Appendix 5.2
6-Digit CDC Codes

BIRTH DEFECTS AND GENETIC DISEASES BRANCH 6-DIGIT CODE

For Reportable Congenital Anomalies


Code modifications developed by Division of Birth Defects and Developmental Disabilities, National Center on Birth Defects and Developmental Disabilities, Centers for Disease Control and Prevention, Public Health Service, U.S. Department of Health and Human Services, Atlanta, Georgia 30333

Doc. No. 6digit88
Version 06/04
Replaces Version 06/93
Explanation of 6-Digit Code

6th Digit Code - Master

- **.000** Blank
- **.001** Left Only
- **.002** Right Only
- **.003** Unilateral Unspecified
- **.004** Bilateral
- **.005**
- **.006**
- **.007** Possible, Probable, Borderline, or Rule Out;
  Defects only diagnosed prenatally should be coded with the last digit 8
  when the prenatal diagnosis is not definitive.
- **.009** Not Otherwise Specified (NOS)

Notes:

An asterisk (*) beside a disease code indicates that the code was created by
CDC.

A pound symbol (#) beside a disease code indicates that the condition or
defect is listed on the MACDP Exclusion List.

A check (T) beside a disease code indicates that an addition/revision was
made since the last printing of the Procedure Manual. Use of the code should
be according to the exclusion list criteria.

The abbreviations NEC and NOS used in this code are defined as not elsewhere
classified and as not otherwise specified, respectively.
CONGENITAL ANOMALIES

Anencephalus and Similar Anomalies

740.0 Anencephalus
- 740.000 Absence of brain
- 740.010 Acrania
- 740.020 Anencephaly
- 740.030 Hemianencephaly, hemicephaly
- 740.080 Other

740.1 Craniorachischisis
- 740.100 Craniorachischisis

740.2 Iniencephaly
- 740.200 Closed iniencephaly
- 740.210 Open iniencephaly
- 740.290 Unspecified iniencephaly

741 Spina Bifida
Includes: Spina bifida aperta (open lesions) myelocele rachischisis
Spina bifida cystica (closed lesions) meningocoele meningomyelocele myelomeningocele
Excludes: Spina bifida occulta (see 756.100) craniorachischisis (see 740.100)

741.0 Spina Bifida with Hydrocephalus
- 741.000 Spina bifida aperta, any site, with hydrocephalus
- 741.010 Spina bifida cystica, any site, with hydrocephalus and Arnold-Chiari malformation
  Arnold-Chiari malformation, NOS
- 741.020 Spina bifida cystica, any site, with stenosed aqueduct of Sylvius
- 741.030 Spina bifida cystica, cervical, with unspecified hydrocephalus
  Spina bifida cystica, cervical, with hydrocephalus but without mention of Arnold-Chiari malformation or aqueduct stenosis
- 741.040 Spina bifida cystica, thoracic, with unspecified hydrocephalus, no mention of Arnold-Chiari
- 741.050 Spina bifida cystica, lumbar, with unspecified hydrocephalus, no mention of Arnold-Chiari
- 741.060 Spina bifida cystica, sacral, with unspecified hydrocephalus, no mention of Arnold-Chiari
- 741.070 Spina bifida of any site with hydrocephalus of late onset
- 741.080 Other spina bifida, meningocele of specified site with hydrocephalus
- 741.085 Spina bifida, meningocele, cervicothoracic, with
hydrocephalus
741.086 Spina bifida, meningocele thoracolumbar, with hydrocephalus
741.087 Spina bifida, meningocele, lumbosacral with hydrocephalus
741.090 Spina bifida of any unspecified type with hydrocephalus

741.9 Spina bifida without mention of hydrocephalus
741.900 Spina bifida (aperta), without hydrocephalus
741.910 Spina bifida (cystica), cervical, without hydrocephalus
741.920 Spina bifida (cystica), thoracic, without hydrocephalus
741.930 Spina bifida (cystica), lumbar, without hydrocephalus
741.940 Spina bifida (cystica), sacral, without hydrocephalus
741.980 Spina bifida, other specified site, without hydrocephalus
Includes: cervicothoracic, thoracolumbar, lumbosacral
741.985 Lipomyelomeningocele
741.990 Spina bifida, site unspecified, without hydrocephalus
(myelocoele, myelomeningocele, meningomyelocele)

742 Other Congenital Anomalies of Nervous System

742.0 Encephalocele
742.000 Occipital encephalocele
742.080 Other encephalocele of specified site (includes midline defects)
742.085 Frontal encephalocele
742.086 Parietal encephalocele
742.090 Unspecified encephalocele

742.1 Microcephalus
742.100 Microcephalus

742.2 Reduction deformities of brain
742.200 Anomalies of cerebrum
742.210 Anomalies of corpus callosum
742.220 Anomalies of hypothalamus
742.230 Anomalies of cerebellum
742.240 Agyria and lissencephaly
742.250 Microgyria, polymicrogyria
742.260 Holoprosencephaly
742.270 Arrhinencephaly
742.280 Other specified reduction defect of brain
742.290 Unspecified reduction defect of brain

742.3 Congenital hydrocephalus
Excludes: hydrocephalus with any condition in 741.9 (use 741.0)
742.300 Anomalies of aqueduct of Sylvius
742.310 Atresia of foramina of Magendie and Luschka
Dandy-Walker syndrome
742.320 Hydranencephaly
742.380 Other specified hydrocephaly
Includes: communicating hydrocephaly
# 742.385 Hydrocephalus secondary to intraventricular hemorrhage (IVH) or CNS bleed
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>742.390</td>
<td>Unspecified hydrocephaly, NOS</td>
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<tr>
<td>742.400</td>
<td>Enlarged brain and/or head megalencephaly</td>
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<tr>
<td>742.410</td>
<td>Enlarged brain and/or head macrocephaly</td>
</tr>
<tr>
<td>742.420</td>
<td>Porencephaly</td>
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<tr>
<td>742.430</td>
<td>Porencephalic cysts</td>
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<tr>
<td>742.440</td>
<td>Cerebral cysts</td>
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<tr>
<td>742.450</td>
<td>Other specified anomalies of brain</td>
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<tr>
<td>742.480</td>
<td>Includes: cortical atrophy</td>
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<tr>
<td>742.485</td>
<td>Cranial nerve defects</td>
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<tr>
<td>742.486</td>
<td>Small brain</td>
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<tr>
<td>742.500</td>
<td>Amyelia</td>
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<tr>
<td>742.510</td>
<td>Hypoplasia and dysplasia of spinal cord</td>
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<tr>
<td>742.520</td>
<td>Diastematomyelia</td>
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<tr>
<td>742.530</td>
<td>Other cauda equina anomalies</td>
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<tr>
<td>742.540</td>
<td>Hydromyelia</td>
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<tr>
<td>742.580</td>
<td>Other specified anomalies of spinal cord and membranes</td>
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<tr>
<td>742.800</td>
<td>Jaw-winking syndrome</td>
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<tr>
<td>742.810</td>
<td>Familial dysautonomia</td>
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<tr>
<td>742.880</td>
<td>Other specified anomalies of nervous system</td>
</tr>
<tr>
<td>742.900</td>
<td>Brain, unspecified anomalies</td>
</tr>
<tr>
<td>742.910</td>
<td>Spinal cord, unspecified anomalies</td>
</tr>
<tr>
<td>742.990</td>
<td>Nervous system, unspecified anomalies</td>
</tr>
</tbody>
</table>
743 Congenital Anomalies of Eye

743.000 Anophthalmos
    agenesis of eye
    cryptophthalmos
743.100 Microphthalmos, small eyes
    aplasia of eye
    hypoplasia of eye
    dysplasia of eye
    rudimentary eye

743.2 Buphthalmos

743.200 Buphthalmos
    congenital glaucoma
    hydrophthalmos
743.210 Enlarged eye, NOS
743.220 Enlarged cornea
    keratoglobus
    congenital megalocornea

743.3 Congenital cataract and lens anomalies

743.300 Absence of lens
    congenital aphakia
743.310 Spherical lens
    Spherophakia
743.320 Cataract, NOS
743.325 Cataract, anterior polar
743.326 Cataract, other specified
743.330 Displaced lens
743.340 Coloboma of lens
743.380 Other specified lens anomalies
743.390 Unspecified lens anomalies

743.4 Coloboma and other anomalies of anterior segments

743.400 Corneal opacity
743.410 Other corneal anomalies
    Excludes: megalocornea (use 743.220)
743.420 Absence of iris
    aniridia
743.430 Coloboma of iris
743.440 Other anomalies of iris
    polycoria
    ectopic pupil
    Peter's anomaly

# Excludes: brushfield spots (use 743.800)

743.450 Blue sclera
    # If <36 weeks gestation, code only if another reportable
defect is present.
    Always code if ≥36 weeks gestation.
743.480 Other specified colobomas and anomalies of anterior segments
743.490 Rieger's anomaly
743.490 Unspecified colobomas and anomalies of anterior eye segments
743.5 Congenital anomalies of posterior segment

743.500 Specified anomalies of vitreous humour
743.510 Specified anomalies of retina
congenital retinal aneurysm
Excludes: Stickler syndrome (use 759.860)
743.520 Specified anomalies of optic disc
hypoplastic optic nerve
coloboma of the optic disc
743.530 Specified anomalies of choroid
743.535 Coloboma of choroid
743.580 Other specified anomalies of posterior segment of eye
743.590 Unspecified anomalies of posterior segment of eye

743.6 Congenital anomalies of eyelids, lacrimal system, and orbit

743.600 Blepharoptosis
congenital ptosis
743.610 Ectropion
743.620 Entropion
# 743.630 Other anomalies of eyelids
absence of eyelashes
long eyelashes
weakness of eyelids
T #
  fused eyelids (exclude if <25 weeks gestation unless another
  reportable defect is present)
743.635 Blepharophimosis
small or narrow palpebral fissures
743.636 Coloboma of the eyelids
743.640 Absence or agenesis of lacrimal apparatus
absence of punctum lacrimale
# 743.650 Stenosis or stricture of lacrimal duct
743.660 Other anomalies of lacrimal apparatus (e.g., cyst)
743.670 Anomalies of orbit

743.8 Other specified anomalies of eye

# 743.800 Other specified anomalies of eye
Includes: exophthalmos
epicanthal folds
antimongoloid slant
upward eye slant
Brushfield spots
Excludes: congenital nystagmus (use 379.500)
retinitis pigmentosa (use 362.700)
ocular albinism (use 270.200)
wide spaced eyes, hypertelorism (use 756.085)
* 743.810 Epibulbar dermoid cyst

743.9 Unspecified anomalies of eye

743.900 Unspecified anomalies of eye
congenital: of eye (any part)
anomaly, NOS
deformity, NOS
744 Congenital Anomalies of Ear, Face, and Neck

744.0 Anomalies of ear causing impairment of hearing

- 744.000 Absence or stricture of auditory canal
- 744.010 Absence of auricle (pinna) absence of ear, NOS
- 744.020 Anomaly of middle ear fusion of ossicles
- 744.030 Anomaly of inner ear
  Includes: congenital anomaly of membranous labyrinth organ of Corti
- 744.090 Unspecified anomalies of ear with hearing impairment
  Includes: congenital deafness, NOS

744.1 Accessory auricle

- # 744.100 Accessory auricle Polyotia
- # 744.110 Preauricular appendage, tag, or lobule (in front of ear canal)
- # 744.120 Other appendage, tag, or lobule include papillomas, ear tags

744.2 Other specified anomalies of ear

- 744.200 Macrotia (enlarged pinna)
- 744.210 Microtia (hypoplastic pinna and absence or stricture of external auditory meatus)
- 744.220 Bat ear
- T # 744.230 Other misshapen ear
  pointed ear elfin pixie-like lop ear cauliflower ear cleft in ear malformed ear absent or decreased cartilage
- 744.240 Misplaced ears
- # 744.245 Low set ears
- # 744.246 Posteriorly rotated ears
- 744.250 Absence or anomaly of eustachian tube
- 744.280 Other specified anomalies of ear (see also 744.230)
  Excludes: Darwin's tubercle

744.3 Unspecified anomalies of ear

- 744.300 Unspecified anomalies of ear
  Congenital: ear (any part) anomaly, deformity, NOS

744.4 Branchial cleft, cyst, or fistula; preauricular sinus

- 744.400 Branchial cleft, sinus, fistula cyst, or pit
- # 744.410 Preauricular sinus, cyst, or pit
- 744.480 Other branchial cleft anomalies
  Includes: dermal sinus of head
- # 744.500 Webbing of neck
Includes: pterygium colli, redundant neck skin folds

744.8 Other unspecified anomalies of face and neck

744.800 Macrostomia (large mouth)
744.810 Microstomia (small mouth)
# 744.820 Macrocheilia (large lips)
# 744.830 Microcheilia (small lips)
744.880 Other specified anomalies of face/neck

744.9 Unspecified anomalies of face and neck

# 744.900 Congenital anomaly of neck, NOS
Includes: short neck
744.910 Congenital anomaly of face, NOS
Abnormal facies
745 Bulbus Cordis Anomalies and Anomalies of Cardiac Septal Closure

745.0 Common truncus (see 747.200 for pseudotruncus)
   745.000 Persistent truncus arteriosus
       absent septum between aorta and pulmonary artery
   745.010 Aortic septal defect
       Includes: aortopulmonary window
       Excludes: atrial septal defect (use 745.590)

745.1 Transposition of great vessels
   745.100 Transposition of great vessels, complete (no VSD)
   745.110 Transposition of great vessels, incomplete (w/ VSD)
       Taussig-Bing syndrome
   745.120 Corrected transposition of great vessels,
       L-transposition, ventri in version
       Excludes: dextrocardia (use 746.800)
   745.180 Other specified transposition of great vessels
       Includes: double outlet right ventricle
   745.190 Unspecified transposition of great vessels

745.2 Tetralogy of Fallot
   745.200 Fallot's tetralogy
   745.210 Fallot's pentalogy
       Fallot's tetralogy plus ASD

745.3 Single ventricle
   745.300 Single ventricle
       Common ventricle
       Cor triloculare biatriatum

745.4 Ventricular septal defect
   745.400 Roger's disease
   745.410 Eisenmenger's syndrome
   745.420 Gerbode defect
   T 745.480 Other specified ventricular septal defect (includes membranous,
       perimembranous, muscular, mid-muscular, crystalline, sub-
       crystalline, subarterial, inlet, conoventricular)
   745.490 VSD (ventricular septal defect), NOS
       Excludes: common atrioventricular canal type (use 745.620)
   745.498 Probable VSD

745.5 Ostium secundum type atrial septal defect
   T # 745.500 Nonclosure of foramen ovale, NOS
       Patent foramen ovale (PFO)
       1) Always code if ≥36 weeks of gestation and defect last noted
          at ≥6 weeks of age.
       2) If ≥36 weeks gestation and defect last noted <6 weeks of age,
          code only if another reportable defect is present.
       3) Never code if <36 weeks gestation regardless of presence of
          other defects.
   745.510 Ostium (septum) secundum defect
745.520  Lutembacher's syndrome
745.580  Other specified atrial septal defect
745.590  ASD (atrial septal defect), NOS
          Auricular septal defect, NOS
          Partial foramen ovale
          PFO vs. ASD

745.6  Endocardial cushion defects

745.600  Ostium primum defects
745.610  Single common atrium, cor triloculare biventriculare
745.620  Common atrioventricular canal
          with ventricular septal defect (VSD)
745.630  Common atrioventricular canal
745.680  Other specified cushion defect
745.690  Endocardial cushion defect, NOS

745.7  Cor biloculare

745.700  Cor biloculare

745.8  Other specified defects of septal closure

745.800  Other specified defects of septal closure

745.9  Unspecified defect of septal closure

745.900  Unspecified defect of septal closure

746  Other Congenital Anomalies of Heart

746.0  Anomalies of pulmonary valve

746.000  Atresia, hypoplasia of pulmonary valve
          See 746.995 if valve is not specified
          (e.g., "pulmonary atresia");
746.010  Stenosis of pulmonary valve
          See 746.995 if valve not specified
          (e.g., "pulmonary stenosis");
          # Excludes: pulmonary infundibular
          stenosis (use 746.830)
746.020  Pulmonary valve insufficiency or regurgitation, congenital
          Code cases designated as 'mild', minimal', 'trivial', or
          'physiologic' only if another reportable defect is present.
          Code all other degrees of insufficiency or regurgitation,
          including those where the degree is not specified, regardless
          of whether another reportable defect is present.
746.080  Other specified anomalies of pulmonary valve
          # Excludes: pulmonary infundibular
          stenosis (use 746.830)
746.090  Unspecified anomaly of pulmonary valve

746.1  Tricuspid atresia and stenosis

746.100  Tricuspid atresia, stenosis, hypoplasia
746.105  Tricuspid valve insufficiency or regurgitation, congenital
          Code cases designated as 'mild', minimal', 'trivial', or
          'physiologic' only if another reportable defect is present.
Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present. Exclude: Ebstein's anomaly (code as 746.200)

746.2 Ebstein's anomaly

746.200 Ebstein's anomaly

746.3 Congenital stenosis of aortic valve

746.300 Congenital stenosis of aortic valve
Includes: congenital aortic stenosis
subvalvular aortic stenosis
Excludes: supravalvular aortic stenosis (747.220)

746.4 Congenital insufficiency of aortic valve

T # 746.400 Aortic valve insufficiency or regurgitation, congenital bicuspid aortic valve.
Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present.
Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.
* 746.480 Other specified anomalies of the aortic valves
Includes: aortic valve atresia
Excludes: supravalvular aortic stenosis (747.220)
* 746.490 Unspecified anomalies of the aortic valves

746.5 Congenital mitral stenosis

746.500 Congenital mitral stenosis
746.505 Absence, atresia, or hypoplasia of mitral valve

746.6 Mitral valve insufficiency or regurgitation, congenital

T # 746.600 Mitral valve insufficiency or regurgitation, congenital
Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present.
Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.

