State of Connecticut Department of Public Health Birth Defect Registry Reporting Form (Revised 4/27/18)

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

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

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

Is a birth defect present in this child? ☐ **Yes** ☐ **No** **Referral to CDHI?** ☐ **Yes** **Date:** _____ ☐ **No** ☐ **Refused** ☐ **Expired**

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

How was PCP notified of birth defect(s): ☐ **we are the PCP** ☐ **Discharge Summary** ☐ **Electronic Health Record (EHR-networked)** ☐ **Electronic Medical Record (EMR-single record)** ☐ **Email/Fax** ☐ **Phone Call** ☐ **Other – specify** _____

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, unspecified

Holoprosencephaly
- ☐Q04.2 Holoprosencephaly

If condition not listed, please specify:
- ☐Q0____ _____
- ☐Q0____ _____

EYE

Anophtlalmia/microphthalmia (Q11.0-Q11.2)
- ☐Q11.0- Cystic eyeball
- ☐Q11.1- Other anophthalmos
- ☐Q11.2 (**D**)- Microphthalmos
- ☐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

Interupted 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, CFHPS Section at 860-509-7720

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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

Bilary 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-Hypospaidas, 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____ _____

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, unspecified 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 & 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 unspec 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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