CHAPTER XVII

Congenital malformations, deformations and chromosomal abnormalities

(Q00-Q99)

inborn errors of metabolism

Q00-Q07 Congenital malformations of the nervous system

Q10-Q18 Congenital malformations of eye, ear, face and neck

Q20-Q28 Congenital malformations of the circulatory system

Q30-Q34 Congenital malformations of the respiratory system

Q35-Q37 Cleft lip and cleft palate

Q38-Q45 Other congenital malformations of the digestive system

Q50-Q56 Congenital malformations of genital organs

Q60-Q64 Congenital malformations of the urinary system

Q65-Q79 Congenital malformations and deformations of the musculoskeletal system

Q80-Q89 Other congenital malformations

Q90-Q99 Chromosomal abnormalities, not elsewhere classified

Congenital malformations of the nervous system

Q00-Q07

Q00 Anencephaly and similar malformations

Q00.0 Anencephaly

Acephaly

Acrania

Amyelencephaly

Hemianencephaly

Hemicephaly

Q00.1 Craniorachischisis

Q00.2 Iniencephaly

Q01 Encephalocele

encephalomyelocele

hydroencephalocele

hydromeningocele, cranial

meningocele, cerebral

meningoencephalocele

Meckel-Gruber syndrome

Q01.0 Frontal encephalocele

Q01.1 Nasofrontal encephalocele

Q01.2 Occipital encephalocele

Q01.8 Encephalocele of other sites

Q01.9 Encephalocele, unspecified

Q02 Microcephaly

Hydromicrocephaly

Micrencephalon

Meckel-Gruber syndrome

Q03 Congenital hydrocephalus

hydrocephalus in newborn

Arnold-Chiari syndrome

hydrocephalus:acquired NOS

hydrocephalus:due to congenital toxoplasmosis

hydrocephalus:acquired, of newborn

hydrocephalus:with spina bifida

Q03.0 Malformations of aqueduct of Sylvius

Aqueduct of Sylvius:anomaly

Aqueduct of Sylvius:obstruction, congenital

Aqueduct of Sylvius:stenosis

Q03.1 Atresia of foramina of Magendie and Luschka

Dandy-Walker syndrome

Q03.8 Other congenital hydrocephalus

Q03.9 Congenital hydrocephalus, unspecified

Q04 Other congenital malformations of brain

cyclopia

macrocephaly

Q04.0 Congenital malformations of corpus callosum

Agenesis of corpus callosum

Q04.1 Arhinencephaly

Q04.2 Holoprosencephaly

Q04.3 Other reduction deformities of brain

Absenceof part of brain

Agenesisof part of brain

Aplasiaof part of brain

Hypoplasiaof part of brain

Agyria

Hydranencephaly

Lissencephaly

Microgyria

Pachygyria

congenital malformations of corpus callosum

Q04.4 Septo-optic dysplasia

Q04.5 Megalencephaly

Q04.6 Congenital cerebral cysts

Porencephaly

Schizencephaly

acquired porencephalic cyst

Q04.8 Other specified congenital malformations of brain

Macrogyria

Q04.9 Congenital malformation of brain, unspecified

Congenital:anomalyNOS of brain

Congenital:deformityNOS of brain

Congenital:disease or lesionNOS of brain

Congenital:multiple anomaliesNOS of brain

Q05 Spina bifida

hydromeningocele (spinal)

meningocele (spinal)

meningomyelocele

myelocele

myelomeningocele

rachischisis

spina bifida (aperta)(cystica)

syringomyelocele

Arnold-Chiari syndrome

spina bifida occulta

Q05.0 Cervical spina bifida with hydrocephalus

Q05.1 Thoracic spina bifida with hydrocephalus

Spina bifida:dorsalwith hydrocephalus

Spina bifida:thoracolumbarwith hydrocephalus

Q05.2 Lumbar spina bifida with hydrocephalus

Lumbosacral 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

Spina bifida:dorsal NOS

Spina bifida:thoracolumbar NOS

Q05.7 Lumbar spina bifida without hydrocephalus

Lumbosacral spina bifida NOS

Q05.8 Sacral spina bifida without hydrocephalus

Q05.9 Spina bifida, unspecified

Q06 Other congenital malformations of spinal cord

Q06.0 Amyelia

Q06.1 Hypoplasia and dysplasia of spinal cord

Atelomyelia

Myelatelia

Myelodysplasia of spinal cord

Q06.2 Diastematomyelia

Q06.3 Other congenital cauda equina malformations

Q06.4 Hydromyelia

Hydrorachis

Q06.8 Other specified congenital malformations of spinal cord

Q06.9 Congenital malformation of spinal cord, unspecified

Congenital:anomalyNOS of spinal cord or meninges

Congenital:deformityNOS of spinal cord or meninges

Congenital:disease or lesionNOS of spinal cord or meninges

Q07 Other congenital malformations of nervous system

familial dysautonomia [Riley-Day]

neurofibromatosis (nonmalignant)

Q07.0 Arnold-Chiari syndrome

Q07.8 Other specified congenital malformations of nervous system

Agenesis of nerve

Displacement of brachial plexus

Jaw-winking syndrome

Marcus Gunn syndrome

Q07.9 Congenital malformation of nervous system, unspecified

Congenital:anomalyNOS of nervous system

Congenital:deformityNOS of nervous system

Congenital:disease or lesionNOS of nervous system

Congenital malformations of eye, ear, face and neck

Q10-Q18

cleft lip and cleft palate

congenital malformation of:cervical spine

congenital malformation of:larynx

congenital malformation of:lip NEC

congenital malformation of:nose

congenital malformation of:parathyroid gland

congenital malformation of:thyroid gland

Q10 Congenital malformations of eyelid, lacrimal apparatus and orbit

cryptophthalmos:NOS

cryptophthalmos:syndrome

Q10.0 Congenital ptosis

Q10.1 Congenital ectropion

Q10.2 Congenital entropion

Q10.3 Other congenital malformations of eyelid

Ablepharon

Absence or agenesis of:cilia

Absence or agenesis of:eyelid

Accessory:eyelid

Accessory:eye muscle

Blepharophimosis, congenital

Coloboma of eyelid

Congenital malformation of eyelid NOS

Q10.4 Absence and agenesis of lacrimal apparatus

Absence of punctum lacrimale

Q10.5 Congenital stenosis and stricture of lacrimal duct

Q10.6 Other congenital malformations of lacrimal apparatus

Congenital malformation of lacrimal apparatus NOS

Q10.7 Congenital malformation of orbit

Q11 Anophthalmos, microphthalmos and macrophthalmos

Q11.0 Cystic eyeball

Q11.1 Other anophthalmos

Agenesisof eye

Aplasiaof eye

Q11.2 Microphthalmos

Cryptophthalmos NOS

Dysplasia of eye

Hypoplasia of eye

Rudimentary eye

cryptophthalmos syndrome

Q11.3 Macrophthalmos

macrophthalmos in congenital glaucoma

Q12 Congenital lens malformations

Q12.0 Congenital cataract

Q12.1 Congenital displaced lens

Q12.2 Coloboma of lens

Q12.3 Congenital aphakia

Q12.4 Spherophakia

Q12.8 Other congenital lens malformations

Q12.9 Congenital lens malformation, unspecified

Q13 Congenital malformations of anterior segment of eye

Q13.0 Coloboma of iris

Coloboma NOS

Q13.1 Absence of iris

Aniridia

Q13.2 Other congenital malformations of iris

Anisocoria, congenital

Atresia of pupil

Congenital malformation of iris NOS

Corectopia

Q13.3 Congenital corneal opacity

Q13.4 Other congenital corneal malformations

Congenital malformation of cornea NOS

Microcornea

Peter anomaly

Q13.5 Blue sclera

Q13.8 Other congenital malformations of anterior segment of eye

Rieger anomaly

Q13.9 Congenital malformation of anterior segment of eye, unspecified

Q14 Congenital malformations of posterior segment of eye

Q14.0 Congenital malformation of vitreous humour

Congenital vitreous opacity

Q14.1 Congenital malformation of retina

Congenital retinal aneurysm

Q14.2 Congenital malformation of optic disc

Coloboma of optic disc

Q14.3 Congenital malformation of choroid

Q14.8 Other congenital malformations of posterior segment of eye

Coloboma of the fundus

Q14.9 Congenital malformation of posterior segment of eye, unspecified

Q15 Other congenital malformations of eye

congenital nystagmus

ocular albinism

retinitis pigmentosa

Q15.0 Congenital glaucoma

Buphthalmos

Glaucoma of newborn

Hydrophthalmos

Keratoglobus, congenital, with glaucoma

Macrocornea with glaucoma

Macrophthalmos in congenital glaucoma

Megalocornea with glaucoma

Q15.8 Other specified congenital malformations of eye

Q15.9 Congenital malformation of eye, unspecified

Congenital:anomalyNOS of eye

Congenital:deformityNOS of eye

Q16 Congenital malformations of ear causing impairment of hearing

congenital deafness

Q16.0 Congenital absence of (ear) auricle

Q16.1 Congenital absence, atresia and stricture of auditory canal (external)

