Q71.93         Unspecified reduction defect of left upper limb, bilateral           Q72.01         Congenital complete absence of left lower limb           Q72.02         Congenital complete absence of left lower limb           Q72.03         Congenital absence of left thigh and lower leg with foot present           Q72.11         Congen		· ·
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Q71.51         Longitudinal reduction defect of left ulna           Q71.52         Longitudinal reduction defect of left ulna           Q71.53         Longitudinal reduction defect of left ulna           Q71.61         Lobster-claw right hand           Q71.62         Lobster-claw left hand           Q71.63         Lobster-claw hand, bilateral           Q71.811         Congenital shortening of right upper limb           Q71.812         Congenital shortening of left upper limb           Q71.813         Congenital shortening of upper limb, bilateral           Q71.891         Other reduction defects of right upper limb           Q71.892         Other reduction defects of left upper limb           Q71.893         Other reduction defects of upper limb, bilateral           Q71.91         Unspecified reduction defect of right upper limb           Q71.92         Unspecified reduction defect of upper limb, bilateral           Q72.03         Congenital complete absence of right lower limb           Q72.04         Congenital complete absence of left lower limb           Q72.05         Congenital complete absence of left lower limb           Q72.11         Congenital absence of left thigh and lower leg with foot present           Q72.12         Congenital absence of both lower leg with foot present           Q72.13         Congenital absenc		
Q71.52         Longitudinal reduction defect of left ulna           Q71.53         Longitudinal reduction defect of ulna, bilateral           Q71.61         Lobster-claw right hand           Q71.62         Lobster-claw left hand           Q71.63         Lobster-claw hand, bilateral           Q71.811         Congenital shortening of right upper limb           Q71.812         Congenital shortening of upper limb, bilateral           Q71.813         Congenital shortening of upper limb, bilateral           Q71.891         Other reduction defects of right upper limb           Q71.892         Other reduction defects of left upper limb           Q71.893         Other reduction defects of upper limb, bilateral           Q71.91         Unspecified reduction defect of left upper limb           Q71.92         Unspecified reduction defect of upper limb, bilateral           Q72.01         Congenital complete absence of right lower limb           Q72.02         Congenital complete absence of left lower limb           Q72.03         Congenital complete absence of lower limb, bilateral           Q72.11         Congenital absence of right thigh and lower leg with foot present           Q72.12         Congenital absence of left thigh and lower leg with foot present           Q72.13         Congenital absence of both lower leg and foot, left lower limb		
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Q71.62         Lobster-claw left hand           Q71.63         Lobster-claw hand, bilateral           Q71.811         Congenital shortening of right upper limb           Q71.812         Congenital shortening of left upper limb           Q71.813         Congenital shortening of upper limb, bilateral           Q71.891         Other reduction defects of right upper limb           Q71.892         Other reduction defects of left upper limb           Q71.893         Other reduction defect of right upper limb           Q71.91         Unspecified reduction defect of right upper limb           Q71.92         Unspecified reduction defect of left upper limb           Q71.93         Unspecified reduction defect of upper limb, bilateral           Q72.01         Congenital complete absence of right lower limb           Q72.02         Congenital complete absence of left lower limb           Q72.03         Congenital complete absence of lower limb, bilateral           Q72.10         Congenital absence of right thigh and lower leg with foot present           Q72.11         Congenital absence of left thigh and lower leg with foot present           Q72.12         Congenital absence of both lower leg with foot present, bilateral           Q72.21         Congenital absence of both lower leg and foot, right lower limb           Q72.22         Congenital absence of both lower leg and		
