7th DIGIT

PROPOSAL

Code	Definition
1	Syndrome or disease
2	Known association
3	Sequence or named complex malformation
4	Component of syndrome, association or sequence or complex malformation
5	Major
6	Major same system
7	Secondary to major anomaly
8	Independant minor
9	Unknown

EUROCAT GUIDE 5

Classification and Coding of Congenital Anomalies

An E.E.C. Concerted Action Project

EUROCAT Central Registry
Department of Epidemiology
Catholic University of Louvain
Brussels, 1990

PRINCIPLES FOR THE NEW EUROCAT CODING SYSTEM.

1. It is mainly a nomenclature and a coding system, not a classification system, but essential improvements in ciassification are made.

2. It is an extension to 6 digits of the British Paediatric Association Classification of Disease and should therefore be compatible with the BPA system.

3. Coding should be as specific as possible. When not already used in the BPA system, new fifth digits can also be assigned. For subcategories of an anomaly already coded to 5 digits in the BPA system, a sixth digit should be assigned.

4. The general rules for the last digit of each code are:

- 0 is not used, (if O is given in the PBA code as a fifth digit, add a 1 as sixth digit)
- 1 to 7 are used for specific conditions,
- 8 is used for 'other specified',
- 9 is used for unspecified.

5. As many synonymns as possible should be included to allow coders to find the nomenclature to which they are accustomed.

6. Syndromes, diseases, associations and sequences with onset of symptoms at birth or during infancy are included.

7. Conditions with Mendelian inheritance are included along with their McKusick code number.

8. Definitions and remarks on exclusions and inclusions should be given where necessary to allow a coder with minimum medical training to translate descriptions in medical records to the appropriate codes.

I. INFECTIOUS AND PARASITIC DISEASES

Exclude all codes in this chapter
Congenital Syphilis see code 7701.64 (chapter XV)

II. NEOPLASMS

77125

Malignant neoplasm of major salivary glands

Excludes : malignant neoplasms of minor salivary glands, which are to be classified according to their anatomical location; if location is not specified, classify to 1429.

1420.-- Parotid gland
1421.-- Submandibular gland
Submaillary gland
1422.-- Sublingual gland
1428.-- Other
1429.-- Site unspecified
Salivary gland (major) NOS

Malignant neoplasm of colon

1530.-- Hepatic flexure
1531.-- Transverse colon
1532.-- Descending colon
1533.-- Sigmoid colon
Sigmoid (flexure)
Excludes : rectosigmoid junction (1540)
1534.-- Caecum
Ileocaecal valve
1535.-- Appendix
1536.-- Ascending colon
1537.-- Splenic flexure
1538.-- Other
1539.-- Colon, unspecified
Large intestine NOS

Malignant neoplasm of liver and intrahepatic bile ducts

1550.-- Liver, primary
Carcinoma : liver, specified as primary
hepatocellular
liver cell
Hepatoblastoma
1551.-- Intrahepatic bile ducts
Canaliculi biliferi
Interlobular : bile ducts
biliary canals
Intrahepatic : canaliculi
biliary passages
bile duct
1552.-- Liver, not specified as primary or secondary

Malignant neoplasm of gallbladder and extrahepatic bile ducts

1560.-- Gallbladder
1561.-- Extrahepatic bile ducts

Biliary duct or passage NOS
Common bile duct
Cystic duct
Hepatic duct
1562.-- Ampulla of Vater
1568.-- Other
1569.-- Biliary tract, part unspecified
Malignant neoplasm involving both intrahepatic and extrahepatic bile ducts

Malignant neoplasm of pancreas

1570.-- Head of pancreas
1571.-- Body of pancreas
1572.-- Tail of pancreas
1573.-- Pancreatic duct
Ducts of : Santorini
Wirsung
1574.-- Islets of Langerhans
Islets of Langerhans, any part of pancreas
1578.-- Other
1579.-- Part unspecified

Malignant neoplasm of retroperitoneum and peritoneum

1580.-- Retroperitoneum
1588.-- Specified parts of peritoneum
Mesentery
Mesocolon
Omentum
Peritoneum : parietal
pelvic
1589.-- Peritoneum, unspecified