Atresia or stricture of osseous meatus

Q16.2 Absence of eustachian tube

Q16.3 Congenital malformation of ear ossicles

Fusion of ear ossicles

Q16.4 Other congenital malformations of middle ear

Congenital malformation of middle ear NOS

Q16.5 Congenital malformation of inner ear

Anomaly:membranous labyrinth

Anomaly:organ of Corti

Q16.9 Congenital malformation of ear causing impairment of hearing, unspecified

Congenital absence of ear NOS

Q17 Other congenital malformations of ear

preauricular sinus

Q17.0 Accessory auricle

Accessory tragus

Polyotia

Preauricular appendage or tag

Supernumerary:ear

Supernumerary:lobule

Q17.1 Macrotia

Q17.2 Microtia

Q17.3 Other misshapen ear

Pointed ear

Q17.4 Misplaced ear

Low-set ears

cervical auricle

Q17.5 Prominent ear

Bat ear

Q17.8 Other specified congenital malformations of ear

Congenital absence of lobe of ear

Q17.9 Congenital malformation of ear, unspecified

Congenital anomaly of ear NOS

Q18 Other congenital malformations of face and neck

cleft lip and cleft palate

conditions classified to Q67.0-Q67.4

congenital malformations of skull and face bones

cyclopia

dentofacial anomalies [including malocclusion]

malformation syndromes affecting facial appearance

persistent thyroglossal duct

Q18.0 Sinus, fistula and cyst of branchial cleft

Branchial vestige

Q18.1 Preauricular sinus and cyst

Fistula (of):auricle, congenital

Fistula (of):cervicoaural

Pretragal sinus and cyst

Q18.2 Other branchial cleft malformations

Branchial cleft malformation NOS

Cervical auricle

Otocephaly

Q18.3 Webbing of neck

Pterygium colli

Q18.4 Macrostomia

Q18.5 Microstomia

Q18.6 Macrocheilia

Hypertrophy of lip, congenital

Q18.7 Microcheilia

Q18.8 Other specified congenital malformations of face and neck

Medial:cystof face and neck

Medial:fistulaof face and neck

Medial:sinusof face and neck

Q18.9 Congenital malformation of face and neck, unspecified

Congenital anomaly NOS of face and neck

Congenital malformations of the circulatory system

Q20-Q28

Q20 Congenital malformations of cardiac chambers and connections

dextrocardia with situs inversus

mirror-image atrial arrangement with situs inversus

Q20.0 Common arterial trunk

Persistent truncus arteriosus

Q20.1 Double outlet right ventricle

Taussig-Bing syndrome

Q20.2 Double outlet left ventricle

Q20.3 Discordant ventriculoarterial connection

Dextrotransposition of aorta

Transposition of great vessels (complete)

Q20.4 Double inlet ventricle

Common ventricle

Cor triloculare biatriatum

Single ventricle

Q20.5 Discordant atrioventricular connection

Corrected transposition

Laevotransposition

Ventricular inversion

Q20.6 Isomerism of atrial appendages

Isomerism of atrial appendages with asplenia or polysplenia

Q20.8 Other congenital malformations of cardiac chambers and connections

Q20.9 Congenital malformation of cardiac chambers and connections, unspecified

Q21 Congenital malformations of cardiac septa

acquired cardiac septal defect

Q21.0 Ventricular septal defect

Q21.1 Atrial septal defect

Coronary sinus defect

Patent or persistent:foramen ovale

Patent or persistent:ostium secundum defect (type II)

Sinus venosus defect

Q21.2 Atrioventricular septal defect

Common atrioventricular canal

Endocardial cushion defect

Ostium primum atrial septal defect (type I)

Q21.3 Tetralogy of Fallot

Ventricular septal defect with pulmonary stenosis or atresia, dextroposition of aorta and hypertrophy of right ventricle.

Q21.4 Aortopulmonary septal defect

Aortic septal defect

Aortopulmonary window

Q21.8 Other congenital malformations of cardiac septa

Eisenmenger defect

Pentalogy of Fallot

Eisenmengercomplex

Eisenmengersyndrome

Q21.9 Congenital malformation of cardiac septum, unspecified

Septal (heart) defect NOS

Q22 Congenital malformations of pulmonary and tricuspid valves

Q22.0 Pulmonary valve atresia

Q22.1 Congenital pulmonary valve stenosis

Q22.2 Congenital pulmonary valve insufficiency

Congenital pulmonary valve regurgitation

Q22.3 Other congenital malformations of pulmonary valve

Congenital malformation of pulmonary valve NOS

Q22.4 Congenital tricuspid stenosis

Tricuspid atresia

Q22.5 Ebstein anomaly

Q22.6 Hypoplastic right heart syndrome

Q22.8 Other congenital malformations of tricuspid valve

Q22.9 Congenital malformation of tricuspid valve, unspecified

Q23 Congenital malformations of aortic and mitral valves

Q23.0 Congenital stenosis of aortic valve

Congenital aortic:atresia

Congenital aortic:stenosis

congenital subaortic stenosis

that in hypoplastic left heart syndrome

Q23.1 Congenital insufficiency of aortic valve

Bicuspid aortic valve

Congenital aortic insufficiency

Q23.2 Congenital mitral stenosis

Congenital mitral atresia

Q23.3 Congenital mitral insufficiency

Q23.4 Hypoplastic left heart syndrome

Atresia, or marked hypoplasia of aortic orifice or valve, with hypoplasia of ascending aorta and defective development of left ventricle (with mitral valve stenosis or atresia).

Q23.8 Other congenital malformations of aortic and mitral valves

Q23.9 Congenital malformation of aortic and mitral valves, unspecified

Q24 Other congenital malformations of heart

endocardial fibroelastosis

Q24.0 Dextrocardia

dextrocardia with situs inversus

isomerism of atrial appendages (with asplenia or polysplenia)

mirror-image atrial arrangement with situs inversus

Q24.1 Laevocardia

Q24.2 Cor triatriatum

Q24.3 Pulmonary infundibular stenosis

Q24.4 Congenital subaortic stenosis

Q24.5 Malformation of coronary vessels

Congenital coronary (artery) aneurysm

Q24.6 Congenital heart block

Q24.8 Other specified congenital malformations of heart

Congenital:diverticulum of left ventricle

Congenital:malformation of:myocardium

Congenital:malformation of:pericardium

Malposition of heart

Uhl disease

Q24.9 Congenital malformation of heart, unspecified

Congenital:anomalyNOS of heart

Congenital:diseaseNOS of heart

Q25 Congenital malformations of great arteries

Q25.0 Patent ductus arteriosus

Patent ductus Botalli

Persistent ductus arteriosus

Q25.1 Coarctation of aorta

Coarctation of aorta (preductal)(postductal)