746.7 Hypoplastic left heart syndrome

746.700 Hypoplastic left heart syndrome
Atresia, or marked hypoplasia of the ascending aorta and defective development of left ventricle (with mitral valve atresia)

746.8 Other specified anomalies of the heart

746.800 Dextrocardia without situs inversus (situs solitus)
Dextrocardia with no mention of situs inversus
Excludes: dextrocardia with situs inversus (use 759.300)
746.810 Levocardia
746.820 Cor triatriatum
746.830 Pulmonary infundibular (subvalvular) stenosis
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>746.840</td>
<td>Trilogy of Fallot</td>
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<tr>
<td>746.850</td>
<td>Anomalies of pericardium</td>
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<td>746.860</td>
<td>Anomalies of myocardium cardiomegaly, congenital, NOS</td>
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<td>cardiomypathy, congenital</td>
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<td></td>
<td>cardiomypathy, hypertrophic</td>
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<td>746.870</td>
<td>Congenital heart block</td>
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<td>746.880</td>
<td>Other specified anomalies of heart</td>
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<td>Includes: ectopia (ectopic) cordis (mesocardia),</td>
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<td>conduction defects, NOS</td>
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<td>746.881</td>
<td>Hypoplastic left ventricle</td>
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<td>Excludes: hypoplastic left heart syndrome (746.700)</td>
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<td>746.882</td>
<td>Hypoplastic right heart (ventricle)</td>
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<td>Uhl's disease</td>
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<td>746.883</td>
<td>Hypoplastic ventricle, NOS</td>
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<td>746.885</td>
<td>Anomalies of coronary artery or sinus</td>
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<td>746.886</td>
<td>Ventricular hypertrophy (right or left)</td>
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<td>746.887</td>
<td>Other defects of the atria</td>
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<td>Excludes: congenital Wolfe-Parkinson-White</td>
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<td>(use 426.705)</td>
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<td>rhythm anomalies (use 426.-, 427.-)</td>
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<td>746.9</td>
<td>Unspecified anomalies of heart</td>
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<td>746.900</td>
<td>Unspecified anomalies of heart valves</td>
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<td>746.910</td>
<td>Anomalous bands of heart</td>
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<td>746.920</td>
<td>Acyanotic congenital heart disease, NOS</td>
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<tr>
<td>746.930</td>
<td>Cyanotic congenital heart disease, NOS</td>
</tr>
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<td>Blue baby</td>
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<tr>
<td>746.990</td>
<td>Unspecified anomaly of heart:</td>
</tr>
<tr>
<td></td>
<td>Includes: congenital heart disease (CHD)</td>
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<tr>
<td>746.995</td>
<td>&quot;Pulmonic&quot; or &quot;pulmonary&quot; atresia, stenosis, or hypoplasia, NOS</td>
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</tbody>
</table>

747 Other Congenital Anomalies of Circulatory System

<table>
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<tr>
<th>Code</th>
<th>Description</th>
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<tbody>
<tr>
<td>T # 747.000</td>
<td>Patent ductus arteriosus (PDA)</td>
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<td>1)Always code if ≥36 weeks of gestation and defect last noted at ≥6 weeks of age.</td>
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<tr>
<td></td>
<td>2)If ≥36 weeks gestation and defect last noted &lt;6 weeks of age, code only if the PDA was treated (e.g. by ligation or indomethacin) or if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td>3)Never code if &lt;36 weeks gestation or if treated with prostaglandins regardless of gestational age.</td>
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<tr>
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<td>(See PDA Tree Appendix)</td>
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<td>747.008</td>
<td>Probable PDA</td>
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747.1 Coarctation of aorta

<table>
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<tbody>
<tr>
<td>747.100</td>
<td>Preductal (proximal) coarctation of aorta</td>
</tr>
<tr>
<td>747.110</td>
<td>Postductal (distal) coarctation of aorta</td>
</tr>
<tr>
<td>747.190</td>
<td>Unspecified coarctation of aorta</td>
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</tbody>
</table>

747.2 Other anomalies of aorta

<table>
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<tr>
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<th>Description</th>
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<tbody>
<tr>
<td>747.200</td>
<td>Atresia of aorta</td>
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<td>absence of aorta</td>
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<tr>
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<td>pseudotruncus arteriosus</td>
</tr>
<tr>
<td>747.210</td>
<td>Hypoplasia of aorta</td>
</tr>
</tbody>
</table>
tubular hypoplasia of aorta
747.215 Interrupted aortic arch
747.220 Supra-aortic stenosis (supravalvular)
   Excludes: aortic stenosis, congenital (see 746.300)
747.230 Persistent right aortic arch
747.240 Aneurysm of sinus of Valsalva
747.250 Vascular ring (aorta)
   double aortic arch
   Includes: vascular ring compression of trachea
747.260 Overriding aorta
747.270 Congenital aneurysm of aorta
   congenital dilatation of aorta
747.280 Other specified anomalies of aorta
747.290 Unspecified anomalies of aorta

747.3 Anomalies of pulmonary artery

747.300 Pulmonary artery atresia, absence or agenesis
   Use 746.995 if artery or valve is not specified
747.310 Pulmonary artery atresia with septal defect
747.320 Pulmonary artery stenosis
   Use 746.995 if artery or valve is not specified
747.325 Peripheral pulmonary artery stenosis
   Includes: peripheral pulmonic stenosis (PPS)
   peripheral pulmonic stenosis (PPS) murmur only if documented by echocardiogram
747.330 Aneurysm of pulmonary artery
747.340 Pulmonary arteriovenous malformation or aneurysm
747.380 Other specified anomaly of pulmonary artery
   Includes: pulmonary artery hypoplasia
747.390 Unspecified anomaly of pulmonary artery

747.4 Anomalies of great veins

747.400 Stenosis of vena cava (inferior or superior)
747.410 Persistent left superior vena cava
747.420 (TAPVR) Total anomalous pulmonary venous return
747.430 Partial anomalous pulmonary venous return
747.440 Anomalous portal vein termination
747.450 Portal vein – hepatic artery fistula
747.480 Other specified anomalies of great veins
747.490 Unspecified anomalies of great veins

747.5 Absence or hypoplasia of umbilical artery

# 747.500 Single umbilical artery

747.6 Other anomalies of peripheral vascular system

747.600 Stenosis of renal artery
747.610 Other anomalies of renal artery
747.620 Arteriovenous malformation (peripheral)
   Excludes: pulmonary (747.340)
   cerebral (747.800)
   retinal (743.510)
747.630 Congenital phlebectasia
    congenital varix
747.640 Other anomalies of peripheral arteries
    Includes: aberrant subclavian artery
747.650 Other anomalies of peripheral veins
    Excludes: Budd-Chiari - occlusion of hepatic vein (use 453.000)
T 747.680 Other anomalies of peripheral vascular system
    Includes: primary pulmonary artery hypertension ONLY if it occurs with another reportable defect
# 747.690 Unspecified anomalies of peripheral vascular system

747.8 Other specified anomalies of circulatory system

747.800 Arteriovenous (malformation) aneurysm of brain
747.810 Other anomalies of cerebral vessels
    Includes: vein of Galen
747.880 Other specified anomalies of circulatory system
    Excludes: congenital aneurysm:
        coronary (746.880)
        peripheral (747.640)
        pulmonary (747.330)
        retinal (743.510)
        ruptured cerebral arteriovenous aneurysm (430.000)
        ruptured cerebral aneurysm (430.000)

747.9 Unspecified anomalies of circulatory system

747.900 Unspecified anomalies of circulatory system
748  Congenital Anomalies of Respiratory System

748.0 Choanal atresia
    748.000 Choanal atresia
        atresia of nares, anterior or posterior
        congenital stenosis

748.1 Other anomalies of nose
    748.100 Agenesis or underdevelopment of nose
    748.110 Accessory nose
    748.120 Fissured, notched, or cleft nose
    748.130 Sinus wall anomalies
    748.140 Perforated nasal septum
    # 748.180 Other specified anomalies of nose
        flat bridge of nose
        wide nasal bridge
        small nose and nostril
        absent nasal septum
    748.185 Tubular nose, single nostril, proboscis
    748.190 Unspecified anomalies of nose
        Excludes: congenital deviation of the nasal
        septum (use 754.020)

748.2 Web of larynx
    748.205 Web of larynx-glottic
    748.206 Web of larynx-subglottic
    748.209 Web of larynx-NOS

748.3 Other anomalies of larynx, trachea, and bronchus
    748.300 Anomalies of larynx and supporting cartilage
    T 748.310 Congenital subglottic stenosis – Never code if chart states
        the condition was acquired or secondary to endotracheal (ET)
        intubation or ventilation
    748.330 Other anomalies of trachea
        Excludes: vascular ring compression of the
        trachea (use 747.250)
    748.340 Stenosis of bronchus
    748.350 Other anomalies of bronchus
    748.360 Congenital laryngeal stridor, NOS
    748.380 Other specified anomalies of larynx and bronchus
    748.385 Cleft larynx, laryngotracheoesophageal cleft
    748.390 Unspecified anomalies of larynx, trachea, and bronchus

748.4 Congenital cystic lung
    748.400 Single cyst, lung or lung cyst
    748.410 Multiple cysts, lung
        Polycystic lung
    748.420 Honeycomb lung
    748.480 Other specified congenital cystic lung
748.5 Agenesis or aplasia of lung

748.500 Agenesis or aplasia of lung

748.510 Hypoplasia of lung; Pulmonary hypoplasia

T Exlude if isolated defect in infants <36 weeks gestation.

748.520 Sequestration of lung

748.580 Other specified dysplasia of lung

Fusion of lobes of lung

* 748.590 Unspecified dysplasia of lung

748.6 Other anomalies of lung

748.600 Ectopic tissues in lung

748.610 Bronchiectasis

748.620 Accessory lobe of lung

748.625 Bilobar right lung or right lung with left lung bronchial pattern

748.690 Other and unspecified anomalies of lung

748.8 Other specified anomalies of respiratory system

748.800 Anomaly of pleura

748.810 Congenital cyst of mediastinum

748.880 Other specified respiratory system anomalies

Includes: congenital lobar emphysema

lymphangiectasia of lungs

748.9 Unspecified anomalies of respiratory system

748.900 Unspecified anomalies of respiratory system

Absence of respiratory organ, NOS

Anomaly of respiratory system, NOS
749  Cleft Palate and Cleft Lip

749.0  Cleft palate alone
   (If description of condition includes Pierre Robin sequence, 
   use additional code, 524.080)

749.000  Cleft hard palate, unilateral
749.010  Cleft hard palate, bilateral
749.020  Cleft hard palate, central
749.030  Cleft hard palate, NOS
749.040  Cleft soft palate, alone unilateral
749.050  Cleft soft palate, alone bilateral
749.060  Cleft soft palate, alone central
749.070  Cleft soft palate, alone, NOS
749.080  Cleft uvula
749.090  Cleft palate, NOS
   palatoschisis

749.1  Cleft lip alone
   Includes: alveolar ridge cleft
   cleft gum
   harelip
749.100  Cleft lip, unilateral
749.110  Cleft lip, bilateral
749.120  Cleft lip, central
749.190  Cleft lip, NOS (fused lip)
   cleft gum

749.2  Cleft lip with cleft palate

749.200  Cleft lip, unilateral, with any cleft palate
749.210  Cleft lip, bilateral, with any cleft palate
749.220  Cleft lip, central, with any cleft palate
749.290  Cleft lip, NOS, with any cleft palate
750 Other Congenital Anomalies of Upper Alimentary Tract

# 750.000 Tongue tie
Ankyloglossia

750.1 Other anomalies of tongue
Excludes: protruding tongue (never a defect)

750.100 Aglossia
Absence of tongue
750.110 Hypoglossia (small tongue)
Microglossia
750.120 MacroGLOSSIA (large tongue)
750.130 Dislocation or displacement of tongue
Glossoptosis
750.140 Cleft tongue or split tongue
750.180 Other specified anomalies of tongue
750.190 Unspecified anomalies of tongue

750.2 Other specified anomalies of mouth and pharynx

750.200 Pharyngeal pouch
750.210 Other pharyngeal anomalies
750.230 Other anomalies of salivary glands or ducts
# 750.240 High arched palate
750.250 Other anomalies of palate
750.260 Lip fistulae or pits
750.270 Other lip anomalies
Includes: notched lip, prominent philtrum, long philtrum
Excludes: cleft lip (see 749)
750.280 Other specified anomalies of mouth and pharynx
Excludes: receding jaw (see 524.0)
large and small mouth (see 744.8)

750.3 Tracheoesophageal (T-E) fistula, esophageal atresia and stenosis

750.300 Esophageal atresia without mention of T-E fistula
750.310 Esophageal atresia with mention of T-E fistula
750.320 Tracheoesophageal fistula without mention of esophageal atresia
750.325 Tracheoesophageal fistula - "H" type
750.330 Bronchoesophageal fistula with or without mention of esophageal atresia
750.340 Stenosis or stricture of esophagus
750.350 Esophageal web
750.380 Other tracheoesophageal anomalies
750.4 Other specified anomalies of esophagus

750.400 Congenital dilatation of esophagus
giant esophagus
750.410 Displacement of esophagus
750.420 Diverticulum of esophagus
esophageal pouch
750.430 Duplication of esophagus
750.480 Other specified anomalies of esophagus

750.5 Congenital hypertrophic pyloric stenosis

# 750.500 Pylorospasm
750.510 Congenital hypertrophic pyloric stenosis
750.580 Other congenital pyloric obstruction

750.6 Congenital hiatus hernia

750.600 Congenital hiatus hernia
Cardia displacement through esophageal hiatus
Partial thoracic stomach
Excludes: congenital diaphragmatic hernia (756.610)

750.7 Other specified anomalies of stomach

750.700 Microgastria
750.710 Megalogastria
750.720 Cardiospasm
achalasia of cardia, congenital
750.730 Displacement or transposition of stomach
750.740 Diverticulum of stomach
750.750 Duplication of stomach
750.780 Other specified anomalies of stomach

750.8 Other specified anomalies of upper alimentary tract

750.800 Other specified anomalies of upper alimentary tract

750.9 Unspecified anomalies of upper alimentary tract

750.900 Unspecified anomalies of mouth and pharynx
750.910 Unspecified anomalies of esophagus
750.920 Unspecified anomalies of stomach
750.990 Unspecified anomalies of upper alimentary tract
751 Other Congenital Anomalies of Digestive System

751.0 Meckel's diverticulum
- 751.000 Persistent omphalomesenteric duct
- Persistent vitelline duct
- # 751.010 Meckel's diverticulum

751.1 Atresia and stenosis of small intestine
- 751.100 Stenosis, atresia or absence of duodenum
- 751.110 Stenosis, atresia or absence of jejunum
- 751.120 Stenosis, atresia or absence of ileum
- 751.190 Stenosis, atresia or absence of small intestine
- 751.195 Stenosis, atresia or absence of small intestine with fistula

751.2 Atresia and stenosis of large intestine, rectum and anal canal
- 751.200 Stenosis, atresia or absence of large intestine
- Stenosis, atresia or absence of appendix
- 751.210 Stenosis, atresia or absence of rectum with fistula
- Stenosis, atresia or absence of rectum without mention of fistula
- Includes: Imperforate anus with fistula
- 751.230 Stenosis, atresia or absence of anus with fistula
- Includes: Imperforate anus without fistula

751.3 Hirschsprung's disease and other congenital functional disorders of the colon
- 751.300 Total intestinal aganglionosis
- 751.310 Long-segment Hirschsprung's disease; aganglionosis beyond the rectum
- 751.320 Short-segment Hirschsprung's disease; aganglionosis involving no more than the anal sphincter and the rectum
- 751.330 Hirschsprung's disease, NOS
- 751.340 Congenital megacolon
- Congenital macrocolon, not aganglionic

751.4 Anomalies of intestinal fixation
- 751.400 Malrotation of cecum and/or colon
- 751.410 Anomalies of mesentery
- 751.420 Congenital adhesions or bands of omentum and peritoneum; Ladd's bands
- 751.490 Other specified and unspecified malrotation
- 751.495 Malrotation of small intestine alone

751.5 Other anomalies of intestine
- 751.500 Duplication of anus, appendix, cecum, or intestine enterogenous cyst
- 751.510 Transposition of appendix, colon, or intestine
- 751.520 Microcolon
- 751.530 Ectopic (displaced) anus
- 751.540 Congenital anal fistula
- 751.550 Persistent cloaca

T = Rev. 6/04
* = code created by CDC
# = on the MACDP Excl List
751.560 Duodenal web
751.580 Other specified anomalies of intestine
   Includes: rectal fissures
751.590 Unspecified anomalies of intestine

751.6 Anomalies of gallbladder, bile ducts, and liver

751.600 Absence or agenesis of liver, total or partial
751.610 Cystic or fibrocystic disease of liver
751.620 Other anomalies of liver
   hepatomegaly
   hepatosplenomegaly (also use code 759.020)
   Excludes: Budd-Chiari (use 453.000)
751.630 Agenesis or hypoplasia of gallbladder
751.640 Other anomalies of gallbladder
duplication of gallbladder
751.650 Agenesis or atresia of hepatic or bile ducts
   Includes: biliary atresia
   Excludes: congenital or neonatal hepatitis
   (use 774.480 or 774.490)
751.660 Choledochal cysts
751.670 Other anomalies of hepatic or bile ducts
751.680 Anomalies of biliary tract, NEC

