



**State of Connecticut Department of Public Health
Birth Defect Registry Reporting Form (Revised 3/20/16)**



Baby's Last Name: _____ **DOB:** _____ **Accession #:** _____ **Sex:** _____

Birth Hospital: _____ **Medical Record #:** _____ **Birth Sequence:** _____

Mother's Last Name: _____ **Mother's First Name:** _____

Is a birth defect present in this child Yes (if yes, please refer to Infoline) No

Referral to Infoline: Yes No Refused Expired
Referral Date: _____

Report Submitted by: _____ **Title:** _____ **Date:** _____ **Reporting Hospital:** _____

CENTRAL NERVOUS SYSTEM

Anencephalus (Q00.0-Q00.1)

- Q00.0-Anencephaly
- Q00.1-Craniorachischisis

Microcephalus

- Q02-Microcephaly

Spina Bifida

(Q05.0-Q05.9, Q07.01, Q07.03 w/o Q00.0-Q00.1)

- 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 (D)-Unspecified spina bifida with hydrocephalus
- Q05.5-Cervical spina bifida without hydrocephalus
- Q05.6-Thoracic spina bifida without hydrocephalus
- Q05.7-Lumbar spina bifida without hydrocephalus
- Q05.8-Sacral spina bifida without hydrocephalus
- Q05.9 (D)-Spina bifida, unspecified
- Q07.01-Arnold-Chiari syndrome with spina bifida
- Q07.03-Arnold-Chiari syndrome with spina bifida and hydrocephalus

Encephalocele (Q01.0-Q01.9)

- Q01.0-Frontal encephalocele
- Q01.1-Nasofrontal encephalocele
- Q01.2-Occipital encephalocele
- Q01.8-encephalocele of other sites
- Q01.9 (D)-encephalocele, unspiced

Holoprosencephaly

- Q04.2 Holoprosencephaly

If condition not listed, please specify:

- Q0 _____
- Q0 _____

EYE

Anophthalmia/micropthalmia (Q11.0-Q11.2)

- Q11.0- Cystic eyeball
- Q11.1- Other anophthalmos
- Q11.2 (D)- Micropthalmos
- Q12.0-Congenital cataract

If condition not listed, please specify:

- Q1 _____
- Q1 _____

EAR

Anotia/microtia (Q16.0, Q17.2)

- Q16.0-Congenital absence of (ear) auricle
- Q17.2-Microtia

If condition not listed, please specify:

- Q1 _____
- Q1 _____

CARDIOVASCULAR

Aortic Valve Stenosis

- Q23.0 (D)-Congenital stenosis of aortic valve

Atrial Septal Defect

- Q21.1 (D)-Atrial septal defect

Atrioventricular septal defect

- Q21.2-Atrioventricular septal defect

Coarctation of aorta

- Q25.1-Coarctation of aorta

Common truncus

- Q20.0 (D)-Common arterial truck

Double outlet right ventricle (DORV)

- Q20.1 (D)-Double outlet right ventricle

Ebstein's anomaly

- Q22.5-Ebstein's anomaly

Hyperplastic left heart syndrome

- Q23.4-Hypoplastic left heart syndrome

Interrupted Aortic Arch (IAA) (Q25.2 and Q25.4)

- Q25.2 (D)-Atresia of aorta
- Q25.4-Other congenital malformations of aorta

Pulmonary Valve Atresia and Stenosis (Q22.0 and Q22.1)

- Q22.0-Pulmonary valve atresia
- Q22.1-Congenital pulmonary valve stenosis

Single Ventricle

- Q20.4 (D)-Double inlet ventricle

Tetralogy of Fallot (TOF)

- Q21.3 (D)-Tetralogy of Fallot

Total anomalous pulmonary venus connection (TAPVC)

- Q26.2 (D)-Total anomalous pulmonary venous connection

Transposition of great arteries (TGA) (Q20.3 and Q20.5)

- Q20.3-Discordant ventriculoarterial connection
- Q20.5-Discordant atrioventricular connection

Tricuspid valve atresia and stenosis

- Q22.4 (D)-Congenital tricuspid stenosis

Ventricular septal defect

- Q21.0 (D)-Ventricular septal defect

If condition not listed, please specify:

- Q2 _____
- Q2 _____

Enter this information into the Maven Newborn Screening System, or

if needed fax to: Department of Public Health, Attn: Karin Davis, Family Health Section at 860-509-7720



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Referral to Infoline: Yes **Referral Date:** _____ No Refused Expired
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OROFACIAL

