Reportable Conditions List (Organized Numerically by ICD-10-CM Code)

ICD-10-CM Diagnosis Codes	Categories of Congenital Anomalies
F84.0 – F84.9*	Pervasive developmental disorders (includes autism, childhood disintegrative disorder, Asperger's, & Rett syndrome; **reported at any age)**
Q00.0 – Q07.9, G90.1	Central nervous system
Q10.0 – Q18.9, Q35.1 – Q38.8	Orofacial
Q20.0 – Q28.9, P29.3	Cardiovascular
Q30.0 – Q34.9	Respiratory
Q39.0 – Q45.9	Gastrointestinal
Q50.01 – Q64.9	Genitourinary
Q65.00 – Q79.9	Musculoskeletal
Q80.0 – Q84.9	Integument
Q85.0 – Q89.9, E78.71, E78.72	Congenital anomalies, other & unspecified
Q86.0, P04.3*	Fetal alcohol syndrome **(reported up to 5 years)**
Q90.0 – Q99.9	Chromosome & syndromes

ICD-10-CM Diagnosis Codes	Other Reportable Conditions
C00.0 – C41.9	Neoplasms
C88.0	Waldenstrom's macroglobulinemia
D47.4, D75.81 – D75.89, D89.2	Thrombophilias
D55.0 – D58.9	Hereditary hemolytic anemias
D61.0 – D61.8	Constitutional aplastic anemia
D66, D67, D68.0 – D68.9	Coagulation defects
D69.3 – D69.42	Primary thrombocytopenia
D70.0 – D76.3	White blood cell diseases
D75.0	Familial polycythemia
D75.81 – D75.89	Other specified diseases of blood & blood-forming organs
D80.0 – D89.9	Immune mechanism disorders
E00.0 – E00.9, E07.89, E25.0 – E25.9	Endocrine newborn screening conditions
E03.4, E07.89	Thyroid-binding globulin deficiency
E07.1, E07.9	Dyshormonogenic goiter
E10 – E11.9	Diabetes mellitus
E29.8	Testicular dysfunction, other
E70.0 – E74.9, E77.1, E80.3, E88.40 – E88.49, C96.5, C96.6, D81.810 – D81.819, H49.811 – H49.819	Metabolic newborn screening conditions
E75.00 – E75.29, G31.81, G31.82, G93.9, G94	Cerebral degenerations usually manifesting in childhood
E76.01 – E76.3	Mucopolysaccharidoses
E84.0 – E84.9	Cystic fibrosis
G12.0 – G12.9	Anterior horn cell disease
G71.0 – G73.7	Muscular dystrophies
H35.179	Retrolental fibroplasia
H35.50 – H35.54	Hereditary retinal dystrophies
H47.61 – H47.619	Cortical blindness
H49.00 – H51.9	Strabismus & other disorders of binocular eye movement
H54.0 – H54.3	Visual impairment
H54.8	Legal blindness, as defined in USA
H55.01	Congenital nystagmus
H90.0 – H91.93	Hearing loss
K40.0 – K40.91	Inguinal hernia
K42.0 – K42.9	Umbilical hernia
P02.8, P02.9	Amniotic band syndrome
M26.00 – M26.10	Anomalies of jaw
N44.00 – N44.04	Torsion of testes
P00.2**,** P04.1**, P04.4, P04.41, P04.49,** P04.9**, P96.1**	**Fetus/newborn exposure to maternal infection or substance**
P35.0 – P37.9, A33	Congenital infections

For additional information, please visit: www.birthdefects.in.gov

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Reportable Conditions List (Organized Alphabetically)

ICD-10-CM Diagnosis Codes	Categories of Congenital Anomalies
Q20.0 – Q28.9, P29.3	Cardiovascular
Q00.0 – Q07.9, G90.1	Central nervous system
Q90.0 – Q99.9	Chromosome & syndromes
Q85.0 – Q89.9, E78.71, E78.72	Congenital anomalies, other & unspecified
Q86.0, P04.3*	Fetal alcohol syndrome **(reported up to 5 years)**
Q39.0 – Q45.9	Gastrointestinal
Q50.01 – Q64.9	Genitourinary
Q80.0 – Q84.9	Integument
Q65.00 – Q79.9	Musculoskeletal
Q10.0 – Q18.9, Q35.1 – Q38.8	Orofacial
F84.0 – F84.9*	Pervasive developmental disorders (includes autism, childhood disintegrative disorder, Asperger's, & Rett syndrome; **reported at any age)**
Q30.0 – Q34.9	Respiratory

ICD-10-CM Diagnosis Codes	Other Reportable Conditions
P02.8, P02.9	Amniotic band syndrome
M26.00 – M26.10	Anomalies of jaw
G12.0 – G12.9	Anterior horn cell disease
E75.00 – E75.29, G31.81, G31.82, G93.9, G94	Cerebral degenerations usually manifesting in childhood
D66, D67, D68.0 –D68.9	Coagulation defects
P35.0 – P37.9, A33	Congenital infections
H55.01	Congenital nystagmus
D61.0 – D61.8	Constitutional aplastic anemia
H47.61 – H47.619	Cortical blindness
E84.0 – E84.9	Cystic fibrosis
E10 – E11.9	Diabetes mellitus
E07.1, E07.9	Dyshormonogenic goiter
E00.0 – E00.9, E07.89, E25.0 – E25.9	Endocrine newborn screening conditions
D75.0	Familial polycythemia
P00.2**,** P04.1**, P04.4, P04.41, P04.49,** P04.9**, P96.1**	**Fetus/newborn exposure to maternal infection or substance**
H90.0 – H91.93	Hearing loss
D55.0 – D58.9	Hereditary hemolytic anemias
H35.50 – H35.54	Hereditary retinal dystrophies
D80.0 – D89.9	Immune mechanism disorders
K40.0 – K40.91	Inguinal hernia
H54.8	Legal blindness, as defined in USA
E70.0 – E74.9, E77.1, E80.3, E88.40 – E88.49, C96.5, - C96.6, D81.810 – D81.819, H49.811 – H49.819	Metabolic newborn screening conditions
E76.01 – E76.3	Mucopolysaccharidoses
G71.0 – G73.7	Muscular dystrophies
C00.0 – C41.9	Neoplasms
D75.81 – D75.89	Other specified diseases of blood & blood-forming organs
D69.3 – D69.42	Primary thrombocytopenia:
H35.179	Retrolental fibroplasia
H49.00 – H51.9	Strabismus & other disorders of binocular eye movement
E29.8	Testicular dysfunction, other
D47.4, D68.51 – D68.69, D75.81 – D75.89, D89.2	Thrombophilias
N44.00 – N44.04	Torsion of testes
E03.4, E07.89	Thyroid-binding globulin deficiency
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H54.0 – H54.3	Visual impairment
C88.0	Waldenstrom's macroglobulinemia
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