751.7 Anomalies of pancreas
Excludes: fibrocystic disease of pancreas (277.000)
diabetes mellitus,
   congenital
   neonatal

751.700 Absence, agenesis or hypoplasia of pancreas
751.710 Accessory pancreas
751.720 Annular pancreas
751.730 Ectopic pancreas
751.740 Pancreatic cyst
751.780 Other specified anomalies of pancreas
751.790 Unspecified anomalies of pancreas

751.8 Other specified anomalies of digestive system

751.800 Absence of alimentary tract, NOS
   (complete or partial)
751.810 Duplication of alimentary tract
751.820 Ectopic digestive organs, NOS
751.880 Other specified anomalies of digestive system

751.9 Unspecified anomalies of digestive system

751.900 Unspecified anomalies of digestive system
   congenital of digestive system, NOS
   anomaly, NOS
   deformity, NOS
   obstruction, NOS
752 Congenital Anomalies of Genital Organs
Excludes: congenital hydrocele (778.600)
testicular feminization syndrome (257.800)
syndromes associated with anomalies in
number and form of chromosomes (758)

752.0 Anomalies of ovaries
752.000 Absence or agenesis of ovaries
752.010 Streak ovary
752.020 Accessory ovary
752.080 Other specified anomalies of ovaries
752.085 Multiple ovarian cysts
752.090 Unspecified anomalies of ovaries

752.1 Anomalies of fallopian tubes and broad ligaments
752.100 Absence of fallopian tube or broad ligament
752.110 Cyst of mesenteric remnant
epooophoron cyst
cyst of Gartner's duct
752.120 Fimbrial cyst
parovarian cyst
752.190 Other and unspecified anomalies of fallopian tube
and broad ligaments

752.2 Doubling of uterus
752.200 Doubling of uterus
doubling of uterus (any degree) or
associated with doubling of cervix and vagina

752.3 Other anomalies of uterus
752.300 Absence or agenesis of uterus
752.310 Displaced uterus
752.320 Fistulae involving uterus with digestive or
urinary tract
Includes: uterointestinal fistula
uterovesical fistula
752.380 Other anomalies of uterus
bicornuate uterus
unicornis uterus
752.390 Unspecified anomalies of uterus

752.4 Anomalies of cervix, vagina, and external female genitalia
752.400 Absence, atresia or agenesis of cervix
752.410 Absence or atresia of vagina, complete or partial
752.420 Congenital rectovaginal fistula
# 752.430 Imperforate hymen
# 752.440 Absence or other anomaly of vulva
fusion of vulva
hypoplastic labia majora - Always code if ≥36 weeks gestation. If
<36 weeks gestation, code only if another reportable defect is
present.
752.450 Absence or other anomaly of clitoris
Includes: clitoromegaly
    enlarged clitoris
    clitoral hypertrophy
    prominent clitoris

752.460 Embryonal cyst of vagina
752.470 Other cyst of vagina, vulva, or canal of Nuck

752.480 Other specified anomalies of cervix, vagina, or external female genitalia
Includes: vaginal tags
    hymenal tags

752.490 Unspecified anomalies of cervix, vagina, or external female genitalia

752.5 Undescended testicle

1) If < 36 weeks gestation, code only if there is a medical/surgical intervention for this problem;
2) If ≥ 36 weeks gestation and defect last noted at < 1 year of age, code only if there was a medical/surgical intervention for this problem or if another reportable defect is present;
3) Always code if ≥ 36 weeks gestation and defect first noted at ≥ 1 of age.

752.500 Undescended testicle, unilateral
    undescended, unpalpable

752.501 Left undescended testicle

752.502 Right undescended testicle

752.514 Undescended testicle, bilateral

752.520 Undescended testicle, NOS (Cryptorchidism)
752.530 Ectopic testis, unilateral and bilateral

752.6 Hypospadias and epispadias

752.600 Hypospadias (alone), NOS
752.605 1°, glandular, coronal
752.606 2°, penile
752.607 3°, perineal, scrotal
752.610 Epispadias
752.620 Congenital chordee (with hypospadias), NOS
752.621 Congenital chordee alone (chordee w/o hypospadias)
752.625 Cong. chordee with 1°, coronal hypospadias
752.626 Cong. chordee with 2°, penile hypospadias
752.627 Cong. chordee with 3°, perineal, scrotal hypospadias

752.7 Indeterminate sex and pseudohermaphroditism
Excludes: pseudohermaphroditism:
    female, with adrenocortical disorder (see 255.200)
    male, with gonadal disorder with specified chromosomal anomaly
        (see 758)

752.700 True hermaphroditism
    ovotestis

752.710 Pseudohermaphroditism, male

752.720 Pseudohermaphroditism, female
    pure gonadal dysgenesis
    Excludes: gonadal agenesis (758.690)

752.730 Pseudohermaphrodite, NOS

752.790 Indeterminate sex, NOS
    ambiguous genitalia
752.8 Other specified anomalies of male genital organs

752.800 Absence of testis
monorchidism, NOS

# 752.810 Aplasia or hypoplasia of testis and scrotum
752.820 Other anomalies of testis and scrotum
polyorchidism
bifid scrotum
Excludes: torsion of the testes or spermatic
cord (use #608.200)

752.830 Atresia of vas deferens
752.840 Other anomalies of vas deferens and prostate
752.850 Absence or aplasia of penis

# 752.860 Other anomalies of penis
absent or hooded foreskin
# redundant foreskin (never a defect)
752.865 Small penis, hypoplastic penis, or micropenis

752.870 Cysts of embryonic remnants
cyst: hydatid of Morgagni
Wolffian duct
appendix testis
752.880 Other specified anomalies of genital organs
microgenitalia
macrogenitalia

752.9 Unspecified anomalies of genital organs

752.900 Unspecified anomalies of genital organs
Congenital: of genital organ, NEC
anomaly, NOS or deformity, NOS
753  Congenital Anomalies of Urinary System

753.0 Renal agenesis and dysgenesis

753.000 Bilateral absence, agenesis, dysplasia, or hypoplasia of kidneys
Potter's syndrome
753.009 Renal agenesis, NOS
753.010 Unilateral absence, agenesis, dysplasia or hypoplasia of kidneys

753.1 Cystic kidney disease

753.100 Renal cyst (single)
753.110 Polycystic kidneys, infantile type
753.120 Polycystic kidneys, adult type
753.130 Polycystic kidneys, NOS
753.140 Medullary cystic disease, juvenile type
753.150 Medullary cystic disease, adult type
Medullary sponge kidney
753.160 Multicystic renal dysplasia
Multicystic kidney
753.180 Other specified cystic disease
Includes: cystic kidneys, NOS

753.2 Obstructive defects of renal pelvis and ureter

753.200 Congenital hydronephrosis
753.210 Atresia, stricture, or stenosis of ureter
Includes: ureteropelvic junction obstruction/stenosis
ureterovesical junction obstruction/stenosis
hypoplastic ureter
753.220 Megaloureter, NOS
Includes: hydroureter
753.290 Other and unspecified obstructive defects of renal pelvis and ureter

753.3 Other specified anomalies of kidney

753.300 Accessory kidney
753.310 Double or triple kidney and pelvis
pyelon duplex or triplex
753.320 Lobulated, fused, or horseshoe kidney
753.330 Ectopic kidney
753.340 Enlarged, hyperplastic or giant kidney
753.350 Congenital renal calculi
753.380 Other specified anomalies of kidney

753.4 Other specified anomalies of ureter

753.400 Absence of ureter
753.410 Accessory ureter
double ureter, duplex collecting system
753.420 Ectopic ureter
753.480 Other specified anomalies of ureter
Includes: ureterocele
753.485 Variations of vesicoureteral reflux
753.5 Exstrophy of urinary bladder

753.500 Exstrophy of urinary bladder
ectopia vesicae
extroversion of bladder

753.6 Atresia and stenosis of urethra and bladder neck

753.600 Congenital posterior urethral valves or posterior urethral obstruction
753.610 Other atresia, or stenosis of bladder neck
753.620 Obstruction, atresia or stenosis of anterior urethra
753.630 Obstruction, atresia or stenosis of urinary meatus
Includes: meatal stenosis
753.690 Other and unspecified atresia and stenosis of urethra and bladder neck

753.7 Anomalies of urachus

T # 753.700 Patent urachus
753.710 Cyst of urachus
753.790 Other and unspecified anomaly of urachus

753.8 Other specified anomalies of bladder and urethra

753.800 Absence of bladder or urethra
753.810 Ectopic bladder
753.820 Congenital diverticulum or hernia of bladder
753.830 Congenital prolapse of bladder (mucosa)
753.840 Double urethra or urinary meatus
753.850 Ectopic urethra or urethral orifice
753.860 Congenital digestive-urinary tract fistulae
rectovesical fistula
753.870 Urethral fistula, NOS
753.880 Other specified anomalies of bladder and urethra

753.9 Unspecified anomalies of urinary system

753.900 Unspecified anomaly of kidney
753.910 Unspecified anomaly of ureter
753.920 Unspecified anomaly of bladder
753.930 Unspecified anomaly of urethra
753.990 Unspecified anomaly of urinary system, NOS
754 Certain Congenital Musculoskeletal Anomalies

754.0 Of skull, face, and jaw
Excludes: dentofacial anomalies (524.0)
   Pierre Robin sequence (524.080)
   syphilitic saddle nose (090.000)
754.000 Asymmetry of face
754.010 Compression (Potter's) facies
   # 754.020 Congenital deviation of nasal septum
      bent nose
   T 754.030 Dolichocephaly
      Always code if ≥36 weeks gestation
      # 754.040 Depressions in skull
         Includes: large fontanelle
         small fontanelle
   754.050 Plagiocephaly
   754.055 Asymmetric head
   T 754.060 Scaphocephaly, no mention of craniosynostosis
      * 754.070 Trigonoccephaly, no mention of craniosynostosis
      Always code if ≥36 weeks gestation
      # 754.080 Other specified skull deformity, no mention of
         craniosynostosis
         Includes: brachycephaly
         acrocephaly
         turricephaly
         oxycephaly
      * 754.090 Deformity of skull, NOS

754.1 Anomalies of sternocleidomastoid muscle
   754.100 Anomalies of sternocleidomastoid muscle
      * Includes: absent or hypoplastic sternocleidomastoid
      contracture of sternocleidomastoid muscle
      sternomastoid tumor
      Excludes: congenital sternocleidomastoid torticollis
      (use 756.860)

754.2 Certain congenital musculoskeletal deformities of spine
   754.200 Congenital postural scoliosis
   754.210 Congenital postural lordosis
   754.220 Congenital postural curvature of spine, NOS

754.3 Congenital dislocation of hip
   754.300 Congenital dislocation of hip
   754.310 Unstable hip
      preluxation of hip
      subluxation of hip
      predislocation status of hip at birth
754.4 Congenital genu recurvatum and bowing of long bones of leg

- 754.400 Bowing, femur
- 754.410 Bowing, tibia and/or fibula
- 754.420 Bow legs, NOS
- 754.430 Genu recurvatum
- 754.440 Dislocation of knee, congenital
- 754.490 Deformity of leg, NOS

754.5 Varus (inward) deformities of feet

- 754.500 Talipes equinovarus
- 754.510 Talipes calcaneovalgus
# 754.520 Metatarsus varus or metatarsus adductus
- 754.530 Complex varus deformities
- 754.590 Unspecified varus deformities of feet

754.6 Valgus (outward) deformities of feet

- 754.600 Talipes calcaneovalgus
- 754.610 Congenital pes planus
- 754.615 Pes valgus
- 754.680 Other specified valgus deformities of foot
- 754.690 Unspecified valgus deformities of foot

754.7 Other deformities of feet

- 754.700 Pes cavus
  Claw foot (use 755.350 for claw foot)
- 754.720 Short Achilles tendon
- 754.730 Clubfoot, NOS
  talipes, NOS
- 754.735 Congenital deformities of foot, NOS
- 754.780 Other specified deformities of ankle and/or toes
  Includes: dorsiflexion of foot
  Excludes: widely spaced 1st and 2nd toes (use 755.600)

754.8 Other specified congenital musculoskeletal deformities

- 754.800 Pigeon chest (pectus carinatum)
- 754.810 Funnel chest (pectus excavatum)
- 754.820 Other anomalies of chest wall
  Includes: deformed chest, barrel chest
- 754.825 Shield chest
- 754.830 Dislocation of elbow
- 754.840 Club hand or fingers
- 754.850 Spade-like hand
- 754.880 Other specified deformity of hands
  (see 755.500 for specified anomalies of fingers)
755 Other Congenital Anomalies of Limbs

755.0 Polydactyly

755.005 Accessory fingers (postaxial polydactyly, Type A)
755.006 Skin tag (postaxial polydactyly, Type B)
755.007 Unspecified finger or skin tag (postaxial polydactyly, NOS)
755.010 Accessory thumbs (preaxial polydactyly)
755.020 Accessory toes (postaxial)
755.030 Accessory big toe (preaxial)
755.090 Accessory digits, NOS (hand/foot not specified)
755.095 Accessory digits hand, NOS (preaxial, postaxial not specified)
755.096 Accessory digits foot, NOS (preaxial, postaxial not specified)

755.1 Syndactyly

755.100 Fused fingers
755.110 Webbed fingers
755.120 Fused toes
755.130 Webbed toes

Code webbing of the second and third toes only if another reportable defect is present. Always code webbing of other toes regardless of whether another reportable defect is present

755.190 Unspecified syndactyly (see below for specified site)
755.191 Unspecified syndactyly thumb and/or fingers, unilateral
755.192 Unspecified syndactyly thumb and/or fingers, bilateral
755.193 Unspecified (webbed vs. fused) syndactyly thumb and/or fingers, NOS
755.194 Unspecified syndactyly toes unilateral
755.195 Unspecified syndactyly toes bilateral
755.196 Unspecified syndactyly toes, NOS
755.199 Unspecified syndactyly (i.e., webbed vs. fused) digits not known

755.2 Reduction defects of upper limb

If description of the condition includes amniotic or constricting bands use additional code, 658.800 (Only use 658.800 if another reportable defect is present)

Excludes shortening of upper limb (use 755.580) or hypoplasia of upper limb (use 755.585)

755.200 Absence of upper limb

Absent: humerus (total or partial), radius, ulna and hand
Includes: amelia of upper limb, NOS
infants with rudimentary or nubbin fingers attached to stump of humerus or shoulder girdle

755.210 Absence of upper arm and forearm

Absent: humerus (total or partial), radius and ulna (total or partial)
Present: hand (total or partial)
Includes: phocomelia of upper limb, NOS;
intercalary reduction defect of upper limb, NOS
755.220 Absence of forearm only or upper arm only
Absent: radius and ulna
Present: humerus, hand (total or partial) or
Absent: humerus
Present: radius, ulna, and hand

755.230 Absence of forearm and hand
Absent: radius and ulna (total or partial) and hand
Includes: infants with rudimentary or nubbin fingers attached to stump of forearm or elbow

755.240 Absence of hand or fingers
Absent: hand or fingers (total or partial) not in conjunction with ray or long bone reduction
Includes: rudimentary or nubbin fingers; absent individual phalanges; absent or missing fingers, NOS
Excludes: isolated absent or hypoplastic thumb (use 755.260)

755.250 Split-hand malformation
Absent: central fingers (third with or without second, fourth) and metacarpals (total or partial)
Includes: monodactyly; lobster-claw hand
Excludes: isolated absent central fingers without metacarpal defects (use 755.240)

755.260 Preaxial longitudinal reduction defect of upper limb
Absent: radius (total or partial) and/or thumb with or without second finger (total or partial)
Includes: isolated absent or hypoplastic thumb; radial ray defect, NOS

755.265 Longitudinal reduction defect of upper limb, NOS
Includes: absent forearm long bone with absent fingers, NOS

755.270 Postaxial longitudinal reduction defect of upper limb
Includes: isolated absent ulna (total or partial); absent fifth with or without fourth finger (total or partial) only if ulna or fifth ± fourth metacarpal also totally or partially absent; ulnar ray defect, NOS

755.280 Other specified reduction defect of upper limb

755.285 Transverse reduction defect of upper limb, NOS
Includes: congenital amputation of upper limb, NOS

755.290 Unspecified reduction defect of upper limb

755.3 Reduction defects of lower limb

If description of condition includes amniotic or constricting bands use additional code, 658.800 (Only use this code if another reportable defect is present)

Excludes shortening of lower limb (use 755.680) and hypoplasia of lower limb (use 755.685)

755.300 Absence of lower limb
Absent: femur (total or partial), tibia, fibula, and foot
Includes: amelia of lower limb, NOS
infants with rudimentary or nubbin toes attached to stump of femur or pelvic girdle