Q25.2 Atresia of aorta

Q25.3 Stenosis of aorta

Supravalvular aortic stenosis

congenital stenosis of aortic valve

Q25.4 Other congenital malformations of aorta

Absenceof aorta

Aplasiaof aorta

Congenital:aneurysmof aorta

Congenital:dilatationof aorta

Aneurysm of sinus of Valsalva (ruptured)

Double aortic arch [vascular ring of aorta]

Hypoplasia of aorta

Persistent:convolutions of aortic arch

Persistent:right aortic arch

hypoplasia of aorta in hypoplastic left heart syndrome

Q25.5 Atresia of pulmonary artery

Q25.6 Stenosis of pulmonary artery

Supravalvular pulmonary stenosis

Q25.7 Other congenital malformations of pulmonary artery

Aberrant pulmonary artery

Agenesisof pulmonary artery

Aneurysm, congenitalof pulmonary artery

Anomalyof pulmonary artery

Hypoplasiaof pulmonary artery

Pulmonary arteriovenous aneurysm

Q25.8 Other congenital malformations of great arteries

Q25.9 Congenital malformation of great arteries, unspecified

Q26 Congenital malformations of great veins

Q26.0 Congenital stenosis of vena cava

Congenital stenosis of vena cava (inferior)(superior)

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

Absence of vena cava (inferior)(superior)

Azygos continuation of inferior vena cava

Persistent left posterior cardinal vein

Scimitar syndrome

Q26.9 Congenital malformation of great vein, unspecified

Anomaly of vena cava (inferior)(superior) NOS

Q27 Other congenital malformations of peripheral vascular system

anomalies of:cerebral and precerebral vessels

anomalies of:coronary vessels

anomalies of:pulmonary artery

congenital retinal aneurysm

haemangioma and lymphangioma

Q27.0 Congenital absence and hypoplasia of umbilical artery

Single umbilical artery

Q27.1 Congenital renal artery stenosis

Q27.2 Other congenital malformations of renal artery

Congenital malformation of renal artery NOS

Multiple renal arteries

Q27.3 Peripheral arteriovenous malformation

Arteriovenous aneurysm

acquired arteriovenous aneurysm

Q27.4 Congenital phlebectasia

Q27.8 Other specified congenital malformations of peripheral vascular system

Aberrant subclavian artery

Absenceof artery or vein NEC

Atresiaof artery or vein NEC

Congenital:aneurysm (peripheral)

Congenital:stricture, artery

Congenital:varix

Q27.9 Congenital malformation of peripheral vascular system, unspecified

Anomaly of artery or vein NOS

Q28 Other congenital malformations of circulatory system

congenital aneurysm:NOS

congenital aneurysm:coronary

congenital aneurysm:peripheral

congenital aneurysm:pulmonary

congenital aneurysm:retinal

ruptured:cerebral arteriovenous malformation

ruptured:malformation of precerebral vessels

Q28.0 Arteriovenous malformation of precerebral vessels

Congenital arteriovenous precerebral aneurysm (nonruptured)

Q28.1 Other malformations of precerebral vessels

Congenital:malformation of precerebral vessels NOS

Congenital:precerebral aneurysm (nonruptured)

Q28.2 Arteriovenous malformation of cerebral vessels

Arteriovenous malformation of brain NOS

Congenital arteriovenous cerebral aneurysm (nonruptured)

Q28.3 Other malformations of cerebral vessels

Congenital:cerebral aneurysm (nonruptured)

Congenital:malformation of cerebral vessels NOS

Q28.8 Other specified congenital malformations of circulatory system

Congenital aneurysm, specified site NEC

Q28.9 Congenital malformation of circulatory system, unspecified

Congenital malformations of the respiratory system

Q30-Q34

Q30 Congenital malformations of nose

congenital deviation of nasal septum

Q30.0 Choanal atresia

Atresiaof nares (anterior)(posterior)

Congenital stenosisof nares (anterior)(posterior)

Q30.1 Agenesis and underdevelopment of nose

Congenital absence of nose

Q30.2 Fissured, notched and cleft nose

Q30.3 Congenital perforated nasal septum

Q30.8 Other congenital malformations of nose

Accessory nose

Congenital anomaly of nasal sinus wall

Q30.9 Congenital malformation of nose, unspecified

Q31 Congenital malformations of larynx

congenital (laryngeal) stridor NOS

Q31.0 Web of larynx

Web of larynx:NOS

Web of larynx:glottic

Web of larynx:subglottic

Q31.1 Congenital subglottic stenosis

Q31.2 Laryngeal hypoplasia

Q31.3 Laryngocele

Q31.5 Congenital laryngomalacia

Q31.8 Other congenital malformations of larynx

Absenceof cricoid cartilage, epiglottis, glottis, larynx or thyroid cartilage

Agenesisof cricoid cartilage, epiglottis, glottis, larynx or thyroid cartilage

Atresiaof cricoid cartilage, epiglottis, glottis, larynx or thyroid cartilage

Cleft thyroid cartilage

Congenital stenosis of larynx NEC

Fissure of epiglottis

Posterior cleft of cricoid cartilage

Q31.9 Congenital malformation of larynx, unspecified

Q32 Congenital malformations of trachea and bronchus

congenital bronchiectasis

Q32.0 Congenital tracheomalacia

Q32.1 Other congenital malformations of trachea

Anomaly of tracheal cartilage

Atresia of trachea

Congenital:dilatationof trachea

Congenital:malformationof trachea

Congenital:stenosisof trachea

Congenital tracheocele

Q32.2 Congenital bronchomalacia

Q32.3 Congenital stenosis of bronchus

Q32.4 Other congenital malformations of bronchus

Absenceof bronchus

Agenesisof bronchus

Atresiaof bronchus

Congenital malformation NOSof bronchus

Diverticulumof bronchus

Q33 Congenital malformations of lung

Q33.0 Congenital cystic lung

Congenital:honeycomb lung

Congenital:lung disease:cystic

Congenital:lung disease:polycystic

cystic lung disease, acquired or unspecified

Q33.1 Accessory lobe of lung

Q33.2 Sequestration of lung

Q33.3 Agenesis of lung

Absence of lung (lobe)

Q33.4 Congenital bronchiectasis

Q33.5 Ectopic tissue in lung

Q33.6 Hypoplasia and dysplasia of lung

pulmonary hypoplasia associated with short gestation

Q33.8 Other congenital malformations of lung

Q33.9 Congenital malformation of lung, unspecified

Q34 Other congenital malformations of respiratory system

Q34.0 Anomaly of pleura

Q34.1 Congenital cyst of mediastinum

Q34.8 Other specified congenital malformations of respiratory system

Atresia of nasopharynx

Q34.9 Congenital malformation of respiratory system, unspecified

Congenital:absenceof respiratory organ

Congenital:anomaly NOSof respiratory organ

Cleft lip and cleft palate

Q35-Q37

Use additional code (Q30.2), if desired, to identify associated malformations of the nose.