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Q71.892       Other reduction defects of left upper limb         Q71.893       Other reduction defects of upper limb, bilateral         Q71.91       Unspecified reduction defect of right upper limb         Q71.92       Unspecified reduction defect of left upper limb         Q71.93       Unspecified reduction defect of upper limb, bilateral         Q72.01       Congenital complete absence of right lower limb         Q72.02       Congenital complete absence of left lower limb         Q72.03       Congenital complete absence of lower limb, bilateral         Q72.03       Congenital complete absence of lower limb, bilateral         Q72.11       Congenital absence of right thigh and lower leg with foot present         Q72.12       Congenital absence of thigh and lower leg with foot present, bilateral         Q72.13       Congenital absence of both lower leg and foot, right lower limb         Q72.21       Congenital absence of both lower leg and foot, left lower limb         Q72.22       Congenital absence of both lower leg and foot, bilateral         Q72.23       Congenital absence of right foot and toe(s)         Q72.31       Congenital absence of right foot and toe(s)         Q72.32       Congenital absence of foot and toe(s)         Q72.33       Congenital absence of foot and toe(s), bilateral         Q72.41       Longitudinal reduction defect of left femur		
Q71.893       Other reduction defects of upper limb, bilateral         Q71.91       Unspecified reduction defect of right upper limb         Q71.92       Unspecified reduction defect of left upper limb         Q71.93       Unspecified reduction defect of upper limb, bilateral         Q72.01       Congenital complete absence of right lower limb         Q72.02       Congenital complete absence of left lower limb         Q72.03       Congenital complete absence of lower limb, bilateral         Q72.03       Congenital complete absence of lower limb, bilateral         Q72.11       Congenital absence of right thigh and lower leg with foot present         Q72.12       Congenital absence of left thigh and lower leg with foot present, bilateral         Q72.13       Congenital absence of both lower leg and foot, right lower limb         Q72.21       Congenital absence of both lower leg and foot, left lower limb         Q72.22       Congenital absence of both lower leg and foot, left lower limb         Q72.23       Congenital absence of both lower leg and foot, left lower limb         Q72.31       Congenital absence of both lower leg and foot, left lower limb         Q72.32       Congenital absence of right foot and toe(s)         Q72.33       Congenital absence of left foot and toe(s)         Q72.41       Longitudinal reduction defect of right femur         Q72.42		• 11
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Q72.52 Longitudinal reduction defect of left tibia Q72.53 Longitudinal reduction defect of tibia, bilateral		
Q72.53 Longitudinal reduction defect of tibia, bilateral		· ·
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Codes	Description
Q72.62	Longitudinal reduction defect of left fibula
Q72.63	Longitudinal reduction defect of fibula, bilateral
Q72.71	Split foot, right lower limb
Q72.71	Split foot, right lower limb
Q72.72	Split foot, left lower limb  Split foot, bilateral
Q72.73	'
Q72.812	Congenital shortening of right lower limb  Congenital shortening of left lower limb
Q72.813	ů ů
	Congenital shortening of lower limb, bilateral
Q72.891	Other reduction defects of right lower limb
Q72.892	Other reduction defects of left lower limb
Q72.893	Other reduction defects of lower limb, bilateral
Q72.91	Unspecified reduction defect of right lower limb
Q72.92	Unspecified reduction defect of left lower limb
Q72.93	Unspecified reduction defect of lower limb, bilateral
Q73.0	Congenital absence of unspecified limb(s)
Q73.1	Phocomelia, unspecified limb(s)
Q73.8	Other reduction defects of unspecified limb(s)
Q74.0	Other congenital malformations of upper limb(s), including shoulder girdle
Q74.1	Congenital malformation of knee
Q74.2	Other congenital malformations of lower limb(s), including pelvic girdle
Q74.3	Arthrogryposis multiplex congenita
Q74.8	Other specified congenital malformations of limb(s)
Q75.0	Craniosynostosis
Q75.1	Craniofacial dysostosis
Q75.2	Hypertelorism
Q75.3	Macrocephaly
Q75.4	Mandibulofacial dysostosis
Q75.5	Oculomandibular dysostosis
Q75.8	Other specified congenital malformations of skull and face bones
Q75.9	Congenital malformation of skull and face bones, unspecified
Q76.0	Spina bifida occulta
Q76.1	Klippel-Feil syndrome