755.310 Absence of thigh and lower leg
<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
</table>
| 755.320 | Absence of lower leg only or femur only  
Absent: tibia and fibula  
Present: femur, foot (total or partial)  
or  
Absent: femur  
Present: tibia, fibula, and foot |
| 755.330 | Absence of lower leg and foot  
Absent: tibia and fibula (total or partial), foot  
Includes: infants with rudimentary or nubbin toes attached to stump of leg or knee |
| 755.340 | Absence of foot or toes  
Absent: foot or toes (total or partial) not in conjunction with ray or long bone reduction  
Includes: rudimentary or nubbin toes; absent individual phalanges; absent or missing toes, NOS  
Excludes: isolated absent or hypoplastic great toe (use 755.365) |
| 755.350 | Split-foot malformation  
Absent: central toes (third with or without second, fourth) and metatarsals (total or partial)  
Includes: monodactyly; lobster claw foot  
Excludes: isolated absent central toes without metatarsal defects (use 755.340)  
Note: preaxial lower limb reductions can occur with split-hand malformations of the upper limb and these lower limb defects should be coded 755.365 |
| 755.360 | Longitudinal reduction defect of lower limb, NOS  
Includes: absent long bone of leg with absent toes, NOS |
| 755.365 | Preaxial longitudinal reduction defect of lower limb  
Absent: tibia (total or partial) and/OR great toe with or without second toe (total or partial)  
Includes: isolated absent or hypoplastic great toe; tibial ray defect, NOS |
| 755.366 | Postaxial longitudinal reduction defect of lower limb  
Includes: isolated absent fibula (total or partial); absent fifth with or without fourth toe (total or partial) only if fibula or fifth ± fourth metatarsal also totally or partially absent; fibular ray defect, NOS |
| 755.380 | Other specified reduction defect of lower limb |
| 755.385 | Transverse reduction defect of lower limb, NOS  
Includes: congenital amputation of lower limb, NOS |
| 755.390 | Unspecified reduction defect of lower limb |
| 755.4 | Reduction defects of unspecified limb  
T  
If description of condition includes amniotic or constricting bands use additional code, 658.800 (note: 658.00 should only be used with another reportable defect) |
| 755.400 | Absence of limb, NOS  
Includes: amelia, NOS |
| 755.410 | Phocomelia, NOS  
Includes: intercalary reduction defect, NOS |
755.420 Transverse reduction defect, NOS
Includes: congenital amputation of unspecified limb
755.430 Longitudinal reduction defect, NOS
Includes: preaxial or postaxial reduction defect, NOS
755.440 Absent digits, not specified whether fingers or toes
755.480 Other specified reduction defect of unspecified limb
755.490 Unspecified reduction defect of unspecified limb

755.5 Other anomalies of upper limb, including shoulder girdle
Includes: complex anomalies involving all or part of upper limb

# 755.500 Anomalies of fingers
Includes: camptodactyly
clinodactyly
macrodactyly
brachydactyly
triphalangeal thumb
incurving fingers
acrocephalosyndactyly (see 756.050)
Apert's syndrome (see 756.055)

755.510 Anomalies of hand
Excludes: simian crease (use 757.200)
755.520 Anomalies of wrist
755.525 Accessory carpal bones
755.526 Madelung's deformity
755.530 Anomalies of forearm, NOS
755.535 Radioulnar dysostosis
755.536 Radioulnar synostosis
755.540 Anomalies of elbow and upper arm
755.550 Anomalies of shoulder
755.555 Cleidocranial dysostosis
755.556 Sprengel's deformity
755.560 Other anomalies of whole arm
755.580 Other specified anomalies of upper limb
Includes: hyperextensibility of upper limb
shortening of arm
755.585 Hypoplasia of upper limb
Includes: hypoplasia of fingers, hands, or arms
Excludes: aplasia or absent upper limb (see 755.2)
755.590 Unspecified anomalies of upper limb
755.6 Other anomalies of lower limb, including pelvic girdle

Includes: complex anomalies involving all or part of lower limb

# 755.600 Anomalies of toes
Includes: overlapping toes
hammer toes
widely spaced first and second toes
755.605 Hallux valgus
755.606 Hallux varus
755.610 Anomalies of foot
Includes: plantar furrow
Excludes: lobster claw foot (use 755.350)
# 755.616 Rocker-bottom foot
755.620 Anomalies of ankle
astragaloscaphoid synostosis
# 755.630 Anomalies of lower leg
angulation of tibia, tibial torsion
(exclude if clubfoot present)
755.640 Anomalies of knee
hyperextended knee
755.645 Genu valgum
755.646 Genu varum
755.647 Absent patella or rudimentary patella
755.650 Anomalies of upper leg
anteversion of femur
755.660 Anomalies of hip
Includes: coxa vara
coxa valga
other abnormalities of hips
755.665 Hip dysplasia, NOS
755.666 Unilateral hip dysplasia
755.667 Bilateral hip dysplasia
755.670 Anomalies of pelvis
fusion of sacroiliac joint
755.680 Other specified anomalies of lower limb
hyperextended legs
shortening of legs
755.685 Hypoplasia of lower limb
Includes: hypoplasia of toes, feet, legs
Excludes: aplasia or absent lower limb (see 755.3)
755.690 Unspecified anomalies of legs
755.8 Other specified anomalies of unspecified limb

755.800 Arthrogryposis multiplex congenita
   Includes: distal arthrogryposis syndrome
   Temporarily includes: flexion contractures of individual joints

755.810 Larsen’s syndrome

755.880 Other specified anomalies of unspecified limb
   Includes: overlapping digits, NOS
   hyperextended joints, NOS
   Excludes: hyperextended knees (use 755.640)

755.9 Unspecified anomalies of unspecified limb

755.900 Unspecified anomalies of unspecified limb
756  Other Congenital Musculoskeletal Anomalies

756.0  Anomalies of skull and face bones
Excludes: skull and face deformities in 754
Pierre Robin sequence (use 524.080)

756.000  Craniosynostosis, NOS
    craniostenosis, NOS
    closed-skull sutures, NOS
756.005  Sagittal craniosynostosis
756.006  Metopic craniosynostosis
756.010  Coronal craniosynostosis
756.020  Lambdoidal craniosynostosis
756.030  Other types of craniosynostosis
    Includes:  basilar craniosynostosis
756.040  Craniofacial dysostosis
    Includes:  Crouzon's disease
756.045  Mandibulofacial dysostosis
    Includes:  Franceschetti syndrome
    Treacher-Collins syndrome
756.046  Other craniofacial syndromes
    Includes:  oculomandibulofacial syndrome
    Hallermann-Streiff syndrome
756.050  Acrocephalosyndactyly, NOS
756.055  Acrocephalosyndactyly types I or II
    Apert syndrome
756.056  Acrocephalosyndactyly type III
756.057  Other specified acrocephalosyndactylies
756.060  Goldenhar syndrome
    oculoauriculovertebral dysplasia
756.065  Hemifacial microsomia
756.080  Other specified skull and face bone anomalies
    Includes:  localized skull defects
#   flat occiput
#   mid-facial hypoplasia
    prominent occiput
    prominent maxilla
    hypotelorism
Excludes:  macrocephaly (use 742.400)
    small chin (see 524.0)
Pierre Robin sequence (use 524.080)
756.085  Hypertelorism, telecanthus, wide set eyes
756.090  Unspecified skull and face bone anomalies
    Excludes:  dentofacial anomalies (524.0)
    skull defects associated with brain anomalies
    such as:
    anencephalus (740.0)
    encephalocele (742.0)
    hydrocephalus (742.3)
    microcephalus (742.100)
756.1 Anomalies of spine
- 756.100 Spina bifida occulta
- 756.110 Klippel-Feil syndrome
  Wildervanck syndrome
- 756.120 Kyphosis
  kyphoscoliosis
- 756.130 Congenital spondylolisthesis
- 756.140 Anomalies of cervical vertebrae
- 756.145 Hemivertebrae (cervical)
- 756.146 Agenesis (cervical)
- 756.150 Anomalies of thoracic vertebrae
- 756.155 Hemivertebrae of thoracic vertebrae
- 756.156 Agenesis of thoracic vertebrae
- 756.160 Anomalies of lumbar vertebrae
- 756.165 Hemivertebrae of lumbar vertebrae
- 756.166 Agenesis of lumbar vertebrae
- 756.170 Sacrococcygeal anomalies
  Includes: agenesis of sacrum
  Excludes: pilonidal sinus (see 685.100)
- 756.179 Sacral mass, NOS
- 756.180 Other specified vertebral anomalies
- 756.185 Hemivertebrae, NOS
- 756.190 Unspecified anomalies of spine

756.2 Cervical rib
- 756.200 Cervical rib
  supernumerary rib in cervical region

756.3 Other anomalies of ribs and sternum
- 756.300 Absence of ribs
- 756.310 Misshapen ribs
- 756.320 Fused ribs
- 756.330 Extra ribs
- 756.340 Other anomalies of ribs
- 756.350 Absence of sternum
- 756.360 Misshapen sternum
- 756.380 Other anomalies of sternum
  Includes: double ossification center in the manubrium,
  bifid sternum, short sternum
  Excludes: thoracic cage, unspecified
  deformed chest (use 754.820)

756.4 Chondrodystrophy
- 756.400 Asphyxiating thoracic dystrophy
  Jeune syndrome
  thoracic-pelvic-phalangeal dysplasia
  Excludes: homozygous achondroplasia
- 756.410 Chondrodysplasia
  Ollier syndrome, enchondromatosis
- 756.420 Chondrodysplasia with hemangioma
  Kast syndrome
  Maffucci syndrome
- 756.430 Achondroplastic dwarfism
- 756.440 Other specified chondrodystrophies
  Excludes: Conradi's (use 756.575)
756.445 Diastrophic dwarfism
756.446 Metatrophic dwarfism
756.447 Thanatophoric dwarfism
756.450 Metaphyseal dysostosis
756.460 Spondyloepiphyseal dysplasia
756.470 Exostosis
    Excludes: Gardner syndrome (see 759.630)
756.480 Other specified chondrodystrophy
756.490 Unspecified chondrodystrophy
    Excludes: lipochondrodystrophy (use 277.510)

756.5 Osteodystrophies

756.500 Osteogenesis imperfecta
756.505 Osteopsathyrosis
756.506 Fragilitas ossium
756.510 Polyostotic fibrous dysplasia
    Albright-McCune-Sternberg syndrome
756.520 Chondroectodermal dysplasia
756.525 Ellis-van Creveld syndrome
756.530 Infantile cortical hyperostosis
    Caffey syndrome
756.540 Osteopetrosis
    Albers-Schonberg syndrome
    marble bones
756.550 Progressive diaphyseal dysplasia
    Engelmann syndrome
    Camurati-Engelmann disease
756.560 Osteopoikilosis
756.570 Multiple epiphyseal dysplasia
756.575 Conradi syndrome
    chondrodysplasia punctata
    Excludes: warfarin embryopathy
756.580 Other specified osteodystrophies
756.590 Unspecified osteodystrophies
756.6 **Anomalies of diaphragm**

- 756.600 Absence of diaphragm
- 756.610 Congenital diaphragmatic hernia
- 756.615 Diaphragmatic hernia (Bochdalek)
- 756.616 Diaphragmatic hernia (Morgagni)
- 756.617 Hemidiaphragm
- 756.620 Eventration of diaphragm
- 756.680 Other specified anomalies of diaphragm
- 756.690 Unspecified anomalies of diaphragm

756.7 **Anomalies of abdominal wall**

- 756.700 Exomphalos, omphalocele
- 756.710 Gastrochisis
  - Excludes: umbilical hernia (553.100)
- 756.720 Prune belly syndrome
- 756.790 Other and unspecified anomalies of abdominal wall
- 756.795 Epigastric hernia

756.8 **Other specified anomalies of muscle, tendon, fascia and connective tissue**

- 756.800 Poland syndrome or anomaly
- 756.810 Other absent or hypoplastic muscle
  - Includes: absent pectoralis major
  - Excludes: prune belly syndrome (use 756.720)
- 756.820 Absent tendon
- 756.830 Nail-patella syndrome
- 756.840 Amyotrophia congenita
- 756.850 Ehlers-Danlos syndrome
- 756.860 Congenital torticollis
  - (see also 754.100, anomalies of sternocleidomastoid muscle)
- 756.880 Other specified anomalies of muscle, tendon, fascia and connective tissue
  - Includes: myopathy, congenital NOS

756.9 **Unspecified anomalies of musculoskeletal system**

- 756.900 Unspecified anomalies of muscle
- 756.910 Unspecified anomalies of tendon
- 756.920 Unspecified anomalies of bone
- 756.930 Unspecified anomalies of cartilage
- 756.940 Unspecified anomalies of connective tissue
- 756.990 Unspecified anomalies of musculoskeletal system
757 Congenital Anomalies of the Integument

757.000 Hereditary edema of legs
   Hereditary trophedema
   Milroy’s disease

757.1 Ichthyosis congenita

757.100 Harlequin fetus
757.110 Collodion baby
757.115 Bullous type
757.120 Sjogren-Larsson syndrome
757.190 Other and unspecified
757.195 Ichthyosis vulgaris
757.196 X-linked ichthyosis
757.197 Ichthyosiform erythroderma

757.2 Dermatoglyphic anomalies

# 757.200 Abnormal palmar creases
   Includes: simian creases, transverse palmar creases

757.3 Other specified anomalies of skin
Excludes: pigmented mole (216.900)
   hemangioma (see 228.0)

757.300 Specified syndromes, not elsewhere classified, involving skin
   anomalies
# 757.310 Skin tags
   Includes: anal tags
   Excludes: preauricular tag (see 744.110)
   vaginal tags (see 752.480)

757.320 Urticaria pigmentosa
757.330 Epidermolysis bullosa
757.340 Ectodermal dysplasia
   Excludes: Ellis-van Creveld syndrome (756.525)
757.345 X-linked type ectodermal dysplasia
757.346 Other specified ectodermal dysplasias
757.350 Incontinentia pigmenti
757.360 Xeroderma pigmentosum
757.370 Cutis laxa hyperelastica
# 757.380 Nevus, not elsewhere classifiable
   Includes: port wine stain or nevus flammeus

T Excludes: hairy nevus (use 216.920)
   Sturge-Weber syndrome (use 759.610)
# 757.385 Birthmark, NOS
# 757.386 Mongolian blue spot
# 757.390 Other specified anomalies of skin
   Includes: cafe au lait spots
   hyperpigmented areas
   skin cysts
   hypoplastic dermal patterns

757.395 Absence of skin

T = Rev. 6/04
* = code created by CDC
# = on the MACDP Excl List
757.4 Specified anomalies of hair
Excludes: kinky hair syndrome (use 759.870)

757.400 Congenital alopecia
Excludes: ectodermal dysplasia (use 757.340)
757.410 Beaded hair
Monilethrix
757.420 Twisted hair
Pili torti
757.430 Taenzer's hair
# 757.450 Persistent or excessive lanugo
Includes: hirsutism
757.480 Other specified anomalies of hair

757.5 Specified anomalies of nails

757.500 Congenital anonychia
Absent nails
757.510 Enlarged or hypertrophic nails
757.515 Onychauxis
757.516 Pachyonychia
757.520 Congenital koilonychia
757.530 Congenital leukonychia
757.540 Club nail
757.580 Other specified anomalies of nails
757.585 Hypoplastic (small) fingernails and/or toenails

757.6 Specified anomalies of breast

757.600 Absent breast with absent nipple
757.610 Hypoplastic breast with hypoplastic nipple
757.620 Accessory (ectopic) breast with nipple
757.630 Absent nipple
T # 757.640 Small nipple (hypoplastic)
Always code if ≥36 weeks gestation
# If <36 weeks gestation, code only if another reportable defect is present
# 757.650 Accessory (ectopic) nipple, supernumerary
# 757.680 Other specified anomalies of breast
Widely spaced nipples
Excludes: inverted nipples (never a defect)

757.8 Other specified anomalies of the integument

757.800 Includes: scalp defects
For specified anomalies of skin see 757.390
For specified anomalies of hair see 757.480
For specified anomalies of nails see 757.580

757.9 Unspecified anomalies of the integument

757.900 Unspecified anomalies of skin
757.910 Unspecified anomalies of hair, NOS
757.920 Unspecified anomalies of nail, NOS
757.990 Unspecified anomalies of the integument, NOS
758  Chromosomal Anomalies

758.0  Down syndrome
Clinical Down syndrome karyotype identified as:

T  758.000  Down syndrome, karyotype trisomy 21, cytogenetics result in record
T  758.008  Down syndrome suspected, cytogenetics pending
T  758.010  Down syndrome, karyotype trisomy G, NOS
T  758.020  Translocation trisomy - duplication of a 21
T  758.030  Translocation trisomy - duplication of a G, NOS
T  758.040  Mosaic Down syndrome
T  758.090  Down syndrome, NOS (i.e. chart states a diagnosis of Trisomy 21 or Downs syndrome, but no cytogenetics result in record)
T  758.098  Down syndrome suspected, cytogenetics never done

758.1  Patau syndrome
Clinical Patau syndrome karyotype identified as:

T  758.100  Patau syndrome, karyotype trisomy 13, cytogenetics result in record
T  758.108  Patau syndrome suspected, cytogenetics pending
T  758.110  Patau syndrome, karyotype trisomy D, NOS
T  758.120  Translocation trisomy - duplication of a 13
T  758.130  Translocation trisomy - duplication of a D, NOS
T  758.190  Patau syndrome, NOS (i.e. chart states a diagnosis of Trisomy 13 or Patau syndrome, but no cytogenetics result in record)
T  758.198  Patau syndrome suspected, cytogenetics pending

758.2  Edwards syndrome
Clinical Edwards syndrome karyotype identified as:

T  758.200  Edwards syndrome, karyotype trisomy 18, cytogenetics result in record
T  758.208  Edwards syndrome suspected, cytogenetics pending
T  758.210  Edwards syndrome, karyotype trisomy E, NOS
T  758.220  Translocation trisomy - duplication of an 18
T  758.230  Translocation trisomy - duplication of an E, NOS
T  758.290  Edwards syndrome, NOS (i.e. chart states a diagnosis of Trisomy 18 or Edwards syndrome, but no cytogenetics result in record)
T  758.295  Edwards phenotype - normal karyotype
T  758.298  Edwards syndrome suspected, cytogenetics pending
758.3 Autosomal deletion syndromes