Robin syndrome

Q35 Cleft palate

fissure of palate

palatoschisis

cleft palate with cleft lip

Q35.1 Cleft hard palate

Q35.3 Cleft soft palate

Q35.5 Cleft hard palate with cleft soft palate

Q35.7 Cleft uvula

Q35.9 Cleft palate, unspecified

Q36 Cleft lip

cheiloschisis

congenital fissure of lip

harelip

labium leporinum

cleft lip with cleft palate

Q36.0 Cleft lip, bilateral

Q36.1 Cleft lip, median

Q36.9 Cleft lip, unilateral

Cleft lip NOS

Q37 Cleft palate with cleft lip

Q37.0 Cleft hard palate with bilateral cleft lip

Q37.1 Cleft hard palate with unilateral cleft lip

Cleft hard palate with cleft lip NOS

Q37.2 Cleft soft palate with bilateral cleft lip

Q37.3 Cleft soft palate with unilateral cleft lip

Cleft soft palate with cleft lip NOS

Q37.4 Cleft hard and soft palate with bilateral cleft lip

Q37.5 Cleft hard and soft palate with unilateral cleft lip

Cleft hard and soft palate with cleft lip NOS

Q37.8 Unspecified cleft palate with bilateral cleft lip

Q37.9 Unspecified cleft palate with unilateral cleft lip

Cleft palate with cleft lip NOS

Other congenital malformations of the digestive system

Q38-Q45

Q38 Other congenital malformations of tongue, mouth and pharynx

macrostomia

microstomia

Q38.0 Congenital malformations of lips, not elsewhere classified

Congenital:fistula of lip

Congenital:malformation of lip NOS

Van der Woude syndrome

cleft lip

cleft lipwith cleft palate

macrocheilia

microcheilia

Q38.1 Ankyloglossia

Tongue tie

Q38.2 Macroglossia

Q38.3 Other congenital malformations of tongue

Aglossia

Bifid tongue

Congenital:adhesionof tongue

Congenital:fissureof tongue

Congenital:malformation NOSof tongue

Hypoglossia

Hypoplasia of tongue

Microglossia

Q38.4 Congenital malformations of salivary glands and ducts

Absence(of) salivary gland or duct

Accessory(of) salivary gland or duct

Atresia(of) salivary gland or duct

Congenital fistula of salivary gland

Q38.5 Congenital malformations of palate, not elsewhere classified

Absence of uvula

Congenital malformation of palate NOS

High arched palate

cleft palate

cleft palatewith cleft lip

Q38.6 Other congenital malformations of mouth

Congenital malformation of mouth NOS

Q38.7 Pharyngeal pouch

Diverticulum of pharynx

pharyngeal pouch syndrome

Q38.8 Other congenital malformations of pharynx

Congenital malformation of pharynx NOS

Q39 Congenital malformations of oesophagus

Q39.0 Atresia of oesophagus without fistula

Atresia of oesophagus NOS

Q39.1 Atresia of oesophagus with tracheo-oesophageal fistula

Atresia of oesophagus with broncho-oesophageal fistula

Q39.2 Congenital tracheo-oesophageal fistula without atresia

Congenital tracheo-oesophageal fistula NOS

Q39.3 Congenital stenosis and stricture of oesophagus

Q39.4 Oesophageal web

Q39.5 Congenital dilatation of oesophagus

Q39.6 Diverticulum of oesophagus

Oesophageal pouch

Q39.8 Other congenital malformations of oesophagus

Absent(of) oesophagus

Congenital displacement(of) oesophagus

Duplication(of) oesophagus

Q39.9 Congenital malformation of oesophagus, unspecified

Q40 Other congenital malformations of upper alimentary tract

Q40.0 Congenital hypertrophic pyloric stenosis

Congenital or infantile:constrictionof pylorus

Congenital or infantile:hypertrophyof pylorus

Congenital or infantile:spasmof pylorus

Congenital or infantile:stenosisof pylorus

Congenital or infantile:strictureof pylorus

Q40.1 Congenital hiatus hernia

Displacement of cardia through oesophageal hiatus

congenital diaphragmatic hernia

Q40.2 Other specified congenital malformations of stomach

Congenital:displacement of stomach

Congenital:diverticulum of stomach

Congenital:hourglass stomach

Duplication of stomach

Megalogastria

Microgastria

Q40.3 Congenital malformation of stomach, unspecified

Q40.8 Other specified congenital malformations of upper alimentary tract

Q40.9 Congenital malformation of upper alimentary tract, unspecified

Congenital:anomalyNOS of upper alimentary tract

Congenital:deformityNOS of upper alimentary tract

Q41 Congenital absence, atresia and stenosis of small intestine

congenital obstruction, occlusion and stricture of small intestine or intestine NOS

meconium ileus

Q41.0 Congenital absence, atresia and stenosis of duodenum

Q41.1 Congenital absence, atresia and stenosis of jejunum

Apple peel syndrome

Imperforate jejunum

Q41.2 Congenital absence, atresia and stenosis of ileum

Q41.8 Congenital absence, atresia and stenosis of other specified parts of small intestine

Q41.9 Congenital absence, atresia and stenosis of small intestine, part unspecified

Congenital absence, atresia and stenosis of intestine NOS

Q42 Congenital absence, atresia and stenosis of large intestine

congenital obstruction, occlusion and stricture of large intestine

Q42.0 Congenital absence, atresia and stenosis of rectum with fistula

Q42.1 Congenital absence, atresia and stenosis of rectum without fistula

Imperforate rectum

Q42.2 Congenital absence, atresia and stenosis of anus with fistula

Q42.3 Congenital absence, atresia and stenosis of anus without fistula

Imperforate anus

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 Other congenital malformations of intestine

