NCADC reporting inclusions and their ICD–10–AM codes, by type of anomaly, 2017

Type of anomaly	ICD–10–AM (tenth edition)
Chromosomal	
Trisomy 21 (Down syndrome)	Q90, Q90.0–Q90.2, Q90.9
Trisomy 18 (Edwards syndrome)	Q91.0–Q91.3
Trisomy 13 (Patau syndrome)	Q91.4–Q91.7
Female sex chromosome (including Turner syndrome)	Q96, Q96.0–Q96.4, Q96.8–Q96.9, Q97.0–Q97.3, Q97.8
Male sex chromosome (including Klinefelter syndrome)	Q98, Q98.0–Q98.8
Other chromosomal	Q92, Q92.0–Q92.8, Q93.0–Q93.8, Q95.1–Q95.5, Q95.8, Q99.0–Q99.2, Q99.8
Circulatory system	
Transposition of the great vessels	Q20.1, Q20.3, Q20.5
Cardiac septal anomalies	Q21.0–Q21.2, Q21.4, Q21.8
Ventricular septal defect	Q21.0
Atrial septal defect	Q21.1
Atrioventricular septal defect	Q21.2
Other cardiac septal defect	Q21.4, Q21.8
Tetralogy of Fallot	Q21.3
Pulmonary valve atresia	Q22.0
Hypoplastic left heart syndrome	Q23.4
Patent ductus arteriosus	Q25.0
Coarctation of aorta	Q25.1
Other circulatory system	Q20.0, Q20.2, Q20.4, Q20.6, Q20.8, Q22, Q22.1–Q22.6, Q22.8, Q23, Q23.0–Q23.3, Q23.8, Q24, Q24.0–Q24.6, Q24.8, Q25.2–Q25.8, Q26.0–Q26.6, Q26.8, Q27.1–Q27.4, Q27.8, Q28.0–Q28.3, Q28.8
Digestive system	
Cleft lip and/or palate	Q35, Q35.1-Q35.3, Q35.9, Q36
Tracheo–oesophageal fistula	Q39.1–Q39.2
Anomalies leading to oesophageal obstruction (without fistula)	Q39.0, Q39.3–Q39.4
Congenital hypertrophic pyloric stenosis	Q40.0
Atresia/stenosis of intestines	Q41.0–Q41.2, Q41.8–Q41.9, Q42.0–Q42.3, Q42.8
Hirschsprung disease	Q43.1
Other digestive system	Q38.7–Q38.8, Q39.5–Q39.6, Q39.8, Q40.1–Q40.3, Q40.8, Q43.2–Q43.4, Q43.6–Q43.8, Q44, Q44.0, Q44.2–Q44.7, Q45.0–Q45.3, Q45.8
Eye, ear, face and neck and integuments	
Anophthalmia and microphthalmia	Q11.1–Q11.2
Microtia (with congenital absence, atresia, and stricture of external auditory canal)	Q17.2 with Q16.1
Other eye, ear, face and neck and integuments	Q10.6–Q10.7, Q11.0, Q12.2, Q13.1, Q13.3–Q13.4, Q13.8, Q15.0, Q16.0, Q16.1 (without Q17.2), Q80, Q80.0–Q80.4, Q80.8, Q81, Q81.0–Q81.2, Q81.8, Q82.0–Q82.4, Q83.0–Q83.1, Q84.0, Q85.0–Q85.1, Q85.8, Q86.0–Q86.2, Q86.8, Q87, Q87.0–Q87.5, Q87.8, Q89.0–Q89.4, Q89.7–Q89.8

(continued)

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Type of anomaly	ICD–10–AM (tenth edition)
Genital organs	
Doubling anomalies of the female genitalia	Q51.1–Q51.4, Q52.1
Hypospadias	Q54, Q54.0–Q54.3, Q54.8–Q54.9
Anomalies related to indeterminate sex	Q56, Q56.0–Q56.4
Other genital organ	Q50.0–Q50.6, Q51.0, Q51.5–Q51.9, Q52.0, Q52.2, Q52.7, Q53.0, Q55.0, Q55.3–Q55.5, Q55.8
Musculoskeletal system	
Congenital hip dislocation	Q65, Q65.0–Q65.2
Talipes	Q66, Q66.00, Q66.01, Q66.1, Q66.4
Polydactyly	Q69, Q69.0–Q69.2, Q69.9
Syndactyly	Q70, Q70.0–Q70.2, Q70.4, Q70.9
Reduction defect of upper limb(s)	Q71, Q71.0–Q71.6, Q71.8–Q71.9
Reduction defect of lower limb(s)	Q72, Q72.0–Q72.9
Congenital diaphragmatic hernia	Q79.0
Exomphalos	Q79.2
Gastroschisis	Q79.3
Other musculoskeletal system	Q65.3–Q65.5, Q65.9, Q67.5, Q73, Q73.0–Q73.1, Q73.8, Q74, Q74.0, Q74.2–Q74.5, Q74.8, Q75, Q75.1, Q76.1–Q76.3, Q76.5, Q76.7, Q77.0–Q77.8, Q78, Q78.0–Q78.6, Q78.8–Q78.9, Q79.1, Q79.4, Q79.6
Nervous system	
Neural tube defects	Q00, Q05, Q05.0–Q05.9, Q07.0, Q01
Anencephaly and related anomalies	Q00, Q00.1–Q00.2
Craniorachischisis	Q00.1
Iniencephaly	Q00.2
Encephalocele	Q01, Q01.0–Q01.2, Q01.8–Q01.9
Spina bifida (including Arnold–Chiari malformation)	Q05, Q05.0–Q05.9, Q07.0
Microcephaly	Q02
Congenital hydrocephalus	Q03, Q03.0–Q03.1, Q03.8–Q03.9
Other nervous system	Q04, Q04.0–Q04.6, Q04.8, Q06.0–Q06.4, Q06.8, Q07.8
Respiratory system	
Choanal atresia	Q30.0
Hypoplasia and dysplasia of lung	Q33.6
Other respiratory system	Q30.1, Q30.3, Q31.0–Q31.3, Q31.8, Q32.0–Q32.4, Q33, Q33.0, Q33.1–Q33.5, Q33.8, Q34.0–Q34.1, Q34.8
Urinary system	
Renal agenesis/hypoplasia	Q60, Q60.0–Q60.6
Cystic kidney disease	Q61, Q61.0–Q61.5, Q61.8–Q61.9
Exstrophy of urinary bladder	Q64.1
Other urinary system	Q62, Q62.0–Q62.8, Q63, Q63.0–Q63.3, Q63.8, Q64.0, Q64.2–Q64.3, Q64.5–Q64.6, Q64.8