Q76.2	Congenital spondylolisthesis
Q76.3	Congenital scoliosis due to congenital bony malformation
Q76.411	Congenital kyphosis, occipito-atlanto-axial region
Q76.412	Congenital kyphosis, cervical region
Q76.413	Congenital kyphosis, cervicothoracic region
Q76.414	Congenital kyphosis, thoracic region
Q76.415	Congenital kyphosis, thoracolumbar region
Q76.425	Congenital lordosis, thoracolumbar region
Q76.426	Congenital lordosis, lumbar region
Q76.427	Congenital lordosis, lumbosacral region
Q76.428	Congenital lordosis, sacral and sacrococcygeal region
Q76.49	Other congenital malformations of spine, not associated with scoliosis
Q76.5	Cervical rib
Q76.6	Other congenital malformations of ribs
Q76.7	Congenital malformation of sternum
Q76.8	Other congenital malformations of bony thorax
Q76.9	Congenital malformation of bony thorax, unspecified
Q77.0	Achondrogenesis
Q77.1	Thanatophoric short stature

Codes	Description
Q77.2	Short rib syndrome
Q77.3	Chondrodysplasia punctata
Q77.4	Achondroplasia
Q77.5	Diastrophic dysplasia
Q77.6	Chondroectodermal dysplasia
Q77.7	Spondyloepiphyseal dysplasia
Q77.8	Other osteochondrodysplasia with defects of growth of tubular bones and spine
	Osteochondrodysplasia with defects of growth of tubular bones and spine,
Q77.9	unspecified
Q78.0	Osteogenesis imperfecta
Q78.1	Polyostotic fibrous dysplasia
Q78.2	Osteopetrosis
Q78.3	Progressive diaphyseal dysplasia
Q78.4	Enchondromatosis
Q78.5	Metaphyseal dysplasia
Q78.6	Multiple congenital exostoses
Q78.8	Other specified osteochondrodysplasias
Q78.9	Osteochondrodysplasia, unspecified
Q79.0	Congenital diaphragmatic hernia
Q79.1	Other congenital malformations of diaphragm
Q79.2	Exomphalos
Q79.3	Gastroschisis
Q79.4	Prune belly syndrome
Q79.51	Congenital hernia of bladder
Q79.60	Ehlers-Danlos syndrome, unspecified
Q79.60 Q79.61	Classical Ehlers-Danlos syndrome  Classical Ehlers-Danlos syndrome
Q79.62	Hypermobile Ehlers-Danlos syndrome
Q79.63 Q79.69	Vascular Ehlers-Danlos syndrome
·	Other Ehlers-Danlos syndromes
Q79.8	Other congenital malformations of musculoskeletal system
Q79.9	Congenital malformation of musculoskeletal system, unspecified
Q80.0	Ichthyosis vulgaris
Q80.1	X-linked ichthyosis
Q80.2	Lamellar ichthyosis
Q80.3	Congenital bullous ichthyosiform erythroderma
Q80.4	Harlequin fetus
Q80.8	Other congenital ichthyosis
Q80.9	Congenital ichthyosis, unspecified
Q81.0	Epidermolysis bullosa simplex
Q81.1	Epidermolysis bullosa letalis
Q81.2	Epidermolysis bullosa dystrophica
Q81.8	Other epidermolysis bullosa
Q81.9	Epidermolysis bullosa, unspecified
Q82.0	Hereditary lymphedema
Q82.1	Xeroderma pigmentosum
Q82.2	Congenital cutaneous mastocytosis
Q82.3	Incontinentia pigmenti
Q82.4	Ectodermal dysplasia (anhidrotic)
Q82.5	Congenital non-neoplastic nevus
Q82.6	Congenital sacral dimple
Q82.8	Other specified congenital malformations of skin
Q83.0	Congenital absence of breast with absent nipple

Codes	Description
Q83.1	Accessory breast
Q83.2	Absent nipple
Q83.3	
·	Accessory nipple
Q83.8	Other congenital malformations of breast
Q84.0	Congenital alopecia
Q84.1	Congenital morphological disturbances of hair, not elsewhere classified
Q84.2	Other congenital malformations of hair
Q84.3	Anonychia
Q84.4	Congenital leukonychia
Q84.5	Enlarged and hypertrophic nails
Q84.6	Other congenital malformations of nails
Q84.8	Other specified congenital malformations of integument
Q85.00	Neurofibromatosis, unspecified
Q85.01	Neurofibromatosis, type 1
Q85.02	Neurofibromatosis, type 2
Q85.03	Schwannomatosis
Q85.09	Other neurofibromatosis
Q85.1	Tuberous sclerosis
Q85.81	PTEN tumor syndrome
Q85.82	Other Cowden syndrome
Q85.83	Von Hippel-Lindau syndrome
Q85.89	Other phakomatoses, not elsewhere classified
Q85.9	Phakomatosis, unspecified
Q86.0	Fetal alcohol syndrome (dysmorphic)
Q86.1	Fetal hydantoin syndrome
Q86.2	Dysmorphism due to warfarin
Q86.8	Other congenital malformation syndromes due to known exogenous causes
Q87.0	Congenital malformation syndromes predominantly affecting facial appearance
Q87.11	Prader-Willi syndrome
Q87.19	Other congen malform synd predom assoc with short stature
Q87.2	Congenital malformation syndromes predominantly involving limbs
Q87.3	Congenital malformation syndromes involving early overgrowth
Q87.40	Marfan's syndrome, unspecified
Q87.410	Marfan's syndrome with aortic dilation
Q87.418	Marfan's syndrome with actite dilation  Marfan's syndrome with other cardiovascular manifestations
Q87.42	Marfan's syndrome with ocular manifestations
Q87.43	Marfan's syndrome with skeletal manifestation
Q87.5	Other congenital malformation syndromes with other skeletal changes