Malignant neoplasm of larynx

1610.-- Glottis
Intrinsic larynx
Laryngeal commissure (anterior)
Laryngeal commissure (posterior)
True vocal cord
Vocal cord NOS
1611.-- Supraglottis
Aryepiglottic fold or interarytenoid fold, laryngeal aspect
Epiglottis (suprahyoid portion) NOS
Extrinsic larynx
False vocal cords
Posterior (laryngeal) surface of epiglottis
Ventricular bands
Excludes : anterior aspect of epiglottis (1464)
aryepiglottic fold or interarytenoid fold :
NOS (1482)
hypopharyngeal aspect (1482)
marginal zone (1482)
1612.-- Subglottis
1613.-- Laryngeal cartilages
Cartilage : arytenoid
cricoid

Cartilage : cuneiform
thyroid

1618.-- Other
1619.-- Larynx, unspecified

Malignant neoplasm of trachea, bronchus and lung

1620.-- Trachea
1622.-- Main bronchus
Carina
Hilus
1623.-- Upper lobe, bronchus or lung
1624.-- Middle lobe, bronchus or lung
1625.-- Lower lobe, bronchus or lung
1628.-- Other
1629.-- Bronchus and lung, unspecified

Malignant neoplasm of thymus, heart and mediastinum

1640.-- Thymus
1641.-- Heart
Pericardium
Excludes : great vessels (1714)
1642.-- Anterior mediastinum
1643.-- Posterior mediastinum
1648.-- Other
1649.-- Mediastinum, part unspecified

Malignant neoplasm of bone and articular cartilage

Excludes : bone marrow NOS (2029)
cartilage : ear (1710)
eyelid (1710)
larynx (1613)
nose (1600)
synovia (171-)

1700.-- Bones of skull and face
Bone : ethmoid
frontal
malar
occipital
orbital
parietal
sphenoid
temporal
Maxilla
Turbinate
Vomer

Excludes : carcinoma, any type other than intraosseous or odontogenic :
maxilla, maxillary (sinus) (1602)
upper jaw bone (1430)
jaw bone (lower) (1701)

1701.-- Lower jaw bone
Mandible
Jaw bone NOS
Excludes : carcinoma, any type other than intraosseous or odontogenic :
jaw bone NOS (1439)
lower (1431)
upper jaw bone (1700)

1702.-- Vertebral column, excluding sacrum and coccyx
Spinal column
Spine
Vertebra

1703.-- Ribs, sternum and clavicle
Costal cartilage
Xiphoid process

1704.-- Long bones of upper limb and scapula
Acromion process
Bone NOS of upper limb
Humerus
Radius
Ulna

1705.-- Upper limb, short bones
Carpal
Cuneiform, wrist
Pisiform
Scaphoid (of hand)
Semilunar or lunate
Trapezium
Trapezoid
Unciform
Metacarpal
Phalanges of hand

1706.-- Pelvic bones, sacrum and coccyx
Coccygeal vertebra
Ilium
Ischium
Pubic bone
Sacral vertebra

1707.-- Lower limb, long bones
Bones NOS of lower limb
Femur
Fibula
Tibia

1708.-- Lower limb, short bones
Patella
Tarsal
Astragalus
Calcaneus
Cuboid
Cuneiform, ankle
Navicular (of ankle)
Metatarsal
Phalanges of foot

1709.-- Site unspecified
Chondrosarcoma, site unspecified
Chondromyxosarcoma, site unspecified
Ewing's sarcoma or tumour, site unspecified
Osteosarcoma, site unspecified

Malignant neoplasm of connective and other soft tissue

Includes : blood vessel
bursa
fascia
ligament, exept uterine
muscle
peripheral, sympathetic and parasympathetic nerves and ganglia
synovia
tendon (sheath)

Excludes : cartilage (of) : articular (170-)
larynx (1613)
nose (1600)
connective tissue of breast (174-)
heart (1641)

1710.-- Head, face and neck
Cartilage of : ear
eyelid

1712.-- Upper limb, including shoulder

1713.-- Lower limb, including hip

1714.-- Thorax
Axilla
Diaphragm
Great vessels

Excludes : thymus, heart and mediastinum (164-)

1715.-- Abdomen
Abdominal wall
Hypochondrium

1716.-- Pelvis
Buttock
Groin
Perineum
Excludes : uterine ligament, any (183-)

1717.-- Trunk, unspecified
Back NOS

1718.-- Other

1719.-- Site unspecified
Fibrosarcoma, site unspecified
Haemangiosarcoma, site unspecified
Leiomysarcoma, site unspecified
Liposarcoma, site unspecified
Lymphangiosarcoma, site unspecified
Myosarcoma, site unspecified
Myxosarcoma, site unspecified
Rhabdomyosarcoma, site unspecified
Sarcoma, site unspecified
Sarcomatosis, site unspecified