- Q30.0-Choanal atresia
- Cleft lip WITH cleft palate (Q37.0-Q37.9)**
- Q37.0**-Cleft hard palate with bilateral cleft lip
- Q37.1-Cleft hard palate with unilateral cleft lip
- Q37.2-Cleft soft palate with bilateral cleft lip
- Q37.3-Cleft soft palate with unilateral cleft lip
- Q37.4-Cleft hard and soft palate with bilateral cleft lip
- Q37.5-Cleft hard and soft palate with unilateral cleft lip
- Q37.8 (D)-Unspecified cleft palate with bilateral cleft lip
- Q37.9 (D)-Unspecified cleft palate with unilateral cleft lip
- Cleft lip alone without cleft palate (Q36.0-Q36.9)**
- Q36.0- Cleft lip, bilateral
- Q36.1- Cleft lip, median
- Q36.9 (D)- Cleft lip, unilateral
- Cleft palate alone (without cleft lip) (Q35.1-Q35.9)**
- 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 (D)-Cleft palate, unspecified

GASTROINTESTINAL

- Biliary atresia (Q44.2-Q44.3)**
- Atresia of bile ducts-Q44.2
- Congenital stenosis and stricture of bile ducts-Q44.3
- Esophageal atresia/tracheoesophageal fistula (Q39.0-Q39.4)**
- Atresia of esophagus without fistula-Q39.0
- Atresia of esophagus w/tracheo-esophageal fistula -Q39.1 (D)
- Congenital tracheo-esophageal fistula without atresia-Q39.2
- Congenital stenosis and stricture of esophagus-Q39.3
- Esophageal web-Q39.4
- Rectal and large intestinal atresia/stenosis (Q42.0-Q42.9)**
- Congenital absence, atresia, & stenosis of rectum with fistula-Q42.0
- Congenital absence, atresia, & 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-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 (D)
- Small intestine atresia/stenosis (Q41.0-Q41.9)**
- 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 parts of small intestine-Q41.8
- Congenital absence, atresia, and stenosis of small intestine, part unspecified-Q41.9
- If condition not listed, please specify:
- Q4 _____
- Q4 _____

GENITOURINARY

- Bladder exstrophy (Q64.10, Q64.19)**
- Q64.10-Exstrophy of urinary bladder, unspecified
- Q64.19 (D)-Other exstrophy of urinary bladder
- Cloacal exstrophy**
- Q64.12-Cloacal exstrophy of urinary bladder
- Congenital posterior**
- Q64.2-Congenital posterior urethral valves
- Hypospadias (Q54.0-Q54.9 excluding Q54.4)**
- Q54.0-Hypospadias, balanic
- Q54.1-Hypospadias, penile
- Q54.2-Hypospadias, penoscrotal
- Q54.3-Hypospadias, perineal
- Q54.8-Other hypospadias
- Q54.9 (D)-Hypospadias, unspecified
- Renal agenesis/hypoplasia (Q60.0-Q60.6)**
- Q60.0-Renal agenesis, unilateral
- Q60.1-Renal agenesis, bilateral
- Q60.2 (D)-Renal agenesis, unspecified
- Q60.3-Renal hypoplasia, unilateral
- Q60.4-Renal hypoplasia, bilateral
- Q60.5-Renal hypoplasia, unspecified
- Q60.6-Potter's syndrome
- If not listed, please specify:
- Q _____
- Q _____

CHROMOSOMAL ABNORMALITIES

- Deletion 22q11**
- Q93.81-DiGeorge syndrome-deletion 22q11 Velo-cardio-facial syndrome
- Trisomy 13-Patau's syndrome (Q91.4-Q91.7)**
- Q91.4-Trisomy 13, non-mosaicism (meiotic nondisjunction)
- Q91.5-Trisomy 13, mosaicism (mitotic nondisjunction)
- Q91.6-Trisomy 13, translocation
- Q91.7 (D)- Trisomy 13, unspecified
- Trisomy 18-Edward syndrome (Q91.0-Q91.3)**
- Q91.0-Trisomy 18, non-mosaicism (meiotic nondisjunction)
- Q91.1-Trisomy 18, mosaicism (mitotic nondisjunction)
- Q91.2-Trisomy 18, translocation
- Q91.3 (D)-Trisomy 18, unspecified
- Trisomy 21-Down syndrome (Q90.0-Q90.9)**
- Q90.0-Trisomy 21, non-mosaicism (meiotic nondisjunction)
- Q90.1-Trisomy 21, mosaicism (mitotic nondisjunction)
- Q90.2-Trisomy 21, translocation
- Q90.9 (D)-Down syndrome, unspecified
- Turner Syndrome (Q96.0-96.9)**
- Q96.0-Karyotype 45,X
- Q96.1-Karyotype 45,X iso (Xq)
- Q96.2-Karyotype 45,X w/abnormal sex chromosome, except iso (Xq)
- Q96.3-Mosaicism, 45,X/46,XX or XY
- Q96.4-Mosaicism, 45,X/other cell line(s) w/abnormal sex chromosome
- Q96.8-Other variants of Turner's syndrome
- Q96.9 (D)-Turner's syndrome, unspecified
- If condition not listed, please specify:
- Q9 _____