Q87.81	Alport syndrome
Q87.82	Arterial tortuosity syndrome  Other appointed expressited molformation avadrames, not alcouple a classified
Q87.89	Other specified congenital malformation syndromes, not elsewhere classified
Q89.01	Asplenia (congenital)
Q89.09	Congenital malformations of spleen
Q89.1	Congenital malformations of adrenal gland
Q89.2	Congenital malformations of other endocrine glands
Q89.3	Situs inversus
Q89.4	Conjoined twins
Q89.7	Multiple congenital malformations, not elsewhere classified
Q89.8	Other specified congenital malformations
Q90.0	Trisomy 21, nonmosaicism (meiotic nondisjunction)
Q90.1	Trisomy 21, mosaicism (mitotic nondisjunction)

090.2         Trisomy 21, translocation           090.9         Down syndrome, unspecified           091.0         Trisomy 18, normosaicism (milotic nondisjunction)           091.1         Trisomy 18, translocation           091.2         Trisomy 18, unspecified           091.3         Trisomy 13, unspecified           091.4         Trisomy 13, mosaicism (melotic nondisjunction)           091.5         Trisomy 13, mosaicism (milotic nondisjunction)           091.6         Trisomy 13, unspecified           092.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           092.1         Trisomy 13, unspecified           092.2         Partial trisomy           092.2         Partial trisomy           092.2         Partial trisomy           092.2         Partial trisomy           092.6         Marker chromosomes in normal individual           092.7         Triploidy and polyploidy           092.8         Other specified trisomies and partial trisomies of autosomes           092.9         Trisomy and partial trisomy of autosomes, unspecified           093.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           093.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           093.2         Chromos	Codes	Description
Down syndrome, unspecified   O91.0   Trisomy 18, nomosaicism (meiotic nondisjunction)   O91.1   Trisomy 18, nomosaicism (meiotic nondisjunction)   O91.2   Trisomy 18, translocation   O91.3   Trisomy 13, unspecified   O91.4   Trisomy 13, unspecified   O91.4   Trisomy 13, nomosaicism (meiotic nondisjunction)   O91.5   Trisomy 13, unspecified   O91.6   Trisomy 13, translocation   O91.7   Trisomy 13, translocation   O91.7   Trisomy 13, unspecified   O92.0   Whole chromosome trisomy, nomosaicism (meiotic nondisjunction)   O92.1   Whole chromosome trisomy, nomosaicism (meiotic nondisjunction)   O92.1   Whole chromosome trisomy, mosaicism (mitotic nondisjunction)   O92.2   Partial trisomy   O92.2   Partial trisomy   O92.2   Partial trisomy   O92.2   Partial trisomy   O92.2   Outplications with other complex rearrangements   O92.5   Outplications   O02.5   Outplications   Outplications   O02.5   Outplications   Outplications   O02.5   Outplications   Outplications		
Op.   Trisomy 18, nonmosaicism (meiotic nondisjunction)		
Q91.1         Trisomy 18, mosaicism (mitotic nondisjunction)           Q91.2         Trisomy 18, unspecified           Q91.3         Trisomy 18, unspecified           Q91.4         Trisomy 13, nonmosaicism (meiotic nondisjunction)           Q91.5         Trisomy 13, unspecified           Q91.6         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (mitotic nondisjunction)           Q93.1         Whole chromosome replaced with ring, dicentric or isochromosome           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome <td></td> <td></td>		
Q91.2         Trisomy 18, translocation           Q91.3         Trisomy 18, unspecified           Q91.4         Trisomy 13, nomosacism (meiotic nondisjunction)           Q91.5         Trisomy 13, mosaicism (mitotic nondisjunction)           Q91.6         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.61         Angelman syndrome           Q93		
Q91.3         Trisomy 18, unspecified           Q91.4         Trisomy 13, nonmosalcism (meiotic nondisjunction)           Q91.6         Trisomy 13, mosalcism (mitotic nondisjunction)           Q91.6         Trisomy 13, translocation           Q91.7         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosalcism (mitotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.6         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (mitotic nondisjunction)           Q93.1         Whole chromosome monosomy, nomosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.5         Angelman syndrome           Q93.8         Other deletions of part of a chromosome      <		
