

Appendix D. Reportable Conditions with ICD-10-CM Diagnosis Codes

Table D.1 Brain Malformations and Neural Tube Defects ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
1. Brain Malformations and Neural Tube Defects	Q00-Q05, Q07
Anencephaly	Q00.0
Craniorachischisis	Q00.1
Iniencephaly	Q00.2
Frontal encephalocele	Q01.0
Nasofrontal encephalocele	Q01.1
Occipital encephalocele	Q01.2
Encephalocele of other sites	Q01.8
Encephalocele, unspecified	Q01.9
Microcephaly	Q02
Malformations of aqueduct of Sylvius	Q03.0
Atresia of foramina of Magendie and Luschka (including Dandy-Walker)	Q03.1
Other congenital hydrocephalus (including obstructive hydrocephaly)	Q03.8
Congenital hydrocephalus, unspecified	Q03.9
Congenital malformations of corpus callosum	Q04.0
Arhinencephaly	Q04.1
Holoprosencephaly	Q04.2
Other reduction deformities of brain	Q04.3
Septo-optic dysplasia of brain	Q04.4
Congenital cerebral cyst (porencephaly, schizencephaly)	Q04.6
Other specified congenital malformations of brain (including ventriculomegaly)	Q04.8
Congenital malformation of brain, unspecified	Q04.9
Cervical spina bifida with hydrocephalus	Q05.0
Thoracic spina bifida with hydrocephalus	Q05.1
Lumbar spina bifida with hydrocephalus	Q05.2
Sacral spina bifida with hydrocephalus	Q05.3
Unspecified spina bifida with hydrocephalus	Q05.4
Cervical spina bifida without hydrocephalus	Q05.5
Thoracic spina bifida without hydrocephalus	Q05.6
Lumbar spina bifida without hydrocephalus	Q05.7
Sacral spina bifida without hydrocephalus	Q05.8
Spina bifida, unspecified	Q05.9
Arnold-Chiari syndrome without spina bifida or hydrocephalus	Q07.00
Arnold-Chiari syndrome with spina bifida	Q07.01
Arnold-Chiari syndrome with hydrocephalus	Q07.02
Arnold-Chiari syndrome with spina bifida and hydrocephalus	Q07.03



Table D.2 Eye Malformations ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
2. Eye Malformations	Q11-Q14, H47
Cystic eyeball	Q11.0
Other anophthalmos	Q11.1
Microphthalmos	Q11.2
Congenital cataract	Q12.0
Coloboma of lens	Q12.2
Coloboma of iris	Q13.0
Rieger's anomaly	Q13.81
Other congenital malformations of anterior segment of eye	Q13.89
Congenital malformation of anterior segment of eye, unspecified	Q13.9
Congenital malformation of retina	Q14.1
Congenital malformation of optic disc	Q14.2
Congenital malformation of choroid	Q14.3
Other congenital malformations of posterior segment of eye	Q14.8
Congenital malformation of posterior segment of eye, unspecified	Q14.9
Optic nerve hypoplasia, right eye	H47.031
Optic nerve hypoplasia, left eye	H47.032
Optic nerve hypoplasia, bilateral	H47.033
Optic nerve hypoplasia, unspecified eye	H47.039



Table D.3 Ear Malformations and Hearing Loss ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
3. Ear Malformations and Hearing Loss	Q16-18, H90, H91, H93
Congenital absence of (ear) auricle (anotia)	Q16.0
Congenital absence, atresia and stricture of auditory canal (external)	Q16.1
Absence of eustachian tube	Q16.2
Congenital malformation of ear ossicles	Q16.3
Other congenital malformations of middle ear	Q16.4
Congenital malformation of inner ear	Q16.5
Congenital malformation of ear causing impairment of hearing, unspecified	Q16.9
Microtia	Q17.2
Misplaced ear (low-set ear)	Q17.4
Congenital malformation of face and neck, unspecified (includes dysmorphic features and low-set ears)	Q18.9
Conductive hearing loss, bilateral	H90.0
Conductive hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side	H90.11
Conductive hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side	H90.12
Conductive hearing loss, unspecified	H90.2
Sensorineural hearing loss, bilateral	H90.3
Sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side	H90.41
Sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side	H90.42
Unspecified sensorineural hearing loss	H90.5
Mixed conductive and sensorineural hearing loss, bilateral	H90.6
Mixed conductive and sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side	H90.71
Mixed conductive and sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side	H90.72
Mixed conductive and sensorineural hearing loss, unspecified	H90.8
Conductive hearing loss, unilateral, right ear with restricted hearing on the contralateral side	H90.A11
Conductive hearing loss, unilateral, left ear with restricted hearing on the contralateral side	H90.A12
Sensorineural hearing loss, unilateral, right ear, with restricted hearing on the contralateral side	H90.A21
Sensorineural hearing loss, unilateral, left ear, with restricted hearing on the contralateral side	H90.A22
Mixed conductive and sensorineural hearing loss, unilateral, right ear with restricted hearing on the contralateral side	H90.A31