758.300 Antimongolism syndrome
Clinical antimongolism syndrome:
karyotype - partial or total deletion of:
21
G, NOS

758.310 Cri du chat syndrome
Clinical Cri du chat syndrome:
karyotype - deletion of:
5
B, NOS

758.320 Wolff-Hirschorn syndrome
Clinical Wolff-Hirschorn syndrome:
karyotype - deletion of:
4
B, NOS

758.330 Deletion of long arm of 13
deletion of long arm of D, NOS

758.340 Deletion of long arm of E
deletion of long arm of 17 or 18

758.350 Deletion of short arm of E
deletion of short arm of 17 or 18

758.360 Monosomy G mosaicism

758.370 Deletion in band 11 of long arm of 22 (22q11 deletions)
Note: Code added for use with births on or after 4/1/2001

758.380 Other loss of autosomal material

758.390 Unspecified autosomal deletion syndromes

758.4 Balanced autosomal translocation in normal individual

758.400 Balanced autosomal translocation in normal individual

758.5 Other conditions due to autosomal anomalies

758.500 Trisomy 8

758.510 Other trisomy C syndromes
Trisomy: 6, 7, 9, 10, 11, 12, or C, NOS

758.520 Other total trisomy syndromes
Trisomy 22
Trisomy, NOS

758.530 Partial trisomy syndromes

758.540 Other translocations
Excludes: balanced translocation in normal individual (use 758.400)

758.580 Other specified anomalies of autosomes, NOS
Includes: marker autosome

758.585 Polyploidy

758.586 Triploidy

758.590 Unspecified anomalies of autosomes
758.6 Gonadal Dysgenesis
Excludes: pure gonadal dysgenesis (752.720)
Noonan syndrome (759.800)

758.600 Turner's phenotype, karyotype 45, X [XO]
758.610 Turner's phenotype, variant karyotypes
karyotype characterized by:
mosaic, including XO
partial X deletion
ring chromosome
Excludes: Turner's phenotype, karyotype normal XX
(use 759.800, Noonan syndrome)

758.690 Turner syndrome, karyotype unspecified, NOS
Bonneville-Ullrich syndrome, NOS

758.7 Klinefelter syndrome

758.700 Klinefelter's phenotype, karyotype 47, XXY
758.710 Klinefelter's phenotype, other karyotype with additional
X chromosomes
XX
XXY
XXXX

758.790 Klinefelter syndrome, NOS

758.8 Other conditions due to sex chromosome anomalies

758.800 Mosaic XO/XY, 45X/46XY
Excludes: with Turner's phenotype (758.610)
758.810 Mosaic XO/XX
Excludes: with Turner's phenotype (758.610)
758.820 Mosaic XY/XXY,46XY/47XXY
Excludes: Klinefelter's phenotype (758.710)
758.830 Mosaic including XXXXY,49XXXXY
Excludes: with Klinefelter's phenotype (use 758.710)
758.840 XYY, male, 47XYY
mosaic XYY male
758.850 XXX female,47XXX
758.860 Additional sex chromosomes, NOS
758.880 Other specified sex chromosome anomaly
Includes: fragile X

758.890 Unspecified sex chromosome anomaly

758.9 Conditions due to anomaly of unspecified chromosomes

758.900 Mosaicism, NOS
758.910 Additional chromosome(s), NOS
758.920 Deletion of chromosome(s), NOS
758.930 Duplication of chromosome(s), NOS
758.990 Unspecified anomaly of chromosome(s)
759 Other and Unspecified Congenital Anomalies

759.0 Anomalies of spleen

759.000 Absence of spleen
    asplenia
759.005 Ivemark syndrome
759.010 Hypoplasia of spleen
# 759.020 Hyperplasia of spleen
    splenomegaly
    hepatosplenoemegaly (also use code 751.620)
759.030 Misshapen spleen
759.040 Accessory spleen
759.050 Ectopic spleen
759.080 Other specified anomalies of spleen
759.090 Unspecified anomalies of spleen

759.1 Anomalies of adrenal gland

759.100 Absence of adrenal gland
759.110 Hypoplasia of adrenal gland
759.120 Accessory adrenal gland
759.130 Ectopic adrenal gland
759.180 Other specified anomaly of adrenal gland
    Excludes: congenital adrenal hyperplasia
    (use 255.200)
759.190 Unspecified anomalies of adrenal gland

759.2 Anomalies of other endocrine glands

759.200 Anomalies of pituitary gland
759.210 Anomalies of thyroid gland
    thyroglossal duct anomalies
    thyroglossal cyst
759.230 Anomalies of parathyroid gland
# 759.240 Anomalies of thymus
    thymic hypertrophy
    absent thymus
759.280 Other specified anomalies of endocrine gland
759.290 Unspecified anomaly of endocrine gland
759.3 Situs inversus

759.300 Dextrocardia with complete situs inversus
759.310 Situs inversus with levocardia
759.320 Situs inversus thoracis
759.330 Situs inversus abdominis
759.340 Kartagener syndrome (triad)
759.390 Unspecified situs inversus
   Excludes: dextrocardia (746.800) not associated with complete situs inversus

759.4 Conjoined twins

759.400 Dicephalus
   two heads
759.410 Craniopagus
   head-joined twins
759.420 Thoracopagus
   thorax-joined twins
759.430 Xiphopagus
   xiphoid- and pelvis-joined twins
759.440 Pygopagus
   buttock-joined twins
759.480 Other specified conjoined twins
759.490 Unspecified conjoined twins

759.5 Tuberous sclerosis

759.500 Tuberous sclerosis
   Bourneville's disease
   epiloia

759.6 Other hamartoses, not elsewhere classified

759.600 Peutz-Jeghers syndrome
759.610 Encephalocutaneous angiomatosis
   Kalischer's disease
   Sturge-Weber syndrome
759.620 Von Hippel-Lindau syndrome
759.630 Gardner syndrome
759.680 Other specified hamartomas
759.690 Unspecified hamartomas

759.7 Multiple congenital anomalies,

759.700 Multiple congenital anomalies,
   anomaly, multiple, NOS
   deformity, multiple, NOS
759.8 Other specified anomalies and syndromes

759.800 Cong malformation syndromes affecting facial appearance
cyclops
    Noonan syndrome
    oral-facial-digital (OFD) syndrome, type I
    Orofaciodigital syndrome, type II (Mohr syndrome)
    Waardenburg syndrome
    whistling face syndrome

759.820 Cong malformation syndromes associated with short stature
    Amsterdam dwarf (Cornelia de Lange syndrome)
    Cockayne syndrome
    Laurence-Moon-Biedl syndrome
    Russell-Silver syndrome
    Seckel syndrome
    Smith-Lemli-Opitz syndrome

759.840 Cong malformation syndromes involving limbs
    Carpenter syndrome
    Holt-Oram syndrome
    Klippel-Trenaunay-Weber syndrome
    Rubinstein-Taybi syndrome
    sirenomelia
    thrombocytopenia-absent radius (TAR) syndrome

759.860 Cong malformation syndromes with other skeletal changes
    Marfan syndrome
    Stickler syndrome

759.870 Cong malformation syndromes with metabolic disturbances
    Alport syndrome
    Beckwith (Wiedemann-Beckwith) syndrome
    leprechaunism
    Menkes syndrome (kinky hair syndrome)
    Prader-Willi syndrome
    Zellweger syndrome

759.890 Other specified anomalies
    Includes: hemihypertrophy
               Meckel-Gruber syndrome

759.9 Congenital anomaly, unspecified

# 759.900 Anomalies of umbilicus
    low-lying umbilicus
    umbilical cord atrophy

759.910 Embryopathy, NEC

759.990 Congenital anomaly, NOS
Other Specified Codes Used in Metro Atlanta Congenital Defects Program

List ordered alphabetically

524.000 Abnormalities of jaw size
micrognathia T
macrognathia T
255.200 Adrenogenital syndrome
# 270.200 Albinism
# 277.620 Alpha-1 antitrypsin deficiency
T # 658.800 Amniotic bands (constricting bands, amniotic cyst)
# 270.600 Arginosuccinic aciduria
# 778.000 Ascites, congenital

216 Benign neoplasm of skin
T (NOTE: All neoplasms should be coded ONLY if another reportable code is present)
Includes: blue nevus pigmented nevus
papilloma dermatofibroma
* syringoadenoma hydrocystoma
* dermoid cyst syringoma
Excludes: skin of female genital organs (use 221.000), skin of male genital organs (use 222.000)
T # 216.200 Benign neoplasm of skin, ear and external auditory canal
Includes: auricle ear
external meatus
auricular canal
external canal
pinna
Excludes: cartilage of ear
T # 216.100 Benign neoplasm of skin, eyelid, including canthus
Excludes: cartilage of eyelid
T # 216.000 Benign neoplasm of skin, lip
Excludes: vermilion border of lip
T # 216.700 Benign neoplasm of skin, lower limb, hip
T # 216.300 Benign neoplasm of skin, other and unspecified parts of face
Includes: cheek, external nose, external eyebrow
 temple
T # 216.800 Benign neoplasm of skin, other specified sites of skin
Excludes: epibulbar dermoid cyst (use 743.810)
T # 216.400 Benign neoplasm of skin, scalp and skin of neck
T # 216.900 Benign neoplasm of skin, site unspecified
# 216.500 Benign neoplasm of skin, trunk, except scrotum
Includes: axillary fold
perianal skin
skin of: chest wall, abdominal wall, groin,
buttock, anus, perineum, back, umbilicus, breast
Excludes: anal canal
anus, NOS
skin of scrotum

T = Rev. 6/04
* = code created by CDC
# = on the MACDP Excl List
Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

T  # 216.600 Benign neoplasm of skin, upper limb, shoulder  
# 221.000 Benign skin neoplasm of female genital organs  
# 222.000 Benign skin neoplasm of male genital organs  
453.000 Budd-Chiari, occlusion of hepatic vein  
427.900 Cardiac arrhythmias, NEC  
# 330.100 Cerebral lipidoses  
  Includes: Tay-Sachs disease, gangliosidosis  
363.200 Chorioretinitis  
279.200 Combined immunodeficiency syndrome  
771.280 Congenital infection, other specified  
  Excludes: human immunodeficiency virus (HIV) infection and acquired immunodeficiency syndrome (AIDS)  
# 277.000 Cystic fibrosis  
  No mention of meconium ileus  
# 277.010 Cystic fibrosis  
  With mention of meconium ileus  
228.100 Cystic hygroma  
  Lymphangioma, any site  
771.100 Cytomegalovirus (CMV) (in utero infections only)  
253.820 Diencephalic syndrome  
279.110 DiGeorge syndrome  
277.400 Disorders of bilirubin excretion  
425.300 Endocardial fibroelastosis  
553.200 Epigastric hernia  
# 767.600 Erb's palsy  
# 368.000 Esotropia  
# 378.000 Exotropia  
# 351.000 Facial palsy  
331.890 Familial degenerative CNS disease  
760.710 Fetal alcohol syndrome  
760.718 Fetal alcohol syndrome, probable  
  Includes: "facies"  
760.750 Fetal hydantoin (Dilantin) syndrome  
# 282.200 Glucose-6-phosphate dehydrogenase (G-6-PD) deficiency  
# 271.000 Glycogen storage diseases  
T  216.920 Hairy nevus  
T  228.0 Hemangioma  
  Include if greater than 4-inches diameter, if multiple hemangiomas, or if cavernous hemangioma  
228.040 Hemangioma, intra-abdominal (Always code regardless of size, type or number)  
228.020 Hemangioma, intracranial (Always code regardless of size, type or number)  
228.090 Hemangioma, of other sites (Always code regardless of size, type or number)  
# 228.000 Hemangioma, of unspecified site. Always code if multiple hemangiomas of any size are present, if one or more cavernous hemangiomas of any size are present, or if a single hemangioma measuring ≥ 4cm in diameter or described as large, huge, or of medical significance is present.  
228.030 Hemangioma, retinal (Always code regardless of size, type or number)
Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

# 228.010 Hemangioma, skin & subcutaneous, NOS Always code if multiple hemangiomas of any size are present, if one or more cavernous hemangiomas of any size are present, or if a single hemangioma measuring $\geq 4$cm in diameter or described as large, huge, or of medical significance is present.

# 286.000 Hemophilia (all types)
774.490 Hepatitis, neonatal, NOS
774.480 Hepatitis, neonatal, other specified
# 282.100 Hereditary elliptocytosis
# 282.000 Hereditary spherocytosis

771.220 Herpes simplex (in utero infections only)
   Includes: encephalitis
   meningoencephalitis
202.300 Histiocytosis, malignant
277.510 Hurler syndrome
   Includes: lipochondrodystrophy
# 778.600 Hydrocele, congenital
# 270.700 Hyperglycinemia
# 251.200 Hypoglycemia, idiopathic
# 252.100 Hypoparathyroidism, congenital
# 275.330 Hypophosphatemic rickets
253.280 Hypopituitarism, congenital
# 243.990 Hypothyroidism, congenital (Exclude even if other defects are present only if the record specifies hypothyroidism of prematurity. Other types of hypothyroidism or hypothyroidism NOS should continue to be on the routine exclusion list.)
345.600 Infantile spasms, congenital
# 550.000 Inguinal hernia or patent processus vaginalis never
   -550.900 code in infants if <36 weeks gestation regardless of the presence of a reportable defect.
   NOTE: for those $\geq 36$ weeks:
   Code in males only if another reportable defect is present;
   Code in females, always code even if found in isolation
208.000 Leukemia, congenital, NOS
214 Lipoma

214.300 Lipoma, intra-abdominal organs
214.200 Lipoma, intrathoracic organs
214.810 Lipoma, lumbar or sacral lipoma
   paraspinous lipoma
214.100 Lipoma, other skin and subcutaneous tissue
214.800 Lipoma, other specified sites
214.000 Lipoma, skin and subcutaneous tissue of face
214.400 Lipoma, spermatic cord
214.900 Lipoma, unspecified site
# 457.800 Lymphatics - other specified disorders of (including chylothorax)
524.000 Macrognathia
# 270.300 Maple syrup urine disease
# 777.600 Meconium peritonitis
# 777.100 Meconium plug syndrome
524.000 Micrognathia
352.600 Moebius syndrome
# 520.600 Natal teeth
239.200 Neck cyst
774.490 Neonatal hepatitis, NOS

T = Rev. 6/04
* = code created by CDC
# = on the MACDP Excl List
Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

774.480 Neonatal hepatitis, other specified
159.800 Neoplasms of the abdomen, other specified
191.000 Neoplasms of the CNS
Includes: medulloblastoma, gliomas
171.800 Neoplasms of the connective tissue
Includes: Ewing's sarcoma
fibrosarcoma
155.000 Neoplasms of the liver
Includes: hepatoblastoma
hemangio-epithelioma
162.800 Neoplasms of the lung
186.000 Neoplasms of the testes
194.000 Neuroblastoma
237.700 Neurofibromatosis
# 379.500 Nystagmus
# 270.100 Phenylketonuria (PKU)
* 524.080 Pierre Robin sequence
# 685.100 Pilonidal sinus (sacrodermal), sacral sinus, sacral dimple
# 277.630 Pseudocholinesterase enzyme deficiency
# 284.000 Red cell aplasia
362.600 Retinal degeneration, peripheral
362.700 Retinitis pigmentosa
190.500 Retinoblastoma
771.000 Rubella, congenital (in utero infections only)
# 685.100 Sacral dimple
T # 216.910 Sebaceous cyst
# 282.600 Sickle cell anemia
# 090.000 Syphilis, congenital (in utero infections only)
238.030 Teratoma, abdomen
238.010 Teratoma, head and face
238.020 Teratoma, neck
238.000 Teratoma, NOS
238.080 Teratoma, other specified
238.040 Teratoma, sacral, coccygeal
257.800 Testicular feminization syndrome
771.090 TORCH infection, unspecified (in utero infections only)
# 608.200 Torsion of the testes or spermatic cord
771.210 Toxoplasmosis (in utero infections only)
# 553.100 Umbilical hernia
# 286.400 von Willebrand disease
335.000 Werdnig-Hoffman disease
189.000 Wilms tumor (nephroblastoma)
426.705 Wolfe-Parkinson-White syndrome, congenital
**Other Specified Codes Used in Metro Atlanta Congenital Defects Program**