Q91.4         Trisomy 13, nonmosaicism (meiotic nondisjunction)           Q91.5         Trisomy 13, mosaicism (intotic nondisjunction)           Q91.6         Trisomy 13, translocation           Q91.7         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (mitotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.3         Deletion of short arm of chromosome 5           Q93.4         Deletion of short arm of chromosome           Q93.5         Angelman syndrome		
Q91.5         Trisomy 13, mosaicism (mitotic nondisjunction)           Q91.6         Trisomy 13, translocation           Q91.7         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.6         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (mitotic nondisjunction)           Q93.1         Whole chromosome monosomy, nomosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.4         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 4           Q93.5         Other deletions for art of a chromosome           Q93.7		
Q91.6         Trisomy 13, translocation           Q91.7         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.81         Velo-cardio-facial syndrome		· · · · · · · · · · · · · · · · · · ·
Q91.7         Trisomy 13, unspecified           Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, nonmosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-faci		
Q92.0         Whole chromosome trisomy, nonmosaicism (meiotic nondisjunction)           Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.5         Angelman syndrome           Q93.5         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.8         Velo-cardio-facial syndrome           Q93.8         Villiams syndrome           Q93.8         Other microdeletions from the autosomes		
Q92.1         Whole chromosome trisomy, mosaicism (mitotic nondisjunction)           Q92.2         Partial trisomy           Q92.61         Duplications with other complex rearrangements           Q92.62         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other deletions from the autosomes           Q93.89         Other deletions from the autosomes           Q93.89         Other deletions from the autosomal individual           Q95.1<		
Q92.2         Partial trisomy           Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonnosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other microdeletions           Q93.84         Other deletions from the autosomes           Q93.89         Other deletions from the autosomes           Q95.0         Balanced translocation and		
Q92.5         Duplications with other complex rearrangements           Q92.61         Marker chromosomes in normal individual           Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.5         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.8         Williams syndrome           Q93.8.2         Williams syndrome           Q93.8.8         Other microdeletions           Q93.8.9         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced autosomal rearrangement in abnormal individual     <		
Q92.61         Marker chromosomes in normal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nomosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other microdeletions           Q93.89         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced sex'autosomal rearrangement in abnormal individual           Q95.3         Balanced rearrangements and structural markers <t< td=""><td></td><td>·</td></t<>		·
Q92.62         Marker chromosomes in abnormal individual           Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.5         Angelman syndrome           Q93.5         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.8         Williams syndrome           Q93.8         Williams syndrome           Q93.8         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced sex/autosomal rearrangement in abnormal individual           Q95.3         Balanced sex/autosomal rearrangement in abnormal individual           Q95.5         Individual with autosomal fragile site <td></td> <td></td>		
Q92.7         Triploidy and polyploidy           Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.59         Other deletions of part of a chromosome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other microdeletions           Q93.89         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced autosomal rearrangement in abnormal individual           Q95.5         Individual with autosomal fragile site           Q96.0         Karyotype 45, X <td< td=""><td></td><td></td></td<>		