Other malignant neoplasm of skin

Includes : malignant neopalsm of :
sebaceous glands
sweat glands

Excludes : malignant melanoma of skin (172-)
skin of genital organs (184-, 187-)

1730.-- Skin of lip
Excludes : vermilion border of lip (140-)

1731.-- Eyelid, including canthus
Excludes : cartilage of eyelid (1710)

1732.-- Ear and external auricular canal
Auricle (ear)
External meatus
Pinna
Excludes : cartilage of ear (1710)

1733.-- Skin of other and unspecified parts of face
Cheek, external
Eyebrow
Nose, external
Temple

1734.-- Scalp and skin of neck

1735.-- Skin of trunk, except scrotum
Axillary fold
Perianal skin
Umbilicus
Skin of :
abdominal wall
anus
back
Skin of :
breast
buttock
chest wall
groin
perineum

Excludes : anal canal (1542)
anus NOS (1543)
skin of scrotum (1877)

1736.-- Skin of upper limb, including shoulder

1737.-- Skin of lower limb, including hip

1738.-- Other

1739.-- Site unspecified

Malignant neoplasm of ovary and other uterine adnexa

1830.-- Ovary
1832.-- Fallopian tube
Oviduct
Uterine tube
1833.-- Broad ligament
1834.-- Parametrium
Uterine ligament NOS
1835.-- Round ligament
1838.-- Other
1839.-- Uterine adnexa, unspecified

Malignant neoplasm of testis

1860.-- Undescended
Ectopic testis
Retained testis
1869.-- Other and unspecified
Testis :
NOS
descended
scrotal

Malignant neoplasm of kidney and other and unspecified urinary organs

1890.-- Kidney, except pelvis/Wilms' tumour(?)
1891.-- Renal pelvis
Pelviureteric junction
Renal calyces
1892.-- Ureter
Excludes : ureteric orifice of bladder (1886)
1893.-- Urethra
Excludes : urethral orifice of bladder (1885)
1894.-- Paraurethral glands
1898.-- Other
1899.-- Site unspecified
Urinary system NOS

Malignant neoplasm of brain

Excludes: cranial nerves (192-)
retrobulbar (1901)
1910.-- Cerebrum, except lobes and ventricles
1911.-- Frontal lobe
1912.-- Temporal lobe
1913.-- Parietal lobe
1914.-- Occipital lobe
1915.-- Ventricle
Floor of ventricle
1916.-- Cerebellum
1917.-- Brain stem
1918.-- Other
1919.-- Brain, unspecified
Astrocytoma, site unspecified
Astroblastoma, site unspecified

Ependymoma, site unspecified
Glioma, site unspecified
Medulloblastoma, site unspecified
Neuroepithelioma, site unspecified
Oligodendroblastoma, site unspecified

Malignant neoplasm of other and unspecified parts of nervous system

Excludes : peripheral, sympathetic and parasympathetic nerves and ganglia (171-)

1920.-- Cranial nerves
1921.-- Cerebral meninges
Meninges NOS
1922.-- Spinal cord
1923.-- Spinal meninges
1928.-- Other
1929.-- Part unspecified
Nervous system (central) NOS
Excludes : meninges NOS (1921)

Malignant neoplasm of other endocrine glands and related structures

Excludes : islets of Langerhans (1574)

1940.-- Suprarenal gland
Adrenal gland, neuroblastoma ?
1941.-- Parathyroid gland
1943.-- Pituitary gland and craniopharyngeal duct
1944.-- Pineal gland
1945.-- Carotid body
1946.-- Aortic body and other paraganglia
1948.-- Other
Pluriglandular involvement NOS
NOTE: If the sites of multiple involvement are known, they should be coded separately.
1949.-- Site unspecified
Endocrine gland NOS

Malignant neoplasm of other and ill-defined sites

Includes : overlapping neoplasms, not elsewhere classified.
Excludes : malignant neoplasm :
lymphatic and haematopoietic tissue (200-208)
unspecified site (199-)

1950.-- Head, face and neck
Cheek NOS
Nose NOS
1951.-- Thorax
Axilla
1952.-- Abdomen
1953.-- Pelvis
Groin
Sites overlapping systems within the pelvis, such as :
rectovesical (septum)
rectovaginal (septum)
1954.-- Upper limb
1955.-- Lower limb
1958.-- Other specified sites