List ordered by 6-digit code number

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td># 090.000</td>
<td>Syphilis, congenital (in utero infections only)</td>
</tr>
<tr>
<td>155.000</td>
<td>Neoplasms of the liver</td>
</tr>
<tr>
<td></td>
<td>Includes: hepatoblastoma</td>
</tr>
<tr>
<td></td>
<td>hemangio-epithelioma</td>
</tr>
<tr>
<td>159.800</td>
<td>Neoplasms of the abdomen</td>
</tr>
<tr>
<td>162.800</td>
<td>Neoplasms of the lung</td>
</tr>
<tr>
<td>171.800</td>
<td>Neoplasms of connective tissue</td>
</tr>
<tr>
<td></td>
<td>Includes: Ewing's sarcoma</td>
</tr>
<tr>
<td></td>
<td>fibrosarcoma</td>
</tr>
<tr>
<td>186.000</td>
<td>Neoplasms of the testes</td>
</tr>
<tr>
<td>189.000</td>
<td>Wilms tumor (nephroblastoma)</td>
</tr>
<tr>
<td>190.500</td>
<td>Retinoblastoma</td>
</tr>
<tr>
<td>191.000</td>
<td>Neoplasms of the CNS</td>
</tr>
<tr>
<td></td>
<td>Includes: gliomas</td>
</tr>
<tr>
<td></td>
<td>medulloblastoma</td>
</tr>
<tr>
<td>194.000</td>
<td>Neuroblastoma</td>
</tr>
<tr>
<td>202.300</td>
<td>Histiocytosis, malignant</td>
</tr>
<tr>
<td>208.000</td>
<td>Leukemia, congenital, NOS</td>
</tr>
<tr>
<td>214</td>
<td>Lipoma</td>
</tr>
<tr>
<td>214.000</td>
<td>Lipoma, skin and subcutaneous tissue of face</td>
</tr>
<tr>
<td>214.100</td>
<td>Lipoma, other skin and subcutaneous tissue</td>
</tr>
<tr>
<td>214.200</td>
<td>Lipoma, intrathoracic organs</td>
</tr>
<tr>
<td>214.300</td>
<td>Lipoma, intra-abdominal organs</td>
</tr>
<tr>
<td>214.400</td>
<td>Lipoma, spermatic cord</td>
</tr>
<tr>
<td>214.800</td>
<td>Lipoma, other specified sites</td>
</tr>
<tr>
<td>214.810</td>
<td>Lipoma, lumbar or sacral lipoma</td>
</tr>
<tr>
<td></td>
<td>paraspinal lipoma</td>
</tr>
<tr>
<td>214.900</td>
<td>Lipoma, unspecified site</td>
</tr>
<tr>
<td>T 216</td>
<td>Benign neoplasm of skin</td>
</tr>
<tr>
<td></td>
<td>(NOTE: All benign neoplasms should be coded ONLY if another reportable code is present)</td>
</tr>
<tr>
<td></td>
<td>Includes: blue nevus pigmented nevus</td>
</tr>
<tr>
<td></td>
<td>papilloma dermatofibroma</td>
</tr>
<tr>
<td></td>
<td>syringoadenoma</td>
</tr>
<tr>
<td></td>
<td>*dermoid cyst</td>
</tr>
<tr>
<td></td>
<td>hydrocystoma</td>
</tr>
<tr>
<td></td>
<td>syringoma</td>
</tr>
<tr>
<td></td>
<td>Excludes: skin of female genital organs (use 221.000), skin of male genital organs (use 222.000)</td>
</tr>
<tr>
<td># 216.000</td>
<td>Skin of lip</td>
</tr>
<tr>
<td></td>
<td>Excludes: vermilion border of lip</td>
</tr>
<tr>
<td># 216.100</td>
<td>Eyelid, including canthus</td>
</tr>
<tr>
<td></td>
<td>Excludes: cartilage of eyelid</td>
</tr>
<tr>
<td># 216.200</td>
<td>Ear and external auditory canal</td>
</tr>
<tr>
<td></td>
<td>Includes: auricle ear, external meatus</td>
</tr>
<tr>
<td></td>
<td>auricular canal, external canal</td>
</tr>
<tr>
<td></td>
<td>pinna</td>
</tr>
<tr>
<td></td>
<td>Excludes: cartilage of ear</td>
</tr>
<tr>
<td># 216.300</td>
<td>Skin of other and unspecified parts of face</td>
</tr>
<tr>
<td></td>
<td>Includes: cheek, external nose, external eyebrow, temple</td>
</tr>
</tbody>
</table>
Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

# 216.400 Scalp and skin of neck
# 216.500 Skin of trunk, except scrotum
   Includes: axillary fold
   perianal skin
   skin of: chest wall
   abdominal wall
groin
buttock
anus
perineum
back
umbilicus
breast
Excludes: anal canal
         anus, NOS
         skin of scrotum
# 216.600 Skin of upper limb, shoulder
# 216.700 Skin of lower limb, hip
# 216.800 Other specified sites of skin
   Excludes: epibulbar dermoid cyst (use 743.810)
# 216.900 Site unspecified
# 216.910 Sebaceous cyst
# 216.920 Hairy nevus
# 221.000 Benign skin neoplasm of female genital organs
# 222.000 Benign skin neoplasm of male genital organs

T  228.0  Hemangioma
   Include if greater than 4-inches diameter, if multiple
   hemangiomas, or if cavernous hemangioma
# 228.000 Hemangioma, of unspecified site
   Always code if multiple hemangiomas of any size are present, if one
   or more cavernous hemangiomas of any size are present, or if a single
   hemangioma measuring ≥ 4cm in diameter or described as large, huge, or
   of medical significance is present.
# 228.010 Hemangioma, skin & subcutaneous, NOS
   Always code if multiple hemangiomas of any size are present, if one
   or more cavernous hemangiomas of any size are present, or if a single
   hemangioma measuring ≥ 4cm in diameter or described as large, huge, or
   of medical significance is present.
   228.020 Hemangioma, intracranial (Always code regardless of size, type or
   number)
   228.030 Hemangioma, retinal (Always code regardless of size, type or number)
   228.040 Hemangioma, intra-abdominal (Always code regardless of size, type or
   number)
   228.090 Hemangioma, of other sites (Always code regardless of size, type or
   number)
   228.100 Cystic hygroma
   Lymphangioma, any site
   237.700 Neurofibromatosis
   238.000 Teratoma, NOS
   238.010 Teratoma, head and face
   238.020 Teratoma, neck
   238.030 Teratoma, abdomen
   238.040 Teratoma, sacral, coccygeal
   238.080 Teratoma, other specified
   239.200 Neck cyst

T = Rev. 6/04
* = code created by CDC
# = on the MACDP Excl List
Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

# 243.990 Hypothyroidism, congenital
   (Exclude even if other defects are present only if the record specifies hypothyroidism of prematurity <36 weeks. Include other types of hypothyroidism and hypothyroidism NOS only when another reportable defect is present regardless of gestational age)
# 251.200 Hypoglycemia, idiopathic
# 252.100 Hypoparathyroidism, congenital
# 253.280 Hypopituitarism, congenital
# 253.820 Diencephalic syndrome
# 255.200 Adrenogenital syndrome (adrenal hyperplasia)
# 257.800 Testicular feminization syndrome
# 270.100 Phenylketonuria (PKU)
# 270.200 Albinism
# 270.300 Maple syrup urine disease
# 270.600 Argininosuccinic aciduria
# 270.700 Hyperglycinemia
# 271.000 Glycogen storage diseases
# 275.330 Hypophosphatemic rickets
# 277.000 Cystic fibrosis with no mention of meconium ileus
# 277.010 Cystic fibrosis with mention of meconium ileus
# 277.400 Disorders of bilirubin excretion
# 277.510 Hurler syndrome
   Includes: lipochondrodystrophy
# 277.620 Alpha-1 antitrypsin deficiency
# 277.630 Pseudocholinesterase enzyme deficiency
# 279.110 DiGeorge syndrome
# 279.200 Combined immunodeficiency syndrome
# 282.000 Hereditary spherocytosis
# 282.100 Hereditary elliptocytosis
# 282.200 Glucose-6-phosphate dehydrogenase (G-6-PD) deficiency
# 282.600 Sickle cell anemia
# 284.000 Red cell aplasia
# 286.000 Hemophilia (all types)
# 286.400 von Willebrand disease
# 330.100 Cerebral lipidoses
   Includes: Tay-Sachs disease
   gangliosidosis
# 331.890 Familial degenerative CNS disease
# 335.000 Werdnig-Hoffman disease
# 345.600 Infantile spasms, congenital
# 351.000 Facial palsy
# 352.600 Moebius syndrome
# 362.600 Retinal degeneration, peripheral
# 362.700 Retinitis pigmentosa
# 363.200 Chorioretinitis
# 368.000 Esotropia
# 378.000 Exotropia
# 379.500 Nystagmus
# 425.300 Endocardial fibroelastosis
# 426.705 Congenital Wolfe-Parkinson-White syndrome
# 427.900 Cardiac arrhythmias, NEC
# 453.000 Budd-Chiari, occlusion of hepatic vein
# 457.800 Other specified disorders of lymphatics (including chylothorax)
### Continued: Other Specified Codes Used in Metro Atlanta Congenital Defects Program

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td># 520.600</td>
<td>Natal teeth</td>
</tr>
<tr>
<td>524.000</td>
<td>Abnormalities of jaw size</td>
</tr>
<tr>
<td></td>
<td>micrognathia</td>
</tr>
<tr>
<td></td>
<td>macrognathia</td>
</tr>
<tr>
<td>* 524.080</td>
<td>Pierre Robin sequence</td>
</tr>
<tr>
<td># 550.000-</td>
<td>Ingual hernia or patent processus vaginalis never</td>
</tr>
<tr>
<td>550.900</td>
<td>code in infants if &lt;36 weeks gestation regardless of the presence of a reportable defect.</td>
</tr>
<tr>
<td>NOTE: for those ≥36 weeks: Code in <strong>males</strong> only if another reportable defect is present; in <strong>females</strong>, always code even if found in isolation</td>
<td></td>
</tr>
<tr>
<td># 553.100</td>
<td>Umbilical hernia</td>
</tr>
<tr>
<td>553.200</td>
<td>Epigastric hernia</td>
</tr>
<tr>
<td># 608.200</td>
<td>Torsion of testes or spermatic cord</td>
</tr>
<tr>
<td>T # 658.800</td>
<td>Amniotic bands (constricting bands, amniotic cyst)</td>
</tr>
<tr>
<td># 685.100</td>
<td>Pilonidal sinus (sacrodermal), sacral sinus, sacral dimple</td>
</tr>
<tr>
<td>760.710</td>
<td>Fetal alcohol syndrome</td>
</tr>
<tr>
<td>760.718</td>
<td>Probable fetal alcohol syndrome</td>
</tr>
<tr>
<td></td>
<td>Includes: &quot;facies&quot;</td>
</tr>
<tr>
<td>760.750</td>
<td>Fetal hydantoin (Dilantin) syndrome</td>
</tr>
<tr>
<td># 767.600</td>
<td>Erb's palsy</td>
</tr>
<tr>
<td>771.000</td>
<td>Congenital infections (in utero infections only)</td>
</tr>
<tr>
<td></td>
<td>Excludes: congenital syphilis (use 090.000)</td>
</tr>
<tr>
<td>771.090</td>
<td>TORCH infection, unspecified</td>
</tr>
<tr>
<td>771.100</td>
<td>Cytomegalovirus (CMV)</td>
</tr>
<tr>
<td>771.210</td>
<td>Toxoplasmosis</td>
</tr>
<tr>
<td>771.220</td>
<td>Herpes simplex</td>
</tr>
<tr>
<td></td>
<td>Includes: encephalitis</td>
</tr>
<tr>
<td></td>
<td>meningoencephalitis</td>
</tr>
<tr>
<td>771.280</td>
<td>Congenital infection, other specified</td>
</tr>
<tr>
<td></td>
<td>Excludes: human immunodeficiency virus (HIV) infection and acquired immunodeficiency syndrome (AIDS)</td>
</tr>
<tr>
<td>774.480</td>
<td>Hepatitis, neonatal, other specified</td>
</tr>
<tr>
<td>774.490</td>
<td>Hepatitis, neonatal, NOS</td>
</tr>
<tr>
<td># 777.100</td>
<td>Meconium plug syndrome</td>
</tr>
<tr>
<td># 777.600</td>
<td>Meconium peritonitis</td>
</tr>
<tr>
<td># 778.000</td>
<td>Ascites, congenital</td>
</tr>
<tr>
<td># 778.600</td>
<td>Hydrocele, congenital</td>
</tr>
</tbody>
</table>

HHS:PHS:CDC:NCBDDD:DBDDD:06/16/04  
Doc. 6digit88, Version 06/04
EXCLUSION LIST for the MACDP
Nonreportable birth defects

Conditions Never to be Reported

The following newborn and infant conditions include those descriptions considered to be excludable or nonreportable conditions in the MACDP. This includes certain biochemical disorders not considered part of the present MACDP case definition.

Alphabetical list of conditions that are never considered to be defects.

Description

Anal fissure
Breast hypertrophy
Bronchopulmonary dysplasia (Wilson-Mikity syndrome)
Cephalohematoma
Chalasia (gastroesophageal reflux)
CNS hemorrhage
 Conjunctivitis
Diastasis recti
Epulis
Gastroesophageal reflux
Gum cysts - Includes epulis, ranula, mucocele
Hydrocephalus secondary to intraventricular hemorrhage (IVH) or CNS bleed
Hip click, with no follow-up or therapy
T
Heart murmur
T
Hyaline membrane disease
T
Intestinal obstruction - requires chart review to determine if cause of obstruction is a reportable defect. If so, code only the cause.
T
Intussusception - requires chart review to determine if cause of intussusception is a reportable defect. If so, code only the cause.
Inverted nipples
Laryngotracheomalacia or tracheomalacia
Meconium stained skin or nails
Mucocele
Neonatal acne
Overriding (overlapping) sutures
Petechiae
Phimosis
Pneumothorax
Protruding tongue
Ranula
Redundant foreskin
Retractile testes
Tracheomalacia
T
Volvulus - requires chart review to determine if cause of volvulus is a reportable defect. If so, code only the cause.
Wilson-Mikity syndrome
EXCLUSION LIST for the MACDP
Nonreportable birth defects

Conditions Which may be Included Under Certain Conditions

The following newborn and infant conditions include those descriptions considered to be excludable or nonreportable conditions in the MACDP, but which may be included under certain circumstances.

The following rules apply to coding these conditions:

A. If a condition or defect listed appears in a chart, singly or in any combination with other defects listed only on the Exclusion List, do not fill out the case record form.

B. If one of these conditions listed accompanies a reportable birth defect (from the 6-digit code manual and not on the exclusion list), then use the listed 6-digit code and record all defects (including those from this list) from the hospital record onto the case abstraction form.

Alphabetical list of conditions requiring no record abstraction to be performed unless associated with a reportable defect. The addition or revision dates of the changes in the list of conditions requiring no record abstraction are shown.

<table>
<thead>
<tr>
<th>Revised/ Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>744.100</td>
<td></td>
<td>Accessory auricle</td>
</tr>
<tr>
<td>757.650</td>
<td></td>
<td>Accessory nipple (supernumerary nipple, or skin tag)</td>
</tr>
<tr>
<td>270.200</td>
<td></td>
<td>Albinism</td>
</tr>
<tr>
<td>277.620</td>
<td></td>
<td>Alpha 1-antitrypsin deficiency</td>
</tr>
<tr>
<td>T 658.800</td>
<td></td>
<td>Amniotic bands (constricting bands, amniotic cyst)</td>
</tr>
<tr>
<td>757.310</td>
<td></td>
<td>Anal tags</td>
</tr>
<tr>
<td>10/1/92 T 746.400</td>
<td></td>
<td>Aortic valve insufficiency or regurgitation, congenital - Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
</tr>
<tr>
<td>270.600</td>
<td></td>
<td>Argininosuccinic aciduria</td>
</tr>
<tr>
<td>T 778.000</td>
<td></td>
<td>Ascites or anasarca, congenital. Includes: hydrops fetalis</td>
</tr>
<tr>
<td>744.220</td>
<td></td>
<td>Bat ear</td>
</tr>
<tr>
<td>T # 216.200</td>
<td></td>
<td>Benign neoplasm of skin, ear and external auditory canal Includes: auricle ear external meatus auricular canal external canal pinna</td>
</tr>
<tr>
<td>Excludes: cartilage of ear</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T # 216.100</td>
<td></td>
<td>Benign neoplasm of skin, eyelid, including canthus</td>
</tr>
<tr>
<td>Excludes: cartilage of eyelid</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T # 216.000</td>
<td></td>
<td>Benign neoplasm of skin, lip</td>
</tr>
<tr>
<td>Excludes: vermillion border of lip</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T # 216.700</td>
<td></td>
<td>Benign neoplasm of skin, lower limb, hip</td>
</tr>
<tr>
<td>T # 216.300</td>
<td></td>
<td>Benign neoplasm of skin, other and unspecified parts of face</td>
</tr>
</tbody>
</table>

= Rev. 6/94
* = code created by CDC
# = on the MACDP Excl List
Includes: cheek, external nose, external eyebrow, temple