Birth Defect	ICD-10-CM
Mixed conductive and sensorineural hearing loss, unilateral, left ear with restricted hearing on the contralateral side	H90.A32
Other specified hearing loss, right ear	H91.8X1
Other specified hearing loss, left ear	H91.8X2
Other specified hearing loss, bilateral	H91.8X3
Other specified hearing loss, unspecified ear	H91.8X9
Unspecified hearing loss, unspecified ear	H91.90
Unspecified hearing loss, right ear	H91.91
Unspecified hearing loss, left ear	H91.92
Unspecified hearing loss, bilateral	H91.93
Other abnormal auditory perceptions, right ear	H93.291
Other abnormal auditory perceptions, left ear	H93.292
Other abnormal auditory perceptions, bilateral	H93.293
Other abnormal auditory perceptions, unspecified ear	H93.299
Disorders of right acoustic nerve	H93.3X1
Disorders of left acoustic nerve	H93.3X2
Disorders of bilateral acoustic nerves	H93.3X3
Disorders of unspecified acoustic nerve	H93.3X9

Table D.4 Congenital Heart Disease ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
4. Congenital Heart Disease	Q20-Q26
Common arterial trunk (TA)	Q20.0
Double outlet right ventricle (DORV, Taussig-Bing anomaly)	Q20.1
Discordant ventriculoarterial connection	Q20.3
Double inlet ventricle (single ventricle)	Q20.4
Discordant atrioventricular connection	Q20.5
Ventricular septal defect (VSD)	Q21.0
Atrial septal defect (ASD)	Q21.1
Atrioventricular septal defect (AVSD)	Q21.2
Tetralogy of Fallot (TOF)	Q21.3
Pulmonary valve atresia	Q22.0
Congenital pulmonary valve stenosis	Q22.1
Tricuspid valve stenosis	Q22.4
Ebstein's anomaly	Q22.5
Congenital stenosis of aortic valve	Q23.0
Hypoplastic left heart syndrome (HLHS)	Q23.4
Congenital malformation of heart, unspecified	Q24.9
Coarctation of aorta	Q25.1
Interruption of aortic arch (IAA)	Q25.21
Other atresia of aorta	Q25.29
Congenital malformation of the aorta, unspecified	Q25.40
Absence and aplasia of aorta	Q25.41
Hypoplasia of aorta	Q25.42
Congenital aneurysm of aorta	Q25.43
Congenital dilation of aorta	Q25.44
Double aortic arch	Q25.45
Tortuous aortic arch	Q25.46
Right aortic arch	Q25.47
Anomalous origin of subclavian artery	Q25.48
Other congenital malformations of aorta	Q25.49
Total anomalous pulmonary venous connection (TAPVC)	Q26.2



Table D.5 Orofacial Malformations ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
5. Orofacial	Q30, Q35-Q37
Choanal atresia	Q30.0
Cleft hard palate	Q35.1
Cleft soft palate	Q35.3
Cleft hard palate with cleft soft palate	Q35.5
Cleft uvula	Q35.7
Cleft palate, unspecified	Q35.9
Cleft lip, bilateral	Q36.0
Cleft lip, median	Q36.1
Cleft lip, unilateral	Q36.9
Cleft hard palate with bilateral cleft lip	Q37.0
Cleft hard palate with unilateral cleft lip	Q37.1
Cleft soft palate with bilateral cleft lip	Q37.2
Cleft soft palate with unilateral cleft lip	Q37.3
Cleft hard and soft palate with bilateral cleft lip	Q37.4
Cleft hard and soft palate with unilateral cleft lip	Q37.5
Unspecified cleft palate with bilateral cleft lip	Q37.8
Unspecified cleft palate with unilateral cleft lip	Q37.9



