

## ICD-9 CM codes relevant to the diagnosis of Down syndrome\*

>>> indicates intervening codes have been left of this list

### 14. CONGENITAL ANOMALIES (740-759)

#### 758 Chromosomal anomalies

Includes: syndromes associated with anomalies in the number and form of chromosomes

Use additional codes for conditions associated with the chromosomal anomalies

##### 758.0 Down's syndrome

Mongolism

Translocation Down's syndrome

Trisomy:

21 or 22

G

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### 3. ENDOCRINE, NUTRITIONAL & METABOLIC DISEASES, & IMMUNITY DISORDERS (240-279)

#### DISORDERS OF THYROID GLAND (240-246)

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#### 244 Acquired hypothyroidism

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##### 244.9 Unspecified hypothyroidism

Hypothyroidism, primary or NOS

Myxedema, primary or NOS

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### NEUROTIC DISORDERS, PERSONALITY DISORDERS, AND OTHER NONPSYCHOTIC MENTAL DISORDERS (300-316)

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#### 315 Specific delays in development

**Excludes:** that due to a neurological disorder (320.0-389.9)

##### 315.0 Specific reading disorder

315.00 Reading disorder, unspecified

315.01 Alexia

315.02 Developmental dyslexia

315.09 Other

Specific spelling difficulty

##### 315.1 Mathematics disorder

Dyscalculia

##### 315.2 Other specific learning difficulties

Disorder of written expression

**Excludes:** specific arithmetical disorder (315.1)

specific reading disorder (315.00-315.09)

- 315.3 Developmental speech or language disorder
  - 315.31 Expressive language disorder
    - Developmental aphasia
    - Word deafness
  - Excludes:**
    - acquired aphasia (784.3)
    - elective mutism (309.83, 313.0, 313.23)
  - 315.32 Mixed receptive-expressive language disorder
  - 315.39 Other
    - Developmental articulation disorder
    - Dyslalia
    - Phonological disorder
  - Excludes:**
    - lisping and lalling (307.9)
    - stammering and stuttering (307.0)
- 315.4 Developmental coordination disorder
  - Clumsiness syndrome
  - Dyspraxia syndrome
  - Specific motor development disorder
- 315.5 Mixed development disorder
- 315.8 Other specified delays in development
- 315.9 Unspecified delay in development
  - Developmental disorder NOS
  - Learning disorder NOS

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#### OTHER DISEASES OF DIGESTIVE SYSTEM (570-579)

#### 579 Intestinal malabsorption

- 579.0 Celiac disease
  - Celiac:
    - crisis
    - infantilism
    - rickets
  - Gee (-Herter) disease
  - Gluten enteropathy
  - Idiopathic steatorrhea
  - Nontropical sprue

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#### OTHER DISEASES OF SKIN AND SUBCUTANEOUS TISSUE (700-709)

#### 704 Diseases of hair and hair follicles

**Excludes:** congenital anomalies (757.4)

- 704.0 Alopecia
  - Excludes:**
    - madarosis (374.55)
    - syphilitic alopecia (091.82)

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- 704.01 Alopecia areata
  - Ophiasis

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#### 14. CONGENITAL ANOMALIES (740-759)

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745 Bulbus cordis anomalies and anomalies of cardiac septal closure

- 745.0 Common truncus
  - Absent septum between aorta and pulmonary artery
  - Communication (abnormal) between aorta and pulmonary artery
  - Aortic septal defect
  - Common aortopulmonary trunk
  - Persistent truncus arteriosus

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- 745.6 Endocardial cushion defects
  - 745.60 Endocardial cushion defect, unspecified type
  - 745.61 Ostium primum defect
    - Persistent ostium primum
  - 745.69 Other
    - Absence of atrial septum
    - Atrioventricular canal type ventricular septal defect
    - Common atrioventricular canal
    - Common atrium

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751 Other congenital anomalies of digestive system

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- 751.1 Atresia and stenosis of small intestine
  - Atresia of:
    - duodenum
    - ileum
    - intestine NOS
  - Congenital:
    - absence of small intestine or intestine NOS
    - obstruction of small intestine or intestine NOS
    - stenosis of small intestine or intestine NOS
    - stricture of small intestine or intestine NOS
  - Imperforate jejunum

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- 751.3 Hirschsprung's disease and other congenital functional disorders of colon
  - Aganglionosis
  - Congenital dilation of colon
  - Congenital megacolon
  - Marcrocolon

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**16. SYMPTOMS, SIGNS, AND ILL-DEFINED CONDITIONS (780-799)**

783 Symptoms concerning nutrition, metabolism, and development

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- 783.3 Feeding difficulties and mismanagement
    - Feeding problem (elderly) (infant)
- Excludes:** feeding disturbance or problems:
  - in newborn (779.3)
  - of nonorganic origin (307.50-307.59)

- 783.4 Lack of expected normal physiological development in childhood  
**Excludes:** delay in sexual development and puberty (259.0)  
 gonadal dysgenesis (758.6)  
 pituitary dwarfism (253.3)  
 slow fetal growth and fetal malnutrition (764.00-764.99)  
 specific delays in mental development (315.0-315.9)
- 783.40 Lack of normal physiological development, unspecified  
 Inadequate development  
 Lack of development
- 783.41 Failure to thrive  
 Failure to gain weight
- 783.42 Delayed milestones  
 Late talker  
 Late walker

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### DISLOCATION (830-839)

**Includes:** displacement  
 subluxation

**Excludes:** congenital dislocation (754.0-755.8)  
 pathological dislocation (718.2)  
 recurrent dislocation (718.3)

The descriptions "closed" and "open," used in the fourth-digit subdivisions, include the following terms:

closed:

complete  
 dislocation NOS  
 partial  
 simple  
 uncomplicated

open:

compound  
 infected  
 with foreign body

**Note:** A dislocation not indicated as closed or open should be classified as closed.

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### 839 Other, multiple, and ill-defined dislocations

- 839.0 Cervical vertebra, closed  
 Cervical spine  
 Neck
- 839.00 Cervical vertebra, unspecified
- 839.01 First cervical vertebra
- 839.02 Second cervical vertebra

\* Abstracted from International Classification of Diseases, 9<sup>th</sup> revision, Clinical Modification, issued by the US Department of Health and Human Services. Available in electronic versions from <http://www.cdc.gov/nchs/icd9.htm>. Published copies of ICD-9-CM are available from a variety of sources and should be found in any medical library.