Malignant neoplasm without specification of site

1990.-- Disseminated
Carcinomatosis, unspecified site (primary) (secondary)
Generalised :
cancer, unspecified site (primary) (secondary)
malignancy, unspecified site (primary) (secondary)
Multiple cancer, unspecified site (primary) (secondary)
1991.-- Other
Carcinoma, unspecified site (primary) (secondary)
Cancer, unspecified site (primary) (secondary)
Malignancy, unspecified site (primary) (secondary)

Lymphosarcoma and reticulosarcoma

2000.-- Reticulosarcoma
2001.-- Lymphosarcoma
Excludes : lymphosarcoma-cell leukaemia (2078)
2002.-- Burkitt's tumour
2008.-- Other named variants
Reticulolymphosarcoma

Hodgkin's disease

NOTE: Two alternative sub-classifications are given :
.0-.2 Parker-Jackson
.4-.7 Rye modification of Lukes-Butler
This departure from the principle that categories should be mutually exclusive is deliberate, since both form of terminology are currently encountered on medical records.

2010.-- Hodgkin's paragranuloma
2011.-- Hodgkin's granuloma
2012.-- Hodgkin's sarcoma
2014.-- Lymphocytic-histiocytic predominance
2015.-- Nodular sclerosis
2016.-- Mixed cellularity
2017.-- Lymphocytic depletion
2019.-- Unspecified

Other malignant neoplasm of lymphoid and histiocytic tissue

2020.-- Nodular lymphoma
Brill-Symmers disease
Follicular lymphoma
2021.-- Mycosis fungoides
2022.-- Sézary's disease
2023.-- Malignant histiocytosis
2024.-- Leukaemic reticuloendotheliosis
Hairy-cell leukaemia
2025.-- Letterer-Siwe disease
Acute differentiated progressive histiocytosis
Acute infantile reticuloendotheliosis
Acute (progressive) histiocytosis X
Acute reticulosis of infancy
2026.-- Malignant mast-cell tumours
Malignant :
mastocytoma
mastocytosis

Mast-cell sarcoma
Excludes : mast-cell leukaemia (2078)

2028.-- Other lymphomas
Lymphoma (malignant) :
NOS
diffuse

2029.-- Other and unspecified

Multiple myeloma and immunoproliferative neoplasms

2030.-- Multiple myeloma
Kahler's disease
Myelomatosis
Excludes : solitary myeloma (2386)

2031.-- Plasma cell leukaemia

2038.-- Other immunoproliferative neoplasms

Lymphoid leukaemia

Includes : leukaemia : lymphatic
lymphocytic

2040.-- Acute
Excludes : acute exacerbation of chronic lymphoid leukaemia (2041)

2041.-- Chronic

2042.-- Subacute

2048.-- Other

2049.-- Unspecified

Myeloid leukaemia

Includes : leukaemia : granulocytic
myelogenous

2050.-- Acute
Excludes : acute exacerbation of chronic myeloid leukaemia (2051)

2051.-- Chronic

2052.-- Subacute

2053.-- Myeloid sarcoma
Chloroma
Granulocytic sarcoma

2058.-- Other

2059.-- Unspecified

Monocytic leukaemia

Includes : monocytoid leukaemia

2060.-- Acute
Excludes : acute exacerbation of chronic monocytic leukaemia (2061)

2061.-- Chronic

2062.-- Subacute

2068.-- Other

2069.-- Unspecified

Other specified leukaemia

Excludes : leukaemic reticuloendotheliosis (2024)
plasma cell leukaemia (2031)

2070.-- Acute erythraemia and erythroleukaemia
Acute erythraemic myelosis
Di Guglielmo's disease
2071.-- Chronic erythraemia
Heilmeyer-Schöner disease
2072.-- Megakaryocytic leukaemia
2078.-- Other
Lymphosarcoma cell leukaemia

Leukaemia of unspecified cell type

2080.-- Acute
Acute leukaemia NOS
Stem cell leukaemia
Blast cell leukaemia
Excludes : acute exacerbation of unspecified leukaemia (2081)
2081.-- Chronic
Chronic leukaemia NOS
2082.-- Subacute
Subacute leukaemia NOS
2088.-- Other
2089.-- Unspecified
Leukaemia NOS