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MUSCULOSKELETAL

- Clubfoot** (Q66.0, Q66.89)
 Q66.0-congenital talipes equinovarus
 Q66.89-Other specified congenital deformities of feet
Craniosynostosis
 Q75.0-Craniosynostosis
Diaphragmatic hernia (Q79.0, Q79.1)
 Q79.0 (D)-Congenital diaphragmatic hernia
 Q79.1-Other congenital malformations of diaphragm
Gastroschisis
 Q79.3-Gastroschisis
Limb deficiencies (reduction defects) (Q71.0-Q71.9)
 Q71.0-Congenital complete absence of unspecified upper limb
 Q71.01-Congenital complete absence of right upper limb
 Q71.02-Congenital complete absence of left upper limb
 Q71.03-Congenital complete absence of upper limb, bilateral
 Q71.10 (D)-Congenital absence of unspecified upper arm and forearm with hand present
 Q71.11-Congenital absence of right upper arm and forearm with hand present
 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
 Q71.20-Congenital absence of both forearm and hand, unsepcified upper limb
 Q71.21-Congenital absence of both forearm and hand, right upper limb
 Q71.22-Congenital absence of both forearm and hand, left upper limb
 Q71.23-Congenital absence of both forearm and hand, bilateral
 Q71.30-Congenital absence of unspecified hand and finger
 Q71.31-Congenital absence of right hand and finger
 Q71.32-Congenital absence of left hand and finger
 Q71.33-Congenital absence of hand and finger, bilateral
 Q71.40 (D)-Longitudinal reduction defect of unspecified radius
 Q71.41-Longitudinal reduction defect of right radius
 Q71.42-Longitudinal reduction defect of left radius
 Q71.43-Longitudinal reduction defect of radius, bilateral
 Q71.50 (D)-Longitudinal reduction defect of unspecified ulna
 Q71.51-Longitudinal reduction defect of right ulna
 Q71.52-Longitudinal reduction defect of left ulna
 Q71.53-Longitudinal reduction defect of ulna, bilateral
 Q71.60-Lobster-claw hand, unspecified hand
 Q71.61-Lobster-claw right hand
 Q71.62-Lobster-claw left hand
 Q71.63-Lobster-claw hand, bilateral
 Q71.90 (D)-Unspecified reduction defect of unspecified 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
 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
 Q71.899 (D)-Other reduction defects of unspecified upper limb

(Q72.0-Q72.9)

- Q72.00-Congenital complete absence of unspecified lower limb
 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 (D)- Congenital absence of unspecified thigh and lower leg with foot present
 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 thigh and lower leg with foot present, bilateral
 Q72.20-Congenital absence of both lower leg and foot, unspecified lower limb
 Q72.21-Congenital absence of both lower leg and foot, right 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, bilateral
 Q72.30-Congenital absence of unspecified foot and toe(s)
 Q72.31-Congenital absence of right foot and toe(s)
 Q72.32-Congenital absence of left foot and toe(s)
 Q72.33-Congenital absence of foot and toe(s), bilateral
 Q72.40 (D)-Longitudinal reduction defect of unspecified femur
 Q72.41-Longitudinal reduction defect of right femur
 Q72.42-Longitudinal reduction defect of left femur
 Q72.43-Longitudinal reduction defect of femur, bilateral
 Q72.50 (D)-Longitudinal reduction defect of unspecified tibia
 Q72.51-Longitudinal reduction defect of unspecified tibia
 Q72.52-Longitudinal reduction defect of unspecified tibia
 Q72.53-Longitudinal reduction defect of unspecified tibia
 Q72.60 (D)-Longitudinal reduction defect of unspecified fibula
 Q72.61-Longitudinal reduction defect of right fibula
 Q72.62-Longitudinal reduction defect of left fibula
 Q72.63-Longitudinal reduction defect of fibula, bilateral
 Q72.70-Split foot, unspecified lower limb
 Q72.71-Split foot, right lower limb
 Q72.72-Split foot, left lower limb
 Q72.73-Split foot, 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
 Q72.899 (D)-Other reduction defects of unspecified lower limb
 Q72.90 (D)-Unspecified reduction defect of unspecified lower limb
 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-Q73.8)

- Q73.0-Congenital absence of unspecified limb(s)
 Q73.1-Phocomelia, unspecified limb(s)
 Q73.8 (D)-Other reduction defects of unspecified limb(s)
Omphalocele
 Q79.2-Exomphalos
 If condition not listed, please specify:
 Q7 _____
 Q7 _____

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