Q92.8         Other specified trisomies and partial trisomies of autosomes           Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonnosaicism (mictic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mictic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other microdeletions           Q93.89         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced autosomal rearrangement in abnormal individual           Q95.3         Balanced sex/autosomal rearrangement in abnormal individual           Q95.5         Individual with autosomal fragile site           Q95.8         Other balanced rearrangements and struct		
Q92.9         Trisomy and partial trisomy of autosomes, unspecified           Q93.0         Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)           Q93.1         Whole chromosome monosomy, mosaicism (mitotic nondisjunction)           Q93.2         Chromosome replaced with ring, dicentric or isochromosome           Q93.3         Deletion of short arm of chromosome 4           Q93.4         Deletion of short arm of chromosome 5           Q93.51         Angelman syndrome           Q93.59         Other deletions of part of a chromosome           Q93.7         Deletions with other complex rearrangements           Q93.81         Velo-cardio-facial syndrome           Q93.82         Williams syndrome           Q93.83         Other microdeletions           Q93.89         Other deletions from the autosomes           Q95.0         Balanced translocation and insertion in normal individual           Q95.1         Chromosome inversion in normal individual           Q95.2         Balanced autosomal rearrangement in abnormal individual           Q95.3         Balanced sex/autosomal rearrangement in abnormal individual           Q95.5         Individual with autosomal fragile site           Q96.0         Karyotype 45, X           Q96.1         Karyotype 45, X iso (Xq)           Q96.2 <td< td=""><td></td><td></td></td<>		
Q93.0       Whole chromosome monosomy, nonmosaicism (meiotic nondisjunction)         Q93.1       Whole chromosome monosomy, mosaicism (mitotic nondisjunction)         Q93.2       Chromosome replaced with ring, dicentric or isochromosome         Q93.3       Deletion of short arm of chromosome 4         Q93.4       Deletion of short arm of chromosome 5         Q93.51       Angelman syndrome         Q93.59       Other deletions of part of a chromosome         Q93.7       Deletions with other complex rearrangements         Q93.81       Velo-cardio-facial syndrome         Q93.82       Williams syndrome         Q93.83       Other microdeletions         Q93.89       Other deletions from the autosomes         Q95.0       Balanced translocation and insertion in normal individual         Q95.1       Chromosome inversion in normal individual         Q95.2       Balanced autosomal rearrangement in abnormal individual         Q95.3       Balanced sex/autosomal rearrangement in abnormal individual         Q95.5       Individual with autosomal fragile site         Q95.8       Other balanced rearrangements and structural markers         Q96.0       Karyotype 46, X iso (Xq)         Q96.1       Karyotype 46, X with abnormal sex chromosome, except iso (Xq)         Q96.2       Karyotype 46, X with abnorma		
Q93.1       Whole chromosome monosomy, mosaicism (mitotic nondisjunction)         Q93.2       Chromosome replaced with ring, dicentric or isochromosome         Q93.3       Deletion of short arm of chromosome 4         Q93.4       Deletion of short arm of chromosome 5         Q93.51       Angelman syndrome         Q93.7       Deletions with other complex rearrangements         Q93.81       Velo-cardio-facial syndrome         Q93.82       Williams syndrome         Q93.89       Other microdeletions         Q93.89       Other deletions from the autosomes         Q95.0       Balanced translocation and insertion in normal individual         Q95.1       Chromosome inversion in normal individual         Q95.2       Balanced autosomal rearrangement in abnormal individual         Q95.3       Balanced sex/autosomal rearrangement in abnormal individual         Q95.5       Individual with autosomal fragile site         Q95.8       Other balanced rearrangements and structural markers         Q96.0       Karyotype 45, X         Q96.1       Karyotype 46, X iso (Xq)         Q96.2       Karyotype 46, X with abnormal sex chromosome, except iso (Xq)         Q96.3       Mosaicism, 45, X/46, XX or XY         Q96.9       Turner's syndrome, unspecified         Q97.0       Ka		