**EXCLUSION LIST** for the MACDP
Nonreportable birth defects

## Alphabetical - Conditions Which may be Included Under Certain Conditions

<table>
<thead>
<tr>
<th>Revised/ Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>216.800</td>
<td>Benign neoplasm of skin, other specified sites of skin. Excludes: epibulbar dermoid cyst (use 743.810)</td>
</tr>
<tr>
<td>T</td>
<td>216.400</td>
<td>Benign neoplasm of skin, scalp and skin of neck</td>
</tr>
<tr>
<td>T</td>
<td>216.900</td>
<td>Benign neoplasm of skin, site unspecified</td>
</tr>
<tr>
<td></td>
<td>216.500</td>
<td>Benign neoplasm of skin, trunk, except scrotum</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Includes: axillary fold, perianal skin, skin of: chest wall, abdominal wall, groin, buttock, anus, perineum, back, umbilicus, breast</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Excludes: anal canal, anus, NOS skin of scrotum</td>
</tr>
<tr>
<td></td>
<td>216.600</td>
<td>Benign neoplasm of skin, upper limb, shoulder</td>
</tr>
<tr>
<td></td>
<td>221.000</td>
<td>Benign skin neoplasm of female genital organs</td>
</tr>
<tr>
<td></td>
<td>222.000</td>
<td>Benign skin neoplasm of male genital organs</td>
</tr>
<tr>
<td></td>
<td>754.020</td>
<td>Bent nose, deviation of nasal septum</td>
</tr>
<tr>
<td></td>
<td>744.820</td>
<td>Big lips</td>
</tr>
<tr>
<td></td>
<td>757.385</td>
<td>Birth mark, NOS</td>
</tr>
<tr>
<td></td>
<td>743.450</td>
<td>Blue sclera - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Brusfield spots</td>
</tr>
<tr>
<td></td>
<td>757.390</td>
<td>Cafe au lait spots</td>
</tr>
<tr>
<td></td>
<td>746.860</td>
<td>Cardiomegaly, congenital NOS</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Cauliflower ear</td>
</tr>
<tr>
<td></td>
<td>330.100</td>
<td>Cerebral lipidoses (e.g., Tay-Sachs, gangliosidoses, etc.)</td>
</tr>
<tr>
<td></td>
<td>756.200</td>
<td>Cervical rib</td>
</tr>
<tr>
<td></td>
<td>755.500</td>
<td>Clinodactyly (in-curving of fifth finger)</td>
</tr>
<tr>
<td></td>
<td>752.520</td>
<td>Cryptorchidism (see undescended testicle)</td>
</tr>
<tr>
<td></td>
<td>277.010</td>
<td>Cystic fibrosis, with mention of meconium ileus</td>
</tr>
<tr>
<td></td>
<td>277.000</td>
<td>Cystic fibrosis, with no mention of meconium ileus</td>
</tr>
<tr>
<td></td>
<td>744.280</td>
<td>Darwin's tubercle</td>
</tr>
<tr>
<td>1/1/96 T</td>
<td>754.030</td>
<td>Dolichocephaly - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Downward eye slant (antimongoloid)</td>
</tr>
<tr>
<td></td>
<td>744.110</td>
<td>Ear tags, preauricular</td>
</tr>
<tr>
<td></td>
<td>744.120</td>
<td>Ear tags, other</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Elfin ear, absent or decreased ear cartilage - if &lt;36 weeks gestation, code only if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Epicanttal folds</td>
</tr>
<tr>
<td></td>
<td>767.600</td>
<td>Erb's palsy</td>
</tr>
<tr>
<td></td>
<td>368.000</td>
<td>Esotropia</td>
</tr>
<tr>
<td></td>
<td>378.000</td>
<td>Exotropia</td>
</tr>
<tr>
<td></td>
<td>351.000</td>
<td>Facial palsy</td>
</tr>
<tr>
<td></td>
<td>757.380</td>
<td>Flammeus nevus or port wine stain</td>
</tr>
<tr>
<td></td>
<td>748.180</td>
<td>Flat bridge of nose</td>
</tr>
<tr>
<td></td>
<td>754.040</td>
<td>Fontanelle (large or small)</td>
</tr>
<tr>
<td></td>
<td>743.630</td>
<td>Fused eyelids - never code if &lt;25 weeks gestation</td>
</tr>
</tbody>
</table>
EXCLUSION LIST for the MACDP
Nonreportable birth defects

**Alphabetical - Conditions Which may be Included Under Certain Conditions**

<table>
<thead>
<tr>
<th>Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>3/4/91</td>
<td>752.440</td>
<td>Fusion of vulva</td>
</tr>
<tr>
<td>3/4/91</td>
<td>282.200</td>
<td>Glucose-6-phosphate dehydrogenase (G-6-PD) deficiency</td>
</tr>
<tr>
<td>3/4/91</td>
<td>271.000</td>
<td>Glycogen storage disease</td>
</tr>
<tr>
<td>3/4/91</td>
<td>748.510</td>
<td>Hypoplasia of lung; pulmonary hypoplasia - exclude only if an isolated defect in infants &lt;36 weeks gestation</td>
</tr>
<tr>
<td>1/1/96</td>
<td>T 752.440</td>
<td>Hypoplastic labia majora - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>3/4/91</td>
<td>T 748.510</td>
<td>Hypoplastic scrotum - exclude if secondary to undescended testes</td>
</tr>
<tr>
<td>1/1/96</td>
<td>T 243.990</td>
<td>Hypothyroidism, congenital (Exclude hypothyroidism of prematurity in infants &lt;36 weeks gestation even if other reportable defects are present. Include other types of hypothyroidism and hypothyroidism NOS when another reportable defect is present regardless of gestational age)</td>
</tr>
<tr>
<td>1/1/96</td>
<td>752.430</td>
<td>Imperforate hymen</td>
</tr>
<tr>
<td>1/1/96</td>
<td>755.500</td>
<td>Incurving fingers (clinodactyly)</td>
</tr>
<tr>
<td>T</td>
<td>550.000-550.901</td>
<td>Inguinal hernia or patent processus vaginalis. Never code in infants &lt;36 weeks gestation regardless of the presence of a reportable defect. For infants ≥36 weeks:</td>
</tr>
<tr>
<td>T</td>
<td>550.902</td>
<td>In males, code only if another reportable defect is present;</td>
</tr>
<tr>
<td></td>
<td></td>
<td>In females, always code even if found in isolation</td>
</tr>
<tr>
<td>3/4/91</td>
<td>757.450</td>
<td>Lanugo, excessive or persistent</td>
</tr>
<tr>
<td>3/4/91</td>
<td>754.040</td>
<td>Large fontanelle</td>
</tr>
<tr>
<td>3/4/91</td>
<td>755.500</td>
<td>Long fingers and toes</td>
</tr>
<tr>
<td>3/4/91</td>
<td>744.245</td>
<td>Low set ears</td>
</tr>
<tr>
<td>3/4/91</td>
<td>744.820</td>
<td>Macrocheilia (big lips)</td>
</tr>
<tr>
<td>3/4/91</td>
<td>270.300</td>
<td>Maple syrup urine disease</td>
</tr>
<tr>
<td>3/4/91</td>
<td>751.010</td>
<td>Meckel's diverticulum</td>
</tr>
<tr>
<td>3/4/91</td>
<td>777.600</td>
<td>Meconium peritonitis</td>
</tr>
<tr>
<td>3/4/91</td>
<td>777.100</td>
<td>Meconium plug</td>
</tr>
</tbody>
</table>
### Alphabetical - Conditions Which may be Included Under Certain Conditions

<table>
<thead>
<tr>
<th>Revised/Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>9/10/90</td>
<td>754.520</td>
<td>Metatarsus varus or adductus</td>
</tr>
<tr>
<td></td>
<td></td>
<td><strong>EXCLUSION LIST</strong> for the MACDP</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Nonreportable birth defects</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7/10/90</td>
<td>744.830</td>
<td>Microcheilia (small lips)</td>
</tr>
<tr>
<td>10/1/92</td>
<td>T</td>
<td>Mitral valve insufficiency or regurgitation, congenital</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Code cases designated as &quot;mild&quot;, minimal&quot;, 'trivial', or 'physiologic' only</td>
</tr>
<tr>
<td></td>
<td></td>
<td>if another reportable defect is present. Code all other degrees of</td>
</tr>
<tr>
<td></td>
<td></td>
<td>insufficiency or regurgitation, including those where the degree is not</td>
</tr>
<tr>
<td></td>
<td></td>
<td>specified, regardless of whether another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td>757.386</td>
<td>Mongolian spots</td>
</tr>
<tr>
<td></td>
<td>743.650</td>
<td>Nasal lacrimal duct obstruction</td>
</tr>
<tr>
<td></td>
<td>520.600</td>
<td>Natal teeth</td>
</tr>
<tr>
<td></td>
<td>745.500</td>
<td>Nonclosure of foramen ovale, NOS (see PFO)</td>
</tr>
<tr>
<td></td>
<td>379.500</td>
<td>Nystagmus</td>
</tr>
<tr>
<td>9/10/90</td>
<td>756.080</td>
<td>Occiput, flat or prominent</td>
</tr>
<tr>
<td>3/5/90</td>
<td>457.800</td>
<td>Other specified disorder of lymphatics, including</td>
</tr>
<tr>
<td></td>
<td></td>
<td>chylothorax</td>
</tr>
<tr>
<td></td>
<td>755.600</td>
<td>Overlapping toes</td>
</tr>
<tr>
<td>10/14/92</td>
<td>T</td>
<td>Patent ductus arteriosus (PDA)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1) Always code if ≥36 weeks of gestation and defect last noted ≥6 weeks of</td>
</tr>
<tr>
<td></td>
<td></td>
<td>age.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2) If ≥36 weeks gestation and defect last noted &lt;6 weeks of age, code only</td>
</tr>
<tr>
<td></td>
<td></td>
<td>if the PDA was treated (e.g. by ligation or indomethacin) or if another</td>
</tr>
<tr>
<td></td>
<td></td>
<td>reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3) Never code if &lt;36 weeks gestation or if treated with prostaglandins</td>
</tr>
<tr>
<td></td>
<td></td>
<td>regardless of gestational age.</td>
</tr>
<tr>
<td>10/14/92</td>
<td>T</td>
<td>Nonclosure of foramen ovale, NOS (see PFO)</td>
</tr>
<tr>
<td></td>
<td>#</td>
<td>Patent foramen ovale (PFO)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1) Always code if ≥36 weeks of gestation and defect last noted ≥6 weeks of</td>
</tr>
<tr>
<td></td>
<td></td>
<td>age.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2) If ≥36 weeks gestation and defect last noted &lt;6 weeks of age, code only</td>
</tr>
<tr>
<td></td>
<td></td>
<td>if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3) Never code if &lt;36 weeks gestation regardless of presence of other defects</td>
</tr>
<tr>
<td></td>
<td>753.700</td>
<td>Patent urachus</td>
</tr>
<tr>
<td></td>
<td>744.820</td>
<td>Patulous lips (wide lips)</td>
</tr>
<tr>
<td>8/1/93</td>
<td>747.325</td>
<td>Peripheral pulmonic stenosis (PPS) murmur - do collect if PPS documented</td>
</tr>
<tr>
<td></td>
<td></td>
<td>by echocardiogram</td>
</tr>
<tr>
<td></td>
<td>270.100</td>
<td>Phenylketonuria (PKU)</td>
</tr>
<tr>
<td></td>
<td>685.100</td>
<td>Pilonidal or sacral dimple</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Pixie-like ear</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Pointed ear</td>
</tr>
<tr>
<td></td>
<td>755.006</td>
<td>Polydactyly in blacks (postaxial, type B), including skin tags on hands or</td>
</tr>
<tr>
<td></td>
<td></td>
<td>feet</td>
</tr>
<tr>
<td></td>
<td>744.246</td>
<td>Posteriorly rotated ears</td>
</tr>
<tr>
<td></td>
<td>744.410</td>
<td>Preauricular sinus, cyst or pit</td>
</tr>
<tr>
<td></td>
<td>744.110</td>
<td>Preauricular tags</td>
</tr>
<tr>
<td></td>
<td>747.680</td>
<td>Primary pulmonary artery hypertension</td>
</tr>
</tbody>
</table>

= Rev. 6/94
* = code created by CDC
# = on the MACDP Excl List
### Alphabetical - Conditions Which may be Included Under Certain Conditions

<table>
<thead>
<tr>
<th>Revised/ Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>10/1/92 T 277.630</td>
<td></td>
<td>Pseudocholinesterase enzyme deficiency</td>
</tr>
<tr>
<td>10/1/92 T 746.020</td>
<td></td>
<td>Pulmonary valve insufficiency or regurgitation, congenital</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
</tr>
<tr>
<td>1/1/96 T 750.500</td>
<td></td>
<td>Pylorospasm (intermittent pyloric stenosis)</td>
</tr>
<tr>
<td>1/1/96 T 751.580</td>
<td></td>
<td>Rectal fissures</td>
</tr>
<tr>
<td>1/1/96 T 284.000</td>
<td></td>
<td>Red cell aplasia</td>
</tr>
<tr>
<td>1/1/96 T 744.500</td>
<td></td>
<td>Redundant neck skin folds</td>
</tr>
<tr>
<td>1/1/96 T 755.616</td>
<td></td>
<td>rocker-bottom feet</td>
</tr>
<tr>
<td>1/1/96 T 685.100</td>
<td></td>
<td>Sacral dimple</td>
</tr>
<tr>
<td>1/1/96 T 754.060</td>
<td></td>
<td>Scaphocephaly, no mention of craniosynostosis</td>
</tr>
<tr>
<td>#</td>
<td></td>
<td>Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>1/1/96 T 754.640</td>
<td></td>
<td>Small nipple (hypoplastic)</td>
</tr>
<tr>
<td>#</td>
<td></td>
<td>Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>10/1/92 T 759.020</td>
<td></td>
<td>Splenomegaly</td>
</tr>
<tr>
<td>7/13/92 090.000</td>
<td></td>
<td>Syphilis, congenital</td>
</tr>
<tr>
<td>7/13/92 759.240</td>
<td></td>
<td>Thymic hypertrophy</td>
</tr>
<tr>
<td>7/13/92 755.630</td>
<td></td>
<td>Tibial torsion</td>
</tr>
<tr>
<td>7/13/92 750.000</td>
<td></td>
<td>Tongue-tie</td>
</tr>
<tr>
<td>10/1/92 T 608.200</td>
<td></td>
<td>Torsion of spermatic cord</td>
</tr>
<tr>
<td>10/1/92 T 746.105</td>
<td></td>
<td>Tricuspid valve insufficiency or regurgitation, congenital</td>
</tr>
<tr>
<td>#</td>
<td></td>
<td>Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
</tr>
<tr>
<td>10/1/92 T 759.900</td>
<td></td>
<td>Umbilical cord atrophy</td>
</tr>
<tr>
<td>553.100</td>
<td></td>
<td>Umbilical hernias (completely covered by skin)</td>
</tr>
</tbody>
</table>
EXCLUSION LIST for the MACDP
Nonreportable birth defects

Alphabetical - Conditions Which may be Included Under Certain Conditions

<table>
<thead>
<tr>
<th>Revised/ Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1/1/93</td>
<td>T 752.500-</td>
<td>Undescended testicle (cryptorchidism)</td>
</tr>
<tr>
<td></td>
<td>T 752.520</td>
<td>1) If &lt; 36 weeks gestation, code only if there is a medical/surgical intervention for this problem;</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2) If ≥36 weeks gestation and defect last noted at &lt;1 year of age, code only if there was a medical/surgical intervention for this problem or if another reportable defect is present</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3) Always code if ≥36 weeks gestation and defect first noted at ≥1 of age.</td>
</tr>
<tr>
<td></td>
<td>748.180</td>
<td>Upturned nose</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Upward eye slant (mongoloid)</td>
</tr>
<tr>
<td></td>
<td>752.460</td>
<td>Vaginal cysts</td>
</tr>
<tr>
<td></td>
<td>752.480</td>
<td>Vaginal tags</td>
</tr>
<tr>
<td></td>
<td>286.400</td>
<td>von Willebrand's disease</td>
</tr>
<tr>
<td>3/14/91</td>
<td>T 755.130</td>
<td>Webbed toes</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Code webbing of the second and third toes only if another reportable defect is present. Always code webbing of other toes regardless of whether another reportable defect is present</td>
</tr>
<tr>
<td></td>
<td>744.500</td>
<td>Webbing of neck</td>
</tr>
<tr>
<td></td>
<td>748.180</td>
<td>Wide nasal bridge</td>
</tr>
<tr>
<td></td>
<td>755.600</td>
<td>Widely spaced first and second toes</td>
</tr>
<tr>
<td></td>
<td>757.680</td>
<td>Widely spaced nipples</td>
</tr>
</tbody>
</table>
### EXCLUSION LIST for the MACDP

**Numerical** list of conditions requiring no record abstraction unless associated with a reportable defect. The addition or revision dates of the changes in the list of conditions requiring no record abstraction are shown.

<table>
<thead>
<tr>
<th>Revised/Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>7/13/92</td>
<td>090.000</td>
<td>Syphilis congenital</td>
</tr>
<tr>
<td></td>
<td>216</td>
<td>Benign neoplasm of skin</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(NOTE: All benign neoplasms should be coded ONLY if another reportable code is present)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Includes:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>blue nevus</td>
</tr>
<tr>
<td></td>
<td></td>
<td>pigmented nevus</td>
</tr>
<tr>
<td></td>
<td></td>
<td>papilloma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>dermatofibroma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>syringoadenoma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>*dermoid cyst</td>
</tr>
<tr>
<td></td>
<td></td>
<td>hydrocystoma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>syringoma</td>
</tr>
<tr>
<td>#</td>
<td>216.000</td>
<td>Skin of lip</td>
</tr>
<tr>
<td>#</td>
<td>216.100</td>
<td>Eyelid, including canthus</td>
</tr>
<tr>
<td>#</td>
<td>216.200</td>
<td>Ear and external auditory canal</td>
</tr>
<tr>
<td>#</td>
<td>216.300</td>
<td>Skin of other and unspecified parts of face</td>
</tr>
<tr>
<td>#</td>
<td>216.400</td>
<td>Scalp and skin of neck</td>
</tr>
<tr>
<td>#</td>
<td>216.500</td>
<td>Skin of trunk, except scrotum</td>
</tr>
<tr>
<td>#</td>
<td>216.600</td>
<td>Skin of upper limb, shoulder</td>
</tr>
<tr>
<td>#</td>
<td>216.700</td>
<td>Skin of lower limb, hip</td>
</tr>
<tr>
<td>#</td>
<td>216.800</td>
<td>Other specified sites of skin</td>
</tr>
<tr>
<td>#</td>
<td>216.900</td>
<td>Site unspecified</td>
</tr>
<tr>
<td>#</td>
<td>216.910</td>
<td>Sebaceous cyst</td>
</tr>
</tbody>
</table>

* = code created by CDC
# = on the MACDP Excl List
= Rev. 6/94
**EXCLUSION LIST** for the MACDP

Numerical list of conditions requiring no record abstraction unless associated with a reportable defect. The addition or revision dates of the changes in the list of conditions requiring no record abstraction are shown.