Benign neoplasm of lip, oral cavity and pharynx

Excludes: cyst of jaw (522-, 526-)
cyst of oral soft tissue (528-)
2100.-- Lip
Frenulum labii
Lip (inner aspect) (mucosa) (vermilion border)
Excludes: labial commissure (2104)
skin of lip (2160)
2101.-- Tongue
Lingual tonsil
2102.-- Major salivary glands
Gland: parotid
sublingual
submandibular
Excludes: benign neoplasms of minor salivary glands which are to be classified according to their anatomical location; if location is not specified, classify to 2104.
2103.-- Floor of mouth
2104.-- Other and unspecified parts of mouth
Gingiva — Oral mucosa
Labial commissure — Palate (hard) (soft)
Oral cavity NOS — Uvula
Excludes: benign odontogenic neoplasms (213-)
mucosa of lips (2100)
nasopharyngeal (posterior) (superior) surface of soft palate (2107)
2105.-- Tonsil
Tonsil (faucial) (palatine)
Excludes: lingual tonsil (2101)
pharyngeal tonsil (2107)
tonsillar : fossa (2106)
pillars (2106)

2106.-- Other parts of oropharynx
Epiglottis, anterior aspect
Tonsillar : fossa
pillars
Vallecula
Excludes : epiglottis : NOS (2121)
suprahyoid portion (2121)
2107.-- Nasopharynx
2108.-- Hypopharynx
2109.-- Pharynx, unspecified

Benign neoplasm of other parts of digestive system

2110.-- Oesophagus
2111.-- Stomach
2112.-- Small intestine, including duodenum
Excludes : ampulla of Vater (2115)
ileocaecal valve (2113)
2113.-- Colon
Excludes : rectosigmoid junction (2114)
2114.-- Rectum and anal canal
Anus NOS
Excludes : anus : margin (2165)
skin (2165)
perianal skin (2165)
2115.-- Liver and biliary passages
2116.-- Pancreas, except islets of Langerhans
2117.-- Islets of Langerhans
Islet cell tumour
2118.-- Retroperitoneum and peritoneum
2119.-- Other and unspecified site
Alimentary canal or tract NOS
Gastrointestinal tract NOS
Intestine NOS
Spleen, not elsewhere classified

Benign neoplasm of respiratory and intrathoracic organs

2120.-- Nasal cavities, middle ear and accessory sinuses
Excludes : auricular canal (external) (2162)
bone of : ear (2130)
nose (2130)
cartilage of ear (2150)
ear (external) (skin) (2162)
nose NOS (2298)
skin (2163)
olfactory bulb (2251)
polyp of : accessory sinus (4718)
ear (3879)
nasal cavity (4710)
posterior margin of septum and choanae (2107)
2121.-- Larynx
Excludes : epiglottis, anterior aspect (2106)
polyp of vocal cord or larynx (4784)
2122.-- Trachea
2123.-- Bronchus and lung
2124.-- Pleura
2125.-- Mediastinum

2126.-- Thymus
2127.-- Heart
Excludes : great vessels (2154)
2128.-- Other specified sites
2129.-- Site unspecified
Respiratory organ NOS
Upper respiratory tract NOS
Excludes: intrathoracic NOS (2298)
thoracic NOS (2298)

Benign neoplasm of bone and articular cartilage

Excludes: cartilage of : ear (2150)
eyelid (2150)
larynx (2121)
nose (2120)
synovia (215-)

2130.-- Bones of skull and face
Excludes : lower jaw bone (2131)
2131.-- Lower jaw bone
2132.-- Vertebral column, excluding sacrum and coccyx
2133.-- Ribs, sternum and clavicle
2134.-- Long bones of upper limb and scapula
2135.-- Upper limb, short bones
2136.-- Pelvic bones, sacrum and coccyx
2137.-- Lower limb, long bones
2138.-- Lower limb, short bones
2139.-- Site unspecified
Chondroma, site unspecified
Osteoma, site unspecified

Other benign neoplasm of connective tissue and other soft tissue

Includes : blood vessel
bursa
fascia
ligament
muscle
peripheral, sympathetic and parasympathetic nerves and ganglia
synovia
tendon (sheath)

Excludes: cartilage : articular (213-)
larynx (2121)
nose (2120)
connective tissue of breast (217)

2150.-- Head, face and neck
2152.-- Upper limb, including shoulder
2153.-- Lower limb, including hip
2154.-- Thorax
Excludes: heart (2127)
mediastinum (2125)
thymus (2126)
2155.-- Abdomen
2156.-- Pelvis
Excludes: uterine : leiomyoma (218)
ligament, any (2210)
2157.-- Trunk, unspecified
Back NOS
2158.-- Other specified sites

2159.-- Site unspecified
Leiomyoma, site unspecified
Myoma, site unspecified
Myxofibroma, site unspecified
Myxoma, site unspecified
Rhabdomyoma, site unspecified