Q93.2       Chromosome replaced with ring, dicentric or isochromosome         Q93.3       Deletion of short arm of chromosome 4         Q93.4       Deletion of short arm of chromosome 5         Q93.51       Angelman syndrome         Q93.59       Other deletions of part of a chromosome         Q93.7       Deletions with other complex rearrangements         Q93.81       Velo-cardio-facial syndrome         Q93.82       Williams syndrome         Q93.83       Other microdeletions         Q93.89       Other deletions from the autosomes         Q95.0       Balanced translocation and insertion in normal individual         Q95.1       Chromosome inversion in normal individual         Q95.2       Balanced autosomal rearrangement in abnormal individual         Q95.3       Balanced sex/autosomal rearrangement in abnormal individual         Q95.5       Individual with autosomal fragile site         Q95.8       Other balanced rearrangements and structural markers         Q96.0       Karyotype 45, X         Q96.1       Karyotype 46, X iso (Xq)         Q96.2       Karyotype 46, X with abnormal sex chromosome, except iso (Xq)         Q96.3       Mosaicism, 45, X/46, XX or XY         Q96.9       Turner's syndrome, unspecified         Q97.0       Karyotype 47, XXX		
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Q97.1 Female with more than three X chromosomes  Q97.2 Mosaicism, lines with various numbers of X chromosomes  Q97.3 Female with 46, XY karyotype	Q96.9	Turner's syndrome, unspecified
Q97.2 Mosaicism, lines with various numbers of X chromosomes  Q97.3 Female with 46, XY karyotype	Q97.0	Karyotype 47, XXX
Q97.3 Female with 46, XY karyotype	Q97.1	Female with more than three X chromosomes
	Q97.2	Mosaicism, lines with various numbers of X chromosomes
Q97.8 Other specified sex chromosome abnormalities, female phenotype	Q97.3	Female with 46, XY karyotype
	Q97.8	Other specified sex chromosome abnormalities, female phenotype
Q98.0 Klinefelter syndrome karyotype 47, XXY	Q98.0	Klinefelter syndrome karyotype 47, XXY
Q98.1 Klinefelter syndrome, male with more than two X chromosomes	Q98.1	Klinefelter syndrome, male with more than two X chromosomes

Codes	Description
Q98.3	Other male with 46, XX karyotype
Q98.4	Klinefelter syndrome, unspecified
Q98.5	Karyotype 47, XYY
Q98.6	Male with structurally abnormal sex chromosome
Q98.7	Male with sex chromosome mosaicism
Q98.8	Other specified sex chromosome abnormalities, male phenotype
Q99.0	Chimera 46, XX/46, XY
Q99.1	46, XX true hermaphrodite
Q99.2	Fragile X chromosome
Q99.8	Other specified chromosome abnormalities

## **Background, Product and Disclaimer Information**

Medical Necessity Guidelines are developed to determine coverage for benefits and are published to provide a better understanding of the basis upon which coverage decisions are made. We make coverage decisions using these guidelines, along with the Member's benefit document, and in coordination with the Member's physician(s) on a case-by-case basis considering the individual Member's health care needs.

Medical Necessity Guidelines are developed for selected therapeutic or diagnostic services found to be safe and proven effective in a limited, defined population of patients or clinical circumstances. They include concise clinical coverage criteria based on current literature review, consultation with practicing physicians in our service area who are medical experts in the particular field, FDA and other government agency policies, and standards adopted by national accreditation organizations. We revise and update Medical Necessity Guidelines annually, or more frequently if new evidence becomes available that suggests needed revisions.

For self-insured plans, coverage may vary depending on the terms of the benefit document. If a discrepancy exists between a Medical Necessity Guideline and a self-insured Member's benefit document, the provisions of the benefit document will govern. For Tufts Health Together (Medicaid), coverage may be available beyond these guidelines for pediatric members under age 21 under the Early and Periodic Screening, Diagnostic and Treatment (EPSDT) benefits of the plan in accordance with 130 CMR 450.140 and 130 CMR 447.000, and with prior authorization.

Treating providers are solely responsible for the medical advice and treatment of Members. The use of this guideline is not a guarantee of payment or a final prediction of how specific claim(s) will be adjudicated. Claims payment is subject to eligibility and benefits on the date of service, coordination of benefits, referral/authorization, utilization management guidelines when applicable, and adherence to plan policies, plan procedures, and claims editing logic.