Table D.6 Gastrointestinal Malformations ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
6. Gastrointestinal	Q39, Q41-42, Q44-Q45
Atresia of esophagus without fistula	Q39.0
Atresia of esophagus with tracheo-esophageal fistula	Q39.1
Congenital tracheo-esophageal fistula without atresia	Q39.2
Congenital stenosis and stricture of esophagus	Q39.3
Esophageal web	Q39.4
Congenital absence, atresia and stenosis of duodenum	Q41.0
Congenital absence, atresia and stenosis of jejunum	Q41.1
Congenital absence, atresia and stenosis of ileum	Q41.2
Congenital absence, atresia and stenosis of other specified parts of small intestine	Q41.8
Congenital absence, atresia and stenosis of small intestine, part unspecified	Q41.9
Congenital absence, atresia and stenosis of rectum with fistula	Q42.0
Congenital absence, atresia and stenosis of rectum without fistula	Q42.1
Congenital absence, atresia and stenosis of anus with fistula	Q42.2
Congenital absence, atresia and stenosis of anus without fistula (includes imperforate anus)	Q42.3
Congenital absence, atresia and stenosis of other parts of large intestine	Q42.8
Congenital absence, atresia and stenosis of large intestine, part unspecified	Q42.9
Atresia of bile ducts	Q44.2
Obstruction of bile ducts	Q44.3
Congenital malformation of digestive system, unspecified	Q45.9



Table D.7 Genitourinary Malformations ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
7. Genitourinary	Q54, Q60, Q64
Hypospadias, balanic	Q54.0
Hypospadias, penile	Q54.1
Hypospadias, penoscrotal	Q54.2
Hypospadias, perineal	Q54.3
Other hypospadias	Q54.8
Hypospadias, unspecified	Q54.9
Renal agenesis, unilateral	Q60.0
Renal agenesis, bilateral	Q60.1
Renal agenesis, unspecified	Q60.2
Renal hypoplasia, unilateral	Q60.3
Renal hypoplasia, bilateral	Q60.4
Renal hypoplasia, unspecified	Q60.5
Potter's syndrome	Q60.6
Exstrophy of urinary bladder, unspecified	Q64.10
Cloacal exstrophy of urinary bladder	Q64.12
Other exstrophy of urinary bladder	Q64.19
Congenital posterior urethral valves	Q64.2
Congenital malformation of urinary system, unspecified	Q64.9



Table D.8 Musculoskeletal Malformations ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
9. Musculoskeletal	Q66, Q68, Q71-Q75, Q79
Congenital talipes equinovarus (clubfoot)	Q66.0
Other specified congenital deformities of feet	Q66.89
Other specified congenital musculoskeletal deformities	Q68.8
Congenital complete absence of unspecified upper limb	Q71.00
Congenital complete absence of right upper limb	Q71.01
Congenital complete absence of left upper limb	Q71.02
Congenital complete absence of upper limb bilateral	Q71.03
Congenital absence of unspecified upper arm and forearm with hand present	Q71.10
Congenital absence of right upper arm and forearm with hand present	Q71.11
Congenital absence of left upper arm and forearm with hand present	Q71.12
Congenital absence of upper arm and forearm with hand present bilateral	Q71.13
Congenital absence of both forearm and hand, unspecified upper limb	Q71.20
Congenital absence of both forearm and hand, right upper limb	Q71.21
Congenital absence of both forearm and hand, left upper limb	Q71.22
Congenital absence of both forearm and hand, bilateral	Q71.23
Congenital absence of unspecified hand and finger	Q71.30
Congenital absence of right hand and finger	Q71.31
Congenital absence of left hand and finger	Q71.32
Congenital absence of hand and finger, bilateral	Q71.33
Longitudinal reduction defect of unspecified radius	Q71.40
Longitudinal reduction defect of right radius	Q71.41
Longitudinal reduction defect of left radius	Q71.42
Longitudinal reduction defect of radius, bilateral	Q71.43
Longitudinal reduction defect of unspecified ulna	Q71.50
Longitudinal reduction defect of right ulna	Q71.51
Longitudinal reduction defect of left ulna	Q71.52
Longitudinal reduction defect of ulna, bilateral	Q71.53
Lobster-claw unspecified hand	Q71.60
Lobster-claw right hand	Q71.61
Lobster-claw left hand	Q71.62
Lobster-claw hand, bilateral	Q71.63
Congenital shortening of right upper limb	Q71.811
Congenital shortening of left upper limb	Q71.812
Congenital shortening of upper limb, bilateral	Q71.813
Congenital shortening of unspecified upper limb	Q71.819
Other reduction defects of right upper limb	Q71.891
Other reduction defects of left upper limb	Q71.892
Other reduction defects of upper limb, bilateral	Q71.893