Q43.0 Meckel diverticulum

Persistent:omphalomesenteric duct

Persistent:vitelline duct

Q43.1 Hirschsprung disease

Aganglionosis

Congenital (aganglionic) megacolon

Q43.2 Other congenital functional disorders of colon

Congenital dilatation of colon

Q43.3 Congenital malformations of intestinal fixation

Congenital adhesions [bands]:omental, anomalous

Congenital adhesions [bands]:peritoneal

Jackson membrane

Malrotation of colon

Rotation:failure ofof caecum and colon

Rotation:incompleteof caecum and colon

Rotation:insufficientof caecum and colon

Universal mesentery

Q43.4 Duplication of intestine

Q43.5 Ectopic anus

Q43.6 Congenital fistula of rectum and anus

congenital fistula:rectovaginal

congenital fistula:urethrorectal

pilonidal fistula or sinus

with absence, atresia and stenosis

Q43.7 Persistent cloaca

Cloaca NOS

Q43.8 Other specified congenital malformations of intestine

Congenital:blind loop syndrome

Congenital:diverticulitis, colon

Congenital:diverticulum, intestine

Dolichocolon

Megaloappendix

Megaloduodenum

Microcolon

Transposition of:appendix

Transposition of:colon

Transposition of:intestine

Q43.9 Congenital malformation of intestine, unspecified

Q44 Congenital malformations of gallbladder, bile ducts and liver

Q44.0 Agenesis, aplasia and hypoplasia of gallbladder

Congenital absence of gallbladder

Q44.1 Other congenital malformations of gallbladder

Congenital malformation of gallbladder NOS

Intrahepatic 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

Accessory hepatic duct

Congenital malformation of bile duct NOS

Duplication:biliary duct

Duplication:cystic duct

Q44.6 Cystic disease of liver

Fibrocystic disease of liver

Q44.7 Other congenital malformations of liver

Accessory liver

Alagille syndrome

Congenital:absence of liver

Congenital:hepatomegaly

Congenital:malformation of liver NOS

Q45 Other congenital malformations of digestive system

congenital:diaphragmatic hernia

congenital:hiatus hernia

Q45.0 Agenesis, aplasia and hypoplasia of pancreas

Congenital absence of pancreas

Q45.1 Annular pancreas

Q45.2 Congenital pancreatic cyst

Q45.3 Other congenital malformations of pancreas and pancreatic duct

Accessory pancreas

Congenital malformation of pancreas or pancreatic duct NOS

diabetes mellitus:congenital

diabetes mellitus:neonatal

fibrocystic disease of pancreas

Q45.8 Other specified congenital malformations of digestive system

Absence (complete)(partial) of alimentary tract NOS

Duplicationof digestive organs NOS

Malposition, congenitalof digestive organs NOS

Q45.9 Congenital malformation of digestive system, unspecified

Congenital:anomalyNOS of digestive system

Congenital:deformityNOS of digestive system

Congenital malformations of genital organs

Q50-Q56

androgen resistance syndrome

syndromes associated with anomalies in the number and form of chromosomes

testicular feminization syndrome

Q50 Congenital malformations of ovaries, fallopian tubes and broad ligaments

Q50.0 Congenital absence of ovary

Turner syndrome

Q50.1 Developmental ovarian cyst

Q50.2 Congenital torsion of ovary

Q50.3 Other congenital malformations of ovary

Accessory ovary

Congenital malformation of ovary NOS

Ovarian streak

Q50.4 Embryonic cyst of fallopian tube

Fimbrial cyst

Q50.5 Embryonic cyst of broad ligament

Cyst:epoophoron

Cyst:Gartner duct

Cyst:parovarian

Q50.6 Other congenital malformations of fallopian tube and broad ligament

Absence(of) fallopian tube or broad ligament

Accessory(of) fallopian tube or broad ligament

Atresia(of) fallopian tube or broad ligament

Congenital malformation of fallopian tube or broad ligament NOS

Q51 Congenital malformations of uterus and cervix

Q51.0 Agenesis and aplasia of uterus

Congenital absence of uterus

Q51.1 Doubling of uterus with doubling of cervix and vagina

Q51.2 Other doubling of uterus

Doubling of uterus NOS

Q51.3 Bicornate uterus

Q51.4 Unicornate uterus

Q51.5 Agenesis and aplasia of cervix

Congenital absence of cervix

Q51.6 Embryonic cyst of cervix

Q51.7 Congenital fistulae between uterus and digestive and urinary tracts

Q51.8 Other congenital malformations of uterus and cervix

Hypoplasia of uterus and cervix

Q51.9 Congenital malformation of uterus and cervix, unspecified

Q52 Other congenital malformations of female genitalia

Q52.0 Congenital absence of vagina

Q52.1 Doubling of vagina

Septate vagina

doubling of vagina with doubling of uterus and cervix

Q52.2 Congenital rectovaginal fistula

cloaca

Q52.3 Imperforate hymen

Q52.4 Other congenital malformations of vagina

Congenital malformation of vagina NOS

Cyst:canal of Nuck, congenital

Cyst:embryonic vaginal

Q52.5 Fusion of labia

Q52.6 Congenital malformation of clitoris

Q52.7 Other congenital malformations of vulva

Congenital:absenceof vulva

Congenital:cystof vulva

Congenital:malformation NOSof vulva

Q52.8 Other specified congenital malformations of female genitalia

Q52.9 Congenital malformation of female genitalia, unspecified

Q53 Undescended testicle

Q53.0 Ectopic testis

Unilateral or bilateral ectopic testes

Q53.1 Undescended testicle, unilateral

Q53.2 Undescended testicle, bilateral

Q53.9 Undescended testicle, unspecified

Cryptorchism NOS

Q54 Hypospadias

epispadias

Q54.0 Hypospadias, balanic

Hypospadias:coronal

Hypospadias:glandular

Q54.1 Hypospadias, penile

Q54.2 Hypospadias, penoscrotal

Q54.3 Hypospadias, perineal

Q54.4 Congenital chordee

Q54.8 Other hypospadias

Q54.9 Hypospadias, unspecified

Q55 Other congenital malformations of male genital organs

congenital hydrocele

hypospadias

Q55.0 Absence and aplasia of testis

Monorchism

Q55.1 Hypoplasia of testis and scrotum

Fusion of testes

Q55.2 Other congenital malformations of testis and scrotum

Congenital malformation of testis or scrotum NOS

Polyorchism

Retractile testis

Testis migrans

Q55.3 Atresia of vas deferens

Q55.4 Other congenital malformations of vas deferens, epididymis, seminal vesicles and prostate

Absence or aplasia of:prostate

Absence or aplasia of:spermatic cord

Congenital malformation of vas deferens, epididymis, seminal vesicles or prostate NOS

Q55.5 Congenital absence and aplasia of penis

Q55.6 Other congenital malformations of penis

Congenital malformation of penis NOS

Curvature of penis (lateral)

Hypoplasia of penis

Q55.8 Other specified congenital malformations of male genital organs

Q55.9 Congenital malformation of male genital organ, unspecified

Congenital:anomalyNOS of male genital organ

Congenital:deformityNOS of male genital organ

Q56 Indeterminate sex and pseudohermaphroditism

pseudohermaphroditism:female, with adrenocortical disorder

pseudohermaphroditism:male, with androgen resistance

pseudohermaphroditism:with specified chromosomal anomaly

Q56.0 Hermaphroditism, not elsewhere classified

Ovotestis

Q56.1 Male pseudohermaphroditism, not elsewhere classified

Male pseudohermaphroditism NOS

Q56.2 Female pseudohermaphroditism, not elsewhere classified

Female pseudohermaphroditism NOS

Q56.3 Pseudohermaphroditism, unspecified

Q56.4 Indeterminate sex, unspecified

Ambiguous genitalia

Congenital malformations of the urinary system

Q60-Q64

Q60 Renal agenesis and other reduction defects of kidney

atrophy of kidney:congenital

atrophy of kidney:infantile

congenital absence of kidney

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 syndrome

Q61 Cystic kidney disease

cyst of kidney (acquired)

Potter syndrome

Q61.0 Congenital single renal cyst

Congenital cyst of kidney (single)

Q61.1 Polycystic kidney, autosomal recessive

Polycystic kidney, infantile type

Q61.2 Polycystic kidney, autosomal dominant

Polycystic kidney, adult type

Q61.3 Polycystic kidney, unspecified

Q61.4 Renal dysplasia

Multicystic:kidney (developmental)

Multicystic:kidney disease

Multicystic:renal dysplasia

polycystic kidney disease

Multicystic:dyplastic kidney

Q61.5 Medullary cystic kidney

Sponge kidney NOS

Q61.8 Other cystic kidney diseases

Fibrocystic:kidney

Fibrocystic:renal degeneration or disease

Q61.9 Cystic kidney disease, unspecified

Meckel-Gruber syndrome

Q62 Congenital obstructive defects of renal pelvis and congenital malformations of ureter

Q62.0 Congenital hydronephrosis

Q62.1 Atresia and stenosis of ureter

Congenital occlusion of:ureter

Congenital occlusion of:ureteropelvic junction

Congenital occlusion of:ureterovesical orifice

Impervious ureter

Q62.2 Congenital megaloureter

Congenital dilatation of ureter

Q62.3 Other obstructive defects of renal pelvis and ureter

Congenital ureterocele

Q62.4 Agenesis of ureter

Absent ureter

Q62.5 Duplication of ureter

Accessoryureter

Doubleureter

Q62.6 Malposition of ureter

Deviation(of) ureter or ureteric orifice

Displacement(of) ureter or ureteric orifice

Ectopic(of) ureter or ureteric orifice

Implantation, anomalous(of) ureter or ureteric orifice

Q62.7 Congenital vesico-uretero-renal reflux

Q62.8 Other congenital malformations of ureter

Anomaly of ureter NOS

Q63 Other congenital malformations of kidney

congenital nephrotic syndrome

Q63.0 Accessory kidney

Q63.1 Lobulated, fused and horseshoe kidney

Q63.2 Ectopic kidney

Congenital displaced kidney

Malrotation of kidney

Q63.3 Hyperplastic and giant kidney

Q63.8 Other specified congenital malformations of kidney

Congenital renal calculi

Q63.9 Congenital malformation of kidney, unspecified

Q64 Other congenital malformations of urinary system

Q64.0 Epispadias

hypospadias

Q64.1 Exstrophy of urinary bladder

Ectopia vesicae

Extroversion of bladder

Q64.2 Congenital posterior urethral valves

Q64.3 Other atresia and stenosis of urethra and bladder neck

Congenital:bladder neck obstruction

Congenital:stricture of:urethra

Congenital:stricture of:urinary meatus

Congenital:stricture of:vesicourethral orifice

Impervious urethra

Q64.4 Malformation of urachus

Cyst of urachus

Patent urachus

Prolapse of urachus

Q64.5 Congenital absence of bladder and urethra

Q64.6 Congenital diverticulum of bladder

Q64.7 Other congenital malformations of bladder and urethra

Accessory:bladder

Accessory:urethra

Congenital:hernia of bladder

Congenital:malformation of bladder or urethra NOS

Congenital:prolapse of:bladder (mucosa)