Benign neoplasm of skin

7573-81

Includes: blue naevus, dermatofibroma, hydrocystoma, pigmented naevus, syringoadenoma, syringoma
Excludes: skin of genital organs (211-, 222-)

2160.-- Skin of lip
Excludes : vermilion border of lip (2100)

2161.-- Eyelid, including canthus
Excludes : cartilage of eyelid (2150)

2162.-- Ear and external auricular canal
Auricle (ear)
External meatus
Pinna
Excludes: cartilage of ear (2150)

2163.-- Skin of other and unspecified parts of face
Cheek, external
Eyebrow
Nose, external
Temple

2164.-- Scalp and skin of neck

2165.-- Skin of trunk, except scrotum
Axillary fold
Perianal skin
Umbilicus
Skin of: abdominal wall, anus, back, breast, buttock, chest wall, groin, perineum
Excludes: anal canal (2114)
anus NOS (2114)
skin of scrotum (2224)

2166.-- Skin of upper limb, including shoulder

2167.-- Skin of lower limb, including hip

2168.-- Other

2169.-- Site unspecified

Benign neoplasm of ovary

Benign neoplasm of male genital organs

2220.-- Testis

2221.-- Penis
Corpus cavernosus
Glans penis
Prepuce

2222.-- Prostate
Excludes: adenomatous hyperplasia ofprostate (600)
prostatic : adenoma (600)
enlargement (600)
hypertrophy (600)
2223.-- Epididymis
2224.-- Scrotum
Skin of scrotum
2228.-- Other specified sites
Seminal vesicle
Spermatic cord
2229.-- Site unspecified

Benign neoplasm of kidney and other urinary organs

2230.-- Kidney, except pelvis
Excludes: renal : calyces (2231)
pelvis (2231)
2231.-- Bladder
2232.-- Ureter
Excludes : ureteric orifice of bladder (2233)
2233.-- Bladder
2238.-- Other specified sites
Paraurethral glands
Urethra
Excludes : urethral orifice of bladder (2233)
2239.-- Site unspecified
Urinary system NOS

Benign neoplasm of brain and other parts of nervous system

Excludes: haemangioma (2280)
peripheral, sympathetic and parasympathetic nerves and ganglia (215-)
retrobulbar (2241)
2250.-- Brain
2251.-- Cranial nerves
2252.-- Cerebral meninges
Meninges NOS
Meningioma (cerebral)
2253.-- Spinal cord
2254.-- Spinal meninges
Spinal meningioma
2258.-- Other
2259.-- Part unspecified
Nervous system (central) NOS
Excludes : meninges NOS (2252)

Benign neoplasm of other endocrine glands and related structures

2270.-- Suprarenal gland
Adrenal gland
2271.-- Parathyroid gland
2273.-- Pituitary gland and craniopharyngeal duct
2274.-- Pineal gland
2275.-- Carotid body
2276.-- Aortic body and other paraganglia
2278.-- Other

2279.-- Site unspecified
Endocrine gland NOS

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173-84

Benign neoplasm of other and unspecified sites

2290.-- Lymph nodes
2298.-- Other specified sites
2299.-- Site unspecified

Carcinoma in situ of digestive organs

2300.-- Lip, oral cavity and pharynx
Gingiva — Oropharynx
Hypopharynx — Salivary gland or duct
Mouth (any part) — Tongue
Nasopharynx
Excludes : aryepiglottic fold or interarytenoid fold, laryngeal aspect (2310)
epiglottis : NOS (2310)
suprahyoid portion (2310)
skin of lip (2320)

2301.-- Oesophagus
2302.-- Stomach
2303.-- Colon
Excludes : rectosigmoid junction (2304)
2304.-- Rectum
2305.-- Anal canal
2306.-- Anus, unspecified
Excludes : anus : margin (2325)
skin (2325)
perianal skin (2325)
2307.-- Other and unspecified parts of intestine
Excludes : ampulla of Vater (2308)
2308.-- Liver and biliary system
2309.-- Other and unspecified digestive organs
Digestive organ NOS
Pancreas

Carcinoma in situ of respiratory system

2310.-- Larynx
Excludes : aryepiglottic fold or interarytenoid fold :
NOS (2300)
hypopharyngeal aspect (2300)
marginal zone (2300)
2311.-- Trachea
2312.-- Bronchus and lung
2318.-- Other specified parts
Accessory sinuses
Middle ear
Nasal cavities
Excludes : ear (external) (skin) (2322)
nose NOS (2348)
skin (2323)
2319.-- Part unspecified
Respiratory organ NOS