Birth Defect	ICD-10-CM
Other reduction defects of unspecified upper limb	Q71.899
Unspecified reduction defect of unspecified upper limb	Q71.90
Unspecified reduction defect of right upper limb	Q71.91
Unspecified reduction defect of left upper limb	Q71.92
Unspecified reduction defect of upper limb, bilateral	Q71.93
Congenital complete absence of unspecified lower limb	Q72.00
Congenital complete absence of right lower limb	Q72.01
Congenital complete absence of left lower limb	Q72.02
Congenital complete absence of lower limb, bilateral	Q72.03
Congenital absence of unspecified thigh and lower leg with foot present	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 thigh and lower leg with foot present, bilateral	Q72.13
Congenital absence of both lower leg and foot, unspecified lower limb	Q72.20
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 unspecified foot and toe(s)	Q72.30
Congenital absence of right foot and toe(s)	Q72.31
Congenital absence of left foot and toe(s)	Q72.32
Congenital absence of foot and toe(s), bilateral	Q72.33
Longitudinal reduction defect of unspecified femur	Q72.40
Longitudinal reduction defect of right femur	Q72.41
Longitudinal reduction defect of left femur	Q72.42
Longitudinal reduction defect of femur, bilateral	Q72.43
Longitudinal reduction defect of unspecified tibia	Q72.50
Longitudinal reduction defect of right tibia	Q72.51
Longitudinal reduction defect of left tibia	Q72.52
Longitudinal reduction defect of tibia, bilateral	Q72.53
Longitudinal reduction defect of unspecified fibula	Q72.60
Longitudinal reduction defect of right fibula	Q72.61
Longitudinal reduction defect of left fibula	Q72.62
Longitudinal reduction defect of fibula, bilateral	Q72.63
Split foot, unspecified lower limb	Q72.70
Split foot, right lower limb	Q72.71
Split foot, left lower limb	Q72.72
Split foot, bilateral	Q72.73
Congenital shortening of right lower limb	Q72.811
Congenital shortening of left lower limb	Q72.812
Congenital shortening of lower limb, bilateral	Q72.813



Congenital shortening of unspecified lower limb	Q72.819
Birth Defect	ICD-10-CM
Other reduction defects of right lower limb	Q72.891
Other reduction defects of left lower limb	Q72.892
Other reduction defects of lower limb, bilateral	Q72.893
Other reductions defects of unspecified lower limb	Q72.899
Unspecified reduction defect of unspecified lower limb	Q72.90
Unspecified reduction defect of right lower limb	Q72.91
Unspecified reduction defect of left lower limb	Q72.92
Unspecified reduction defect of lower limb, bilateral	Q72.93
Congenital absence of unspecified limb(s)	Q73.0
Phocomelia, unspecified limb(s)	Q73.1
Other reduction defects of unspecified limb(s)	Q73.8
Arthrogryposis multiplex congenita	Q74.3
Craniosynostosis	Q75.0
Congenital diaphragmatic hernia	Q79.0
Other congenital malformations of diaphragm	Q79.1
Exomphalos (omphalocele)	Q79.2
Gastroschisis	Q79.3
Congenital malformation of musculoskeletal system, unspecified	Q79.9



Table D.9 Chromosomal Abnormalities, Sequences, and Syndromes ICD-10-CM Diagnosis Codes

Birth Defect	ICD-10-CM
10. Chromosomal Abnormalities, Sequences, and Syndromes	Q89-91, Q96, Q99
Multiple congenital malformations, not elsewhere classified	Q89.7
Congenital malformation, unspecified	Q89.9
Trisomy 21, nonmosaicism	Q90.0
Trisomy 21, mosaicism	Q90.1
Trisomy 21, translocation	Q90.2
Down syndrome, unspecified (Trisomy 21)	Q90.9
Trisomy 18, nonmosaicism	Q91.0
Trisomy 18, mosaicism	Q91.1
Trisomy 18, translocation	Q91.2
Trisomy 18, unspecified	Q91.3
Trisomy 13, nonmosaicism	Q91.4
Trisomy 13, mosaicism	Q91.5
Trisomy 13, translocation	Q91.6
Trisomy 13, unspecified	Q91.7
Velo-cardio-facial syndrome (including DiGeorge syndrome)	Q93.81
Turner's syndrome - Karyotype 45, X	Q96.0
Turner's syndrome - Karyotype 46, X iso (Xq)	Q96.1
Turner's syndrome - Karyotype 46, X with abnormal sex chromosome, except iso (Xq)	Q96.2
Turner's syndrome - Karyotype 45, X/46, XX or XY	Q96.3
Turner's syndrome - Karyotype 45, X/other cell line(s) with abnormal sex chromosome	Q96.4
Other variants of Turner's syndrome	Q96.8
Turner's syndrome, unspecified	Q96.9
Chromosomal abnormality, unspecified	Q99.9