Congenital:prolapse of:urethra

Congenital:prolapse of:urinary meatus

Congenital:urethrorectal fistula

Double:urethra

Double:urinary meatus

Q64.8 Other specified congenital malformations of urinary system

Q64.9 Congenital malformation of urinary system, unspecified

Congenital:anomalyNOS of urinary system

Congenital:deformityNOS of urinary system

Congenital malformations and deformations of the musculoskeletal system

Q65-Q79

Q65 Congenital deformities of hip

clicking hip

Q65.0 Congenital dislocation of hip, unilateral

Q65.1 Congenital dislocation of hip, bilateral

Q65.2 Congenital dislocation of hip, unspecified

Q65.3 Congenital subluxation of hip, unilateral

Q65.4 Congenital subluxation of hip, bilateral

Q65.5 Congenital subluxation of hip, unspecified

Q65.6 Unstable hip

Dislocatable hip

Subluxatable hip

Q65.8 Other congenital deformities of hip

Anteversion of femoral neck

Congenital acetabular dysplasia

Congenital coxa:valga

Congenital coxa:vara

Q65.9 Congenital deformity of hip, unspecified

Q66 Congenital deformities of feet

reduction defects of feet

valgus deformities (acquired)

varus deformities (acquired)

Q66.0 Talipes equinovarus

Q66.1 Talipes calcaneovarus

Q66.2 Metatarsus varus

Q66.3 Other congenital varus deformities of feet

Hallux varus, congenital

Q66.4 Talipes calcaneovalgus

Q66.5 Congenital pes planus

Flat foot:congenital

Flat foot:rigid

Flat foot:spastic (everted)

Q66.6 Other congenital valgus deformities of feet

Metatarsus valgus

Q66.7 Pes cavus

Q66.8 Other congenital deformities of feet

Clubfoot NOS

Hammer toe, congenital

Talipes:NOS

Talipes:asymmetric

Tarsal coalition

Vertical talus

Q66.9 Congenital deformity of feet, unspecified

Q67 Congenital musculoskeletal deformities of head, face, spine and chest

congenital malformation syndromes classified to Q87.-

Potter syndrome

Q67.0 Facial asymmetry

Q67.1 Compression facies

Q67.2 Dolichocephaly

Q67.3 Plagiocephaly

Q67.4 Other congenital deformities of skull, face and jaw

Depressions in skull

Deviation of nasal septum, congenital

Hemifacial atrophy or hypertrophy

Squashed or bent nose, congenital

dentofacial anomalies [including malocclusion]

syphilitic saddle nose

Q67.5 Congenital deformity of spine

Congenital scoliosis:NOS

Congenital scoliosis:postural

infantile idiopathic scoliosis

scoliosis due to congenital bony malformation

Q67.6 Pectus excavatum

Congenital funnel chest

Q67.7 Pectus carinatum

Congenital pigeon chest

Q67.8 Other congenital deformities of chest

Congenital deformity of chest wall NOS

Q68 Other congenital musculoskeletal deformities

reduction defects of limb(s)

Q68.0 Congenital deformity of sternocleidomastoid muscle

Congenital (sternomastoid) torticollis

Contracture of sternocleidomastoid (muscle)

Sternomastoid tumour (congenital)

Q68.1 Congenital deformity of hand

Congenital clubfinger

Spade-like hand (congenital)

Q68.2 Congenital deformity of knee

Congenital:dislocation of knee

Congenital:genu recurvatum

Q68.3 Congenital bowing of femur

anteversion of femur (neck)

Q68.4 Congenital bowing of tibia and fibula

Q68.5 Congenital bowing of long bones of leg, unspecified

Q68.8 Other specified congenital musculoskeletal deformities

Congenital:deformity of:clavicle

Congenital:deformity of:elbow

Congenital:deformity of:forearm

Congenital:deformity of:scapula

Congenital:dislocation of:elbow

Congenital:dislocation of:shoulder

Q69 Polydactyly

Q69.0 Accessory finger(s)

Q69.1 Accessory thumb(s)

Q69.2 Accessory toe(s)

Accessory hallux

Q69.9 Polydactyly, unspecified

Supernumerary digit(s) NOS

Q70 Syndactyly

Q70.0 Fused fingers

Complex syndactyly of fingers with synostosis

Q70.1 Webbed fingers

Simple syndactyly of fingers without synostosis

Q70.2 Fused toes

Complex syndactyly of toes with synostosis

Q70.3 Webbed toes

Simple syndactyly of toes without synostosis

Q70.4 Polysyndactyly

Q70.9 Syndactyly, unspecified

Symphalangy NOS

Q71 Reduction defects of upper limb

Q71.0 Congenital complete absence of upper limb(s)

Q71.1 Congenital absence of upper arm and forearm with hand present

Q71.2 Congenital absence of both forearm and hand

Q71.3 Congenital absence of hand and finger(s)

Q71.4 Longitudinal reduction defect of radius

Clubhand (congenital)

Radial clubhand

Q71.5 Longitudinal reduction defect of ulna

Q71.6 Lobster-claw hand

Q71.8 Other reduction defects of upper limb(s)

Congenital shortening of upper limb(s)

Q71.9 Reduction defect of upper limb, unspecified

Q72 Reduction defects of lower limb

Q72.0 Congenital complete absence of lower limb(s)

Q72.1 Congenital absence of thigh and lower leg with foot present

Q72.2 Congenital absence of both lower leg and foot

Q72.3 Congenital absence of foot and toe(s)

Q72.4 Longitudinal reduction defect of femur

Proximal femoral focal deficiency

Q72.5 Longitudinal reduction defect of tibia

Q72.6 Longitudinal reduction defect of fibula

Q72.7 Split foot

Q72.8 Other reduction defects of lower limb(s)

Congenital shortening of lower limb(s)

Q72.9 Reduction defect of lower limb, unspecified

Q73 Reduction defects of unspecified limb

Q73.0 Congenital absence of unspecified limb(s)

Amelia NOS

Q73.1 Phocomelia, unspecified limb(s)

Phocomelia NOS

Q73.8 Other reduction defects of unspecified limb(s)

Longitudinal reduction deformity of unspecified limb(s)

Ectromelia NOSof limb(s) NOS

Hemimelia NOSof limb(s) NOS

Reduction defectof limb(s) NOS

Q74 Other congenital malformations of limb(s)

polydactyly

reduction defect of limb

syndactyly

Q74.0 Other congenital malformations of upper limb(s), including shoulder girdle

Accessory carpal bones

Cleidocranial dysostosis

Congenital pseudarthrosis of clavicle

Macrodactylia (fingers)

Madelung deformity

Radioulnar synostosis

Sprengel deformity

Triphalangeal thumb

Q74.1 Congenital malformation of knee

Congenital:absence of patella

Congenital:dislocation of patella

Congenital:genu:valgum

Congenital:genu:varum

Rudimentary patella

congenital:dislocation of knee

congenital:genu recurvatum

nail patella syndrome

Q74.2 Other congenital malformations of lower limb(s), including pelvic girdle

Congenital:fusion of sacroiliac joint

Congenital:malformation (of):ankle (joint)

Congenital:malformation (of):sacroiliac (joint)

anteversion of femur (neck)

Q74.3 Arthrogryposis multiplex congenita

Q74.8 Other specified congenital malformations of limb(s)

Q74.9 Unspecified congenital malformation of limb(s)

Congenital anomaly of limb(s) NOS

Q75 Other congenital malformations of skull and face bones

congenital malformation of face NOS

congenital malformation syndromes classified to Q87.-

dentofacial anomalies [including malocclusion]

musculoskeletal deformities of head and face

skull defects associated with congenital anomalies of brain such as:anencephaly

skull defects associated with congenital anomalies of brain such as:encephalocele

skull defects associated with congenital anomalies of brain such as:hydrocephalus

skull defects associated with congenital anomalies of brain such as:microcephaly

Q75.0 Craniosynostosis

Acrocephaly

Imperfect fusion of skull

Oxycephaly

Trigonocephaly

Q75.1 Craniofacial dysostosis

Crouzon disease

Q75.2 Hypertelorism

Q75.3 Macrocephaly

Q75.4 Mandibulofacial dysostosis

Syndrome:Franceschetti

Syndrome:Treacher-Collins

Q75.5 Oculomandibular dysostosis

Q75.8 Other specified congenital malformations of skull and face bones

Absence of skull bone, congenital

Congenital deformity of forehead

Platybasia

Q75.9 Congenital malformation of skull and face bones, unspecified

Congenital anomaly of:face bones NOS

Congenital anomaly of:skull NOS

Q76 Congenital malformations of spine and bony thorax

congenital musculoskeletal deformities of spine and chest

Q76.0 Spina bifida occulta

meningocele (spinal)

spina bifida (aperta)(cystica)