Carcinoma in situ of skin

Includes : pigment cells

2320.-- Skin of lip

Excludes : vermilion border of lip (2300)

2321.-- Eyelid, including canthus

2322.-- Ear and external auricular canal

2323.-- Skin of other and unspecified parts of face

2324.-- Scalp and skin of neck

2325.-- Skin of trunk, except scrotum

Anus, margin
Axillary fold
Perianal skin
Umbilicus

Skin of : abdominal wall, anus, back, breast, buttock, chest wall, groin, perineum

Excludes : anal canal (2305)
anus NOS (2306)
skin of genital organs (2333, 2235, 2336)

2326.-- Skin of upper limb, including shoulder

2327.-- Skin of lower limb, including hip

2328.-- Other specified sites

2329.-- Site unspecified

Carcinoma in situ of breast and genitourinary system

2330.-- Breast

Excludes : skin of breast (2325)

2331.-- Cervix uteri

2332.-- Other and unspecified parts of uterus

2333.-- Other and unspecified female genital organs

2334.-- Prostate

2335.-- Penis

2336.-- Other and unspecified male genital organs

2337.-- Bladder

2339.-- Other and unspecified urinary organs

Carcinoma in situ of other and unspecified sites

2340.-- Eye

Excludes : eyelid (skin) (2321)

2348.-- Other specified sites

Endocrine gland (any)

2349.-- Site unspecified

Carcinoma in situ NOS

Neoplasm of uncertain behaviour of digestive and respiratory systems

2350x.-- Major salivary glands

Gland : parotid
sublingual
submandibular

2351.-- Lip, oral cavity and pharynx

Gingiva
Hypopharynx
Mouth
Nasopharynx
Oropharynx
Tongue

Excludes: aryepiglottic fold or interarytenoid fold, laryngeal aspect (2356)
epiglottis :
NOS (2356)
suprahyoid portion (2356)
skin of lip (2382)

2352.-- Stomach, intstines and rectum

2353.-- Liver and biliary passages
Ampulla of Vater — Gallbladder
Bile ducts (any) — Liver

2354.-- Retroperitoneum and peritoneum

2355.-- Other and unspecified digestive organs
Anal : canal — Anus NOS
sphincter — Oesophagus
Pancreas
Excludes: anus : margin (2382)
skin (2382)
perianal skin (2382)

2356.-- Larynx
Excludes: aryepiglottic fold or interarytenoid fold :
NOS (2351)
hypopharyngeal (2351)
marginal zone (2351)

2357.-- Trachea, bronchus and lung

2358.-- Pleura, thymus and mediastinum

2359.-- Other and unspecified respiratory organs
Accessory sinuses — Nasal cavities
Middle ear — Respiratory organ NOS
Excludes: ear (external) (skin) (2382)
nose (2388)
skin (2382)

Neoplasm of uncertain behaviour of genitourinary organs

2360.-- Uterus

2361.-- Placenta
Chorioadenoma (destruens)
Malignant hydatid(iform) mole

2362.-- Ovary

2363.-- Other and unspecified female genital organs

2364.-- Testis

2365.-- Prostate

2366.-- Other and unspecified male genital organs

2367.-- Bladder

2369.-- Other and unspecified urinary organs
Kidney
Ureter
Urethra

Neoplasm of uncertain behaviour of endocrine glands and nervous system

2370.-- Pituitary gland and craniopharyngeal duct

2371.-- Pineal gland

2372.-- Suprarenal gland
Adrenal gland

2373.-- Paraganglia

2374.-- Other and unspecified endocrine glands
Parathyroid gland
Thyroid gland
2375.-- Brain and spinal cord
2376.-- Meninges
Meninges : NOS
cerebral
spinal
2377.-- Neurofibromatosis
von Recklinghausen's disease
2379.-- Other and unspecified parts of nervous system
Cranial nerves