<table>
<thead>
<tr>
<th>Revised/Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>221.000</td>
<td>221.000</td>
<td>Benign skin neoplasm of female genital organs</td>
</tr>
<tr>
<td>222.000</td>
<td>222.000</td>
<td>Benign skin neoplasm of male genital organs</td>
</tr>
<tr>
<td>T 243.990</td>
<td></td>
<td>Hypothyroidism, congenital (Exclude even if other defects are present only if the record specifies hypothyroidism of prematurity &lt;36 weeks. Include other types of hypothyroidism and hypothyroidism NOS only when another reportable defect is present regardless of gestational age.</td>
</tr>
<tr>
<td>251.200</td>
<td></td>
<td>Hypoglycemia, idiopathic</td>
</tr>
<tr>
<td>252.100</td>
<td></td>
<td>Hypoparathyroidism, congenital</td>
</tr>
<tr>
<td>270.100</td>
<td></td>
<td>Phenylketonuria (PKU)</td>
</tr>
<tr>
<td>270.200</td>
<td></td>
<td>Albinism</td>
</tr>
<tr>
<td>270.300</td>
<td></td>
<td>Maple syrup urine disease</td>
</tr>
<tr>
<td>270.600</td>
<td></td>
<td>Argininosuccinic aciduria</td>
</tr>
<tr>
<td>270.700</td>
<td></td>
<td>Hyperglycinemia</td>
</tr>
<tr>
<td>271.000</td>
<td></td>
<td>Glycogen storage diseases</td>
</tr>
<tr>
<td>275.330</td>
<td></td>
<td>Hypophosphatemic rickets</td>
</tr>
<tr>
<td>277.000</td>
<td></td>
<td>Cystic fibrosis, with no mention of meconium ileus</td>
</tr>
<tr>
<td>277.010</td>
<td></td>
<td>Cystic fibrosis, with mention of meconium ileus</td>
</tr>
<tr>
<td>277.620</td>
<td></td>
<td>Alpha 1-antitrypsin deficiency</td>
</tr>
<tr>
<td>277.630</td>
<td></td>
<td>Pseudocholinesterase enzyme deficiency</td>
</tr>
<tr>
<td>282.000</td>
<td></td>
<td>Hereditary spherocytosis</td>
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<tr>
<td>282.100</td>
<td></td>
<td>Hereditary elliptocytosis</td>
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<tr>
<td>282.200</td>
<td></td>
<td>Glucose-6-phosphate dehydrogenase (G-6-PD) deficiency</td>
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<tr>
<td>282.600</td>
<td></td>
<td>Sickle cell anemia</td>
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<tr>
<td>284.000</td>
<td></td>
<td>Red cell aplasia</td>
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<tr>
<td>286.000</td>
<td></td>
<td>Hemophilia</td>
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<tr>
<td>286.400</td>
<td></td>
<td>von Willebrand's disease</td>
</tr>
<tr>
<td>330.100</td>
<td></td>
<td>Cerebral lipidoses (e.g., Tay-Sachs, gangliosidoses, etc.)</td>
</tr>
<tr>
<td>351.000</td>
<td></td>
<td>Facial palsy</td>
</tr>
<tr>
<td>368.000</td>
<td></td>
<td>Esotropia</td>
</tr>
<tr>
<td>378.000</td>
<td></td>
<td>Exotropia</td>
</tr>
<tr>
<td>379.500</td>
<td></td>
<td>Nystagmus</td>
</tr>
<tr>
<td>3/5/90</td>
<td>457.800</td>
<td>Other specified disorder of lymphatics, including chylothorax</td>
</tr>
<tr>
<td>520.600</td>
<td></td>
<td>Natal teeth</td>
</tr>
<tr>
<td>T 550.000-550.900</td>
<td></td>
<td>Inguinal hernia or patent processus vaginalis never code in infants if &lt;36 weeks gestation regardless of the presence of a reportable defect.</td>
</tr>
<tr>
<td>NOTE: for those ≥36 weeks:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>in <strong>males</strong>, code only if another reportable defect is present;</td>
<td></td>
<td></td>
</tr>
<tr>
<td>in <strong>females</strong>, always code even if found in isolation</td>
<td></td>
<td></td>
</tr>
<tr>
<td>553.100</td>
<td></td>
<td>Umbilical hernias (completely covered by skin)</td>
</tr>
<tr>
<td>608.200</td>
<td></td>
<td>Torsion of spermatic cord</td>
</tr>
<tr>
<td>608.200</td>
<td></td>
<td>Torsion of testes</td>
</tr>
<tr>
<td>658.800</td>
<td></td>
<td>Amniotic bands (constricting bands, amniotic cyst)</td>
</tr>
<tr>
<td>685.100</td>
<td></td>
<td>Pilonidal or sacral dimple</td>
</tr>
</tbody>
</table>

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# = on the MACDP Excl List

= Rev. 6/94

Revised 6/93 (EXCL1088)
Replaces 6/89 Exclusion List
**Exclusion List** for the MACDP  
Nonreportable birth defects

**Numerical - Conditions Which may be Included Under Certain Conditions**

<table>
<thead>
<tr>
<th>Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>743.450</td>
<td>Blue sclera - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td></td>
<td>743.630</td>
<td>Fused eyelids - never code if &lt;25 weeks gestation unless another reportable defect is present</td>
</tr>
<tr>
<td></td>
<td>743.650</td>
<td>Nasal lacrimal duct obstruction</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Brushfield spots</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Downward eye slant (antimongoloid)</td>
</tr>
<tr>
<td></td>
<td>743.800</td>
<td>Upward eye slant (mongoloid)</td>
</tr>
<tr>
<td></td>
<td>744.100</td>
<td>Accessory auricle</td>
</tr>
<tr>
<td></td>
<td>744.110</td>
<td>Ear tags, preauricular</td>
</tr>
<tr>
<td></td>
<td>744.120</td>
<td>Ear tags, other</td>
</tr>
<tr>
<td></td>
<td>744.220</td>
<td>Bat ear</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Cauliflower ear</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Elfin ear, absent or decreased ear cartilage If &lt;36 weeks gestation, code only if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Lop ear</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Pixie-like ear</td>
</tr>
<tr>
<td></td>
<td>744.230</td>
<td>Pointed ear</td>
</tr>
<tr>
<td></td>
<td>744.245</td>
<td>Low set ears</td>
</tr>
<tr>
<td></td>
<td>744.246</td>
<td>Posteriorly rotated ears</td>
</tr>
<tr>
<td></td>
<td>744.280</td>
<td>Darwin's tubercle</td>
</tr>
<tr>
<td></td>
<td>744.410</td>
<td>Preauricular sinus, cyst or pit</td>
</tr>
<tr>
<td></td>
<td>744.500</td>
<td>Redundant neck skin folds</td>
</tr>
<tr>
<td></td>
<td>745.500</td>
<td>Webbing of neck</td>
</tr>
<tr>
<td></td>
<td>744.820</td>
<td>Macrocheilia (big lips)</td>
</tr>
<tr>
<td></td>
<td>744.820</td>
<td>Patulous lips (wide lips)</td>
</tr>
<tr>
<td></td>
<td>744.830</td>
<td>Microcheilia (small lips)</td>
</tr>
<tr>
<td></td>
<td>744.900</td>
<td>Short neck</td>
</tr>
<tr>
<td></td>
<td>745.500</td>
<td>Nonclosure of foramen ovale, NOS (see PFO)</td>
</tr>
<tr>
<td>10/14/92</td>
<td>745.500</td>
<td>Patent foramen ovale (PFO)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1) Always code if ≥36 weeks of gestation and defect last noted at ≥6 weeks of age.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2) If ≥36 weeks gestation and defect last noted &lt;6 weeks of age, code only if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3) Never code if &lt;36 weeks gestation regardless of presence of other defects.</td>
</tr>
</tbody>
</table>
| 10/1/92  | 746.020 | Pulmonary valve insufficiency or regurgitation, congenital - Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.
### EXCLUSION LIST for the MACDP

**Nonreportable birth defects**

<table>
<thead>
<tr>
<th>Revised/Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>10/1/92 T 746.105</td>
<td>Tricuspid valve insufficiency or regurgitation, congenital - Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
<td></td>
</tr>
<tr>
<td>10/1/92 T 746.400</td>
<td>Aortic valve insufficiency or regurgitation, congenital - Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
<td></td>
</tr>
<tr>
<td>10/1/92 T 746.600</td>
<td>Mitral valve insufficiency or regurgitation, congenital - Code cases designated as 'mild', minimal', 'trivial', or 'physiologic' only if another reportable defect is present. Code all other degrees of insufficiency or regurgitation, including those where the degree is not specified, regardless of whether another reportable defect is present.</td>
<td></td>
</tr>
<tr>
<td>10/1/92 T 746.860</td>
<td>Cardiomegaly, congenital NOS</td>
<td></td>
</tr>
<tr>
<td>10/1/92 T 746.990</td>
<td>Heart murmur - if chart review does not confirm a heart defect within 6 months, do not code as a defect even if other codable defects are present</td>
<td></td>
</tr>
<tr>
<td>10/14/92 T 747.000</td>
<td>Patent ductus arteriosus (PDA) 1)Always code if ≥36 weeks of gestation and defect last noted at ≥6 weeks of age. 2)If ≥36 weeks gestation and defect last noted &lt;6 weeks of age, code only if the PDA was treated (e.g., by ligation or indomethicin) or if another reportable defect is present. 3)Never code if &lt;36 weeks gestation or if treated with prostaglandins regardless of gestational age.</td>
<td></td>
</tr>
<tr>
<td>8/1/93 747.325</td>
<td>Peripheral pulmonic stenosis (PPS) murmur - do collect if PPS documented by echocardiogram</td>
<td></td>
</tr>
<tr>
<td>8/1/93 747.500</td>
<td>Single umbilical artery</td>
<td></td>
</tr>
<tr>
<td>8/1/93 747.680</td>
<td>Primary pulmonary artery hypertension</td>
<td></td>
</tr>
<tr>
<td>8/1/93 778.000</td>
<td>Ascites or anasarca. Includes: hydrops fetalis</td>
<td></td>
</tr>
<tr>
<td>8/1/93 748.180</td>
<td>Flat bridge of nose</td>
<td></td>
</tr>
<tr>
<td>8/1/93 748.180</td>
<td>Upturned nose</td>
<td></td>
</tr>
<tr>
<td>8/1/93 748.180</td>
<td>Wide nasal bridge</td>
<td></td>
</tr>
<tr>
<td>3/4/91 T 748.510</td>
<td>Hypoplasia of lung; pulmonary hypoplasia - exclude if isolated defect in infants &lt;36 weeks gestation. Tongue-tie</td>
<td></td>
</tr>
<tr>
<td>3/4/91 750.000</td>
<td>High arched palate</td>
<td></td>
</tr>
<tr>
<td>3/4/91 750.240</td>
<td>Pylorospasm (intermittent pyloric stenosis)</td>
<td></td>
</tr>
<tr>
<td>3/4/91 751.010</td>
<td>Meckel's diverticulum</td>
<td></td>
</tr>
</tbody>
</table>
### EXCLUSION LIST for the MACDP
Nonreportable birth defects

#### Numerical - Conditions Which may be Included Under Certain Conditions

<table>
<thead>
<tr>
<th>Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1/1/96</td>
<td>T 752.440</td>
<td>Hypoplastic labia majora - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>3/4/91</td>
<td>752.450</td>
<td>Prominent clitoris</td>
</tr>
<tr>
<td>1/1/93</td>
<td>T 752.460</td>
<td>Vaginal cysts</td>
</tr>
<tr>
<td>1/1/93</td>
<td>T 752.450</td>
<td>Vaginal tags</td>
</tr>
<tr>
<td>1/1/93</td>
<td>752.480</td>
<td>Hymenal tags</td>
</tr>
<tr>
<td>1/1/93</td>
<td>T 752.500-</td>
<td>Undescended testicle (cryptorchidism)</td>
</tr>
<tr>
<td>1/1/93</td>
<td>T 752.520</td>
<td>1) If &lt; 36 weeks gestation, code only if there is a medical/surgical intervention for this problem; 2) If ≥36 weeks gestation and defect last noted at &lt;1 year of age, code only if there was a medical/surgical intervention for this problem or if another reportable defect is present 3) Always code if ≥36 weeks gestation and defect first noted at ≥1 of age.</td>
</tr>
<tr>
<td>1/1/93</td>
<td>752.520</td>
<td>Cryptorchidism (see undescended testicle)</td>
</tr>
<tr>
<td>1/1/93</td>
<td>752.810</td>
<td>Hypoplastic scrotum - exclude if secondary to undescended testes</td>
</tr>
<tr>
<td>1/1/96</td>
<td>753.700</td>
<td>Patent urachus</td>
</tr>
<tr>
<td>1/1/96</td>
<td>754.020</td>
<td>Bent nose, deviation of nasal septum</td>
</tr>
<tr>
<td>1/1/96</td>
<td>T 754.030</td>
<td>Dolichocephaly - if &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>1/1/93</td>
<td>754.040</td>
<td>Fontanelle (large or small)</td>
</tr>
<tr>
<td>1/1/96</td>
<td>754.060</td>
<td>Scaphocephaly, no mention of craniosynostosis If &lt;36 weeks gestation, code only if another reportable defect is present. Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td>1/1/93</td>
<td>754.520</td>
<td>Metatarsus varus or adductus</td>
</tr>
<tr>
<td>1/1/93</td>
<td>755.006</td>
<td>Polydactyly in blacks (postaxial, type B), including skin tags on hands or feet</td>
</tr>
<tr>
<td>3/14/91</td>
<td>T 755.130</td>
<td>Webbed toes Code webbing of the second and third toes only if another reportable defect is present. Always code webbing of other toes regardless of whether another reportable defect is present</td>
</tr>
<tr>
<td>755.500</td>
<td></td>
<td>Clinodactyly (incurving of fifth finger)</td>
</tr>
<tr>
<td>755.500</td>
<td></td>
<td>Long fingers and toes</td>
</tr>
<tr>
<td>755.600</td>
<td></td>
<td>Overlapping toes</td>
</tr>
<tr>
<td>755.600</td>
<td></td>
<td>Widely spaced first and second toes</td>
</tr>
<tr>
<td>755.616</td>
<td></td>
<td>Rocker-bottom feet</td>
</tr>
<tr>
<td>755.630</td>
<td></td>
<td>Tibial torsion</td>
</tr>
<tr>
<td>756.080</td>
<td></td>
<td>Occiput, flat or prominent</td>
</tr>
<tr>
<td>756.200</td>
<td></td>
<td>Cervical rib</td>
</tr>
<tr>
<td>757.200</td>
<td></td>
<td>Sidney line</td>
</tr>
<tr>
<td>757.200</td>
<td></td>
<td>Simian crease (transverse palmar crease)</td>
</tr>
</tbody>
</table>
**EXCLUSION LIST** for the MACDP
Nonreportable birth defects

**Numerical – Conditions Which may be Included Under Certain Conditions**

<table>
<thead>
<tr>
<th>Revised/Changed Date</th>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>1/1/96 T</td>
<td>757.310</td>
<td>Anal tags</td>
</tr>
<tr>
<td></td>
<td>757.380</td>
<td>Flammeus nevus or port wine stain</td>
</tr>
<tr>
<td></td>
<td>757.385</td>
<td>Birth mark, NOS</td>
</tr>
<tr>
<td></td>
<td>757.386</td>
<td>Mongolian spots</td>
</tr>
<tr>
<td></td>
<td>757.390</td>
<td>Cafe au lait spots</td>
</tr>
<tr>
<td></td>
<td>757.390</td>
<td>Skin cysts</td>
</tr>
<tr>
<td></td>
<td>757.450</td>
<td>Lanugo, excessive or persistent</td>
</tr>
<tr>
<td>9/10/90</td>
<td>757.640</td>
<td>Small nipple (hypoplastic)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>If &lt;36 weeks gestation, code only if another reportable defect is present.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Always code if ≥36 weeks gestation.</td>
</tr>
<tr>
<td></td>
<td>757.650</td>
<td>Accessory nipple (supernumerary nipple, or skin tag)</td>
</tr>
<tr>
<td></td>
<td>757.680</td>
<td>Widely spaced nipples</td>
</tr>
<tr>
<td></td>
<td>759.020</td>
<td>Splenomegaly</td>
</tr>
<tr>
<td></td>
<td>759.240</td>
<td>Thymic hypertrophy</td>
</tr>
<tr>
<td></td>
<td>759.900</td>
<td>Umbilical cord atrophy</td>
</tr>
<tr>
<td></td>
<td>767.600</td>
<td>Erb's palsy</td>
</tr>
<tr>
<td></td>
<td>777.100</td>
<td>Meconium plug</td>
</tr>
<tr>
<td></td>
<td>777.600</td>
<td>Meconium peritonitis</td>
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<tr>
<td></td>
<td>778.000</td>
<td>Ascites or anasarca, congenital</td>
</tr>
<tr>
<td></td>
<td>778.600</td>
<td>Hydrocele, congenital</td>
</tr>
</tbody>
</table>

= Rev. 6/94
* = code created by CDC
# = on the MACDP Excl List
MACDP Decision Tree for Determining Whether to Include Patent Ductus Arteriosus (PDA)

Is the child on prostaglandins?  -------> Yes  -------> Never code
  |  
  |  No
  |  
  |  What is the gestational age?  -------> < 36 wks ---> Never code
  |  
  |  > 36 wks
  |  
  |  How old was the child when defect was last noted?  -------> > 6 wks ----> Always code
  |  
  |  < 6 wks
  |  
  |  Has the PDA been treated? (e.g., by ligation or indomethacin)  -------> Yes  -------> Always code
  |  
  |  No
  |  
Include if other defects are present, i.e., exclusion list.
MACDP Decision Tree for Determining Whether to Include Patent Foramen Ovale (PFO)

What is the gestational age?  
---------->  < 36 wks ---> Never code  

|  
|  
|  
|  
> 36 wks  

|  
|  
|  
|  
How old was the child when defect was last noted?  
---------->  > 6 wks ---> Always code  

|  
|  
|  
|  
< 6 wks  

|  
|  
|  
Include if other defects are present, i.e., exclusion list.

May 22, 1996