Q76.1 Klippel-Feil syndrome

Cervical fusion syndrome

Q76.2 Congenital spondylolisthesis

Congenital spondylolysis

spondylolisthesis (acquired)

spondylolysis (acquired)

Q76.3 Congenital scoliosis due to congenital bony malformation

Hemivertebra fusion or failure of segmentation with scoliosis

Q76.4 Other congenital malformations of spine, not associated with scoliosis

Congenital:absence of vertebraunspecified or not associated with scoliosis

Congenital:fusion of spineunspecified or not associated with scoliosis

Congenital:kyphosisunspecified or not associated with scoliosis

Congenital:lordosisunspecified or not associated with scoliosis

Congenital:malformation of lumbosacral (joint) (region)unspecified or not associated with scoliosis

Hemivertebraunspecified or not associated with scoliosis

Malformation of spineunspecified or not associated with scoliosis

Platyspondylisisunspecified or not associated with scoliosis

Supernumerary vertebraunspecified or not associated with scoliosis

Q76.5 Cervical rib

Supernumerary rib in cervical region

Q76.6 Other congenital malformations of ribs

Accessory rib

Congenital:absence of rib

Congenital:fusion of ribs

Congenital:malformation of ribs NOS

short rib syndrome

Q76.7 Congenital malformation of sternum

Congenital absence of sternum

Sternum bifidum

Q76.8 Other congenital malformations of bony thorax

Q76.9 Congenital malformation of bony thorax, unspecified

Q77 Osteochondrodysplasia with defects of growth of tubular bones and spine

mucopolysaccharidosis

Q77.0 Achondrogenesis

Hypochondrogenesis

Q77.1 Thanatophoric short stature

Q77.2 Short rib syndrome

Asphyxiating thoracic dysplasia [Jeune]

Q77.3 Chondrodysplasia punctata

Q77.4 Achondroplasia

Hypochondroplasia

Osteosclerosis congenita

Q77.5 Dystrophic dysplasia

Q77.6 Chondroectodermal dysplasia

Ellis-van Creveld syndrome

Q77.7 Spondyloepiphyseal dysplasia

Q77.8 Other osteochondrodysplasia with defects of growth of tubular bones and spine

Q77.9 Osteochondrodysplasia with defects of growth of tubular bones and spine, unspecified

Q78 Other osteochondrodysplasias

Q78.0 Osteogenesis imperfecta

Fragilitas ossium

Osteopsathyrosis

Q78.1 Polyostotic fibrous dysplasia

Albright(-McCune)(-Sternberg) syndrome

Q78.2 Osteopetrosis

Albers-Sch?nberg syndrome

Q78.3 Progressive diaphyseal dysplasia

Camurati-Engelmann syndrome

Q78.4 Enchondromatosis

Maffucci syndrome

Ollier disease

Q78.5 Metaphyseal dysplasia

Pyle syndrome

Q78.6 Multiple congenital exostoses

Diaphyseal aclasis

Q78.8 Other specified osteochondrodysplasias

Osteopoikilosis

Q78.9 Osteochondrodysplasia, unspecified

Chondrodystrophy NOS

Osteodystrophy NOS

Q79 Congenital malformations of the musculoskeletal system, not elsewhere classified

congenital (sternomastoid) torticollis

Q79.0 Congenital diaphragmatic hernia

congenital hiatus hernia

Q79.1 Other congenital malformations of diaphragm

Absence of diaphragm

Congenital malformation of diaphragm NOS

Eventration of diaphragm

Q79.2 Exomphalos

Omphalocele

umbilical hernia

Q79.3 Gastroschisis

Q79.4 Prune belly syndrome

Q79.5 Other congenital malformations of abdominal wall

umbilical hernia

Q79.6 Ehlers-Danlos syndrome

Q79.8 Other congenital malformations of musculoskeletal system

Absence of:muscle

Absence of:tendon

Accessory muscle

Amyotrophia congenita

Congenital:constricting bands

Congenital:shortening of tendon

Poland syndrome

Q79.9 Congenital malformation of musculoskeletal system, unspecified

Congenital:anomaly NOSof musculoskeletal system NOS

Congenital:deformity NOSof musculoskeletal system NOS

Other congenital malformations

Q80-Q89

Q80 Congenital ichthyosis

Refsum disease

Q80.0 Ichthyosis vulgaris

Q80.1 X-linked ichthyosis

Q80.2 Lamellar ichthyosis

Collodion baby

Q80.3 Congenital bullous ichthyosiform erythroderma

Q80.4 Harlequin fetus

Q80.8 Other congenital ichthyosis

Q80.9 Congenital ichthyosis, unspecified

Q81 Epidermolysis bullosa

Q81.0 Epidermolysis bullosa simplex

Cockayne syndrome

Q81.1 Epidermolysis bullosa letalis

Herlitz syndrome

Q81.2 Epidermolysis bullosa dystrophica

Q81.8 Other epidermolysis bullosa

Q81.9 Epidermolysis bullosa, unspecified

Q82 Other congenital malformations of skin

acrodermatitis enteropathica

congenital erythropoietic porphyria

pilonidal cyst or sinus

Sturge-Weber(-Dimitri) syndrome

Q82.0 Hereditary lymphoedema

Q82.1 Xeroderma pigmentosum

Q82.2 Mastocytosis

Urticaria pigmentosa

malignant mastocytosis

Q82.3 Incontinentia pigmenti

Q82.4 Ectodermal dysplasia (anhidrotic)

Ellis-van Creveld syndrome

Q82.5 Congenital non-neoplastic naevus

Birthmark NOS

Naevus:flammeus

Naevus:portwine

Naevus:sanguineous

Naevus:strawberry

Naevus:vascular NOS

Naevus:verrucous

caf? au lait spots

lentigo

naevus:NOS

naevus:araneus

naevus:melanocytic

naevus:pigmented

naevus:spider

naevus:stellar

Q82.8 Other specified congenital malformations of skin

Abnormal palmar creases

Accessory skin tags

Benign familial pemphigus [Hailey-Hailey]

Cutis laxa (hyperelastica)

Dermatoglyphic anomalies

Inherited keratosis palmaris et plantaris

Keratosis follicularis [Darier-White]

Ehlers-Danlos syndrome

Q82.9 Congenital malformation of skin, unspecified

Q83 Congenital malformations of breast

absence of pectoral muscle

Q83.0 Congenital absence of breast with absent nipple

Q83.1 Accessory breast

Supernumerary breast

Q83.2 Absent nipple

Q83.3 Accessory nipple

Supernumerary nipple

Q83.8 Other congenital malformations of breast

Hypoplasia of breast

Q83.9 Congenital malformation of breast, unspecified

Q84 Other congenital malformations of integument

Q84.0 Congenital alopecia

Congenital atrichosis

Q84.1 Congenital morphological disturbances of hair, not elsewhere classified

Beaded hair

Monilethrix

Pili annulati

Menkes kinky hair syndrome

Q84.2 Other congenital malformations of hair

Congenital:hypertrichosis

Congenital:malformation of hair NOS

Persistent lanugo

Q84.3 Anonychia

nail patella syndrome

Q84.4 Congenital leukonychia

Q84.5 Enlarged and hypertrophic nails

Congenital onychauxis

Pachyonychia

Q84.6 Other congenital malformations of nails

Congenital:clubnail

Congenital:koilonychia

Congenital:malformation of nail NOS

Q84.8 Other specified congenital malformations of integument

Aplasia cutis congenita

Q84.9 Congenital malformation of integument, unspecified

Congenital:anomaly NOSof integument NOS

Congenital:deformity NOSof integument NOS

Q85 Phakomatoses, not elsewhere classified

ataxia telangiectasia [Louis-Bar]

familial dysautonomia [Riley-Day]

Q85.0 Neurofibromatosis (nonmalignant)

Von Recklinghausen disease

Q85.1 Tuberous sclerosis

Bourneville disease

Epiloia

Q85.8 Other phakomatoses, not elsewhere classified

Syndrome:Peutz-Jeghers

Syndrome:Sturge-Weber(-Dimitri)

Syndrome:von Hippel-Lindau

Meckel-Gruber syndrome

Q85.9 Phakomatosis, unspecified

Hamartosis NOS

Q86 Congenital malformation syndromes due to known exogenous causes, not elsewhere classified

iodine-deficiency-related hypothyroidism

nonteratogenic effects of substances transmitted via placenta or breast milk

Q86.0 Fetal alcohol syndrome (dysmorphic)

Q86.1 Fetal hydantoin syndrome

Meadow syndrome

Q86.2 Dysmorphism due to warfarin

Q86.8 Other congenital malformation syndromes due to known exogenous causes

Q87 Other specified congenital malformation syndromes affecting multiple systems

Q87.0 Congenital malformation syndromes predominantly affecting facial appearance

Acrocephalopolysyndactyly

Acrocephalosyndactyly [Apert]

Cryptophthalmos syndrome

Cyclopia

Syndrome:Goldenhar

Syndrome:Moebius

Syndrome:oro-facial-digital

Syndrome:Robin

Whistling face

Q87.1 Congenital malformation syndromes predominantly associated with short stature

Syndrome:Aarskog

Syndrome:Cockayne

Syndrome:De Lange

Syndrome:Dubowitz

Syndrome:Noonan

Syndrome:Prader-Willi

Syndrome:Robinow-Silverman-Smith

Syndrome:Russell-Silver

Syndrome:Seckel

Syndrome:Smith-Lemli-Opitz

Ellis-van Creveld syndrome

Q87.2 Congenital malformation syndromes predominantly involving limbs

Syndrome:Holt-Oram

Syndrome:Klippel-Tr?naunay-Weber

Syndrome:nail patella

Syndrome:Rubinstein-Taybi

Syndrome:sirenomelia

Syndrome:thrombocytopenia with absent radius [TAR]

Syndrome:VATER

Q87.3 Congenital malformation syndromes involving early overgrowth

Syndrome:Beckwith-Wiedemann

Syndrome:Sotos

Syndrome:Weaver

Q87.4 Marfan syndrome

Q87.5 Other congenital malformation syndromes with other skeletal changes

Q87.8 Other specified congenital malformation syndromes, not elsewhere classified

Syndrome:Alport

Syndrome:Laurence-Moon(-Bardet)-Biedl

Syndrome:Zellweger

Q89 Other congenital malformations, not elsewhere classified

Q89.0 Congenital malformations of spleen

Asplenia (congenital)

Congenital splenomegaly

isomerism of atrial appendages (with asplenia or polysplenia)

Q89.1 Congenital malformations of adrenal gland

congenital adrenal hyperplasia

Q89.2 Congenital malformations of other endocrine glands

Congenital malformation of parathyroid or thyroid gland

Persistent thyroglossal duct

Thyroglossal cyst

Q89.3 Situs inversus

Dextrocardia with situs inversus

Mirror-image atrial arrangement with situs inversus

Situs inversus or transversus:abdominalis

Situs inversus or transversus:thoracis

Transposition of viscera:abdominal

Transposition of viscera:thoracic

dextrocardia NOS

laevocardia

Q89.4 Conjoined twins

Craniopagus

Dicephaly

Double monster

Pygopagus

Thoracopagus

Q89.7 Multiple congenital malformations, not elsewhere classified

Monster NOS

Multiple congenital:anomalies NOS

Multiple congenital:deformities NOS

congenital malformation syndromes affecting multiple systems

Q89.8 Other specified congenital malformations

Q89.9 Congenital malformation, unspecified

Congenital:anomaly NOS

Congenital:deformity NOS

Chromosomal abnormalities, not elsewhere classified

Q90-Q99

Q90 Down syndrome

Q90.0 Trisomy 21, meiotic nondisjunction

Q90.1 Trisomy 21, mosaicism (mitotic nondisjunction)

Q90.2 Trisomy 21, translocation

Q90.9 Down syndrome, unspecified

Trisomy 21 NOS

Q91 Edwards syndrome and Patau syndrome

Q91.0 Trisomy 18, meiotic nondisjunction

Q91.1 Trisomy 18, mosaicism (mitotic nondisjunction)

Q91.2 Trisomy 18, translocation

Q91.3 Edwards syndrome, unspecified

Q91.4 Trisomy 13, meiotic nondisjunction

Q91.5 Trisomy 13, mosaicism (mitotic nondisjunction)

Q91.6 Trisomy 13, translocation

Q91.7 Patau syndrome, unspecified

Q92 Other trisomies and partial trisomies of the autosomes, not elsewhere classified

unbalanced translocations and insertions

trisomies of chromosomes 13, 18, 21

Q92.0 Whole chromosome trisomy, meiotic nondisjunction

Q92.1 Whole chromosome trisomy, mosaicism (mitotic nondisjunction)

Q92.2 Major partial trisomy

Whole arm or more duplicated.

Q92.3 Minor partial trisomy

Less than whole arm duplicated.

Q92.4 Duplications seen only at prometaphase

Q92.5 Duplications with other complex rearrangements

Q92.6 Extra marker chromosomes

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 Monosomies and deletions from the autosomes, not elsewhere classified

Q93.0 Whole chromosome monosomy, meiotic nondisjunction

Q93.1 Whole chromosome monosomy, mosaicism (mitotic nondisjunction)

Q93.2 Chromosome replaced with ring or dicentric

Q93.3 Deletion of short arm of chromosome 4

Wolff-Hirschorn syndrome

Q93.4 Deletion of short arm of chromosome 5

Cri-du-chat syndrome

Q93.5 Other deletions of part of a chromosome

Angelman syndrome

Q93.6 Deletions seen only at prometaphase

Q93.7 Deletions with other complex rearrangements

Q93.8 Other deletions from the autosomes

Q93.9 Deletion from autosomes, unspecified

Q95 Balanced rearrangements and structural markers, not elsewhere classified

Robertsonian and balanced reciprocal translocations and insertions

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

Q95.4 Individuals with marker heterochromatin

Q95.5 Individuals with autosomal fragile site

Q95.8 Other balanced rearrangements and structural markers

Q95.9 Balanced rearrangement and structural marker, unspecified

Q96 Turner syndrome

Noonan syndrome

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

Q96.4 Mosaicism, 45,X/other cell line(s) with abnormal sex chromosome

Q96.8 Other variants of Turner syndrome

Q96.9 Turner syndrome, unspecified

Q97 Other sex chromosome abnormalities, female phenotype, not elsewhere classified

Turner syndrome

Q97.0 Karyotype 47,XXX

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

Q97.8 Other specified sex chromosome abnormalities, female phenotype

Q97.9 Sex chromosome abnormality, female phenotype, unspecified

Q98 Other sex chromosome abnormalities, male phenotype, not elsewhere classified

Q98.0 Klinefelter syndrome karyotype 47,XXY

Q98.1 Klinefelter syndrome, male with more than two X chromosomes

Q98.2 Klinefelter syndrome, male with 46,XX karyotype

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

Q98.9 Sex chromosome abnormality, male phenotype, unspecified

Q99 Other chromosome abnormalities, not elsewhere classified

Q99.0 Chimera 46,XX/46,XY

Chimera 46,XX/46,XY true hermaphrodite

Q99.1 46,XX true hermaphrodite

46,XX with streak gonads

46,XY with streak gonads

Pure gonadal dysgenesis

Q99.2 Fragile X chromosome

Fragile X syndrome

Q99.8 Other specified chromosome abnormalities

Q99.9 Chromosomal abnormality, unspecified