Neoplasm of uncertain behaviour of other and unspecified sites and tissues

2380.-- Bone and articular cartilage
Excludes : cartilage : ear (2381)
eyelid (2381)
larynx (2356)
nose (2359)
synovia (2381)
2381.-- Connective and other soft tissue
Excludes : cartilage (of) : articular (2380)
larynx (2356)
nose (2359)
connective tissue of breast (2383)
2382.-- Skin
Excludes : anus NOS (2355)
skin of genital organs (2363, 2366)
vermilion border of lip (2351)
2383.-- Breast
Excludes : skin of breast (2382)
2384.-- Polycythaemia vera
2385.-- Histiocytic and mast cells
Mast-cell tumour NOS
Mastocytoma NOS
2386.-- Plasma cells
Plasmacytoma NOS
Solitary myeloma
2387.-- Other lymphatic and haematopoietic tissues
Disease :
lymphoproliferative (chronic) NOS
myeloproliferative (chronic) NOS
Idiopathic thrombocythaemia
Megakaryocytic myelosclerosis
Myelosclerosis with myeloid metaplasia
Panmyelosis (acute)
Excludes : myelofibrosis (2988)
myelosclerosis NOS (2898)
myelosis : NOS (2059)
megakaryocytic (2072)
2388.-- Other specified sites
Eye
Heart
Excludes : eyelid (skin) (2382)
cartilage (2381)

2389.-- Site unspecified

Neoplasm of unspecified nature

Includes : "growth" NOS
neoplasm NOS
new growth NOS
tumour NOS

2390.-- Digestive system
Excludes : anus : margin (2392)
skin (2392)
perianal skin (2392)

2391.-- Respiratory system

2392.-- Bone, soft tissue and skin
Excludes : anal canal (2390)
anus NOS (2390)
cartilage : larynx (2391)
nose (2391)
connective tissue of breast (2393)
skin of genital organs (2395)
vermilion border of lip (2398)

2393.-- Breast
Excludes : skin of breast (2392)

2394.-- Bladder

2395.-- Other genitourinary organs

2396.-- Brain
Excludes : cerebral meninges (2397)
cranial nerves (2397)

2397.-- Endocrine glands, and other parts of nervous system
Excludes : peripheral, sympathetic and parasympathetic nerves and ganglia (2392)

2398.-- Other specified sites
Excludes : eyelid (skin) (2392)
cartilage (2392)
great vessels (2392)
optic nerve (2397)

2399.-- Site unspecified

III. ENDOCRINE, NUTRITIONAL AND METABOLIC DISEASES AND IMMUNITY DISORDERS

Congenital hypothyroidism

2439.0- Pendred's syndrome
2439.1- Goitrous cretin
Excludes : congenital athyrosis (see code 7592.11)
Excludes : sublingual ectopic thyroid (see code 7592.12)
2439.8- Other specified congenital hypothyroidism
2439.9- Unspecified congenital hypothyroidism

Diabetes mellitus

2500.-- Diabetes mellitus without mention of complications
2501.-- Diabetes with ketoacidosis
2502.-- Diabetes with coma
2509.-- Diabetes with unspecified complications

Other disorders of pancreatic internal secretion

(2510.-- Hypoglycaemic coma)
2511.-- Other hyperinsulinism
2512.-- Hypoglycaemia, unspecified
2519.-- Unspecified disorders of pancreatic internal secretion

Disorders of parathyroid gland

2520.-- Hyperparathyroidism
2521.-- Hypoparathyroidism
2528.-- Other specified disorders of parathyroid
2529.-- Unspecified disorders of parathyroid

Disorders of the pituitary gland and its hypothalmic gland

2530.-- Acromegaly and gigantism
2531.-- Other anterior pituitary hyperfunction
2532.0- Panhypopituitarism idiopathic

Pituitary dwarfism
2533.0- Isolated deficiency growth hormone
2533.1- Other pituitary dwarfism

Other anterior pituitary disorders
2534.0- Deficiency of follicle-stimulating hormone (FSH)
2534.1- Deficiency of luteinising hormone (LH)
2534.2- Deficiency of thyroid-stimulating hormone (TSH)
2534.9- Unspecified disorders of anterior pituitary gland

Diabetes insipidus
2535.0- Diabetes insipidus due to disorders in the pituitary gland
2535.1- Other diabetes insipidus

Other disorders
2538.0- Cyst of Rathke's pouch
2538.1- Adiposogenital dystrophy
2538.2- Other disorders of pituitary

Disorders of adrenal glands

Cushing's syndrome
2550.-- Cushing's syndrome

Adreno genital disorders

Congenital adrenal hyperplasia
2552.0- Defective synthesis of 21 ß Hydroxylase
2552.1- Defective synthesis of 11 ß Hydroxylase
2552.2- Defective synthesis of 3 ß Hydroxysteroid dehydrogenase
2552.3- Defective synthesis of 17-20 Desmolase
2552.4- Defective synthesis of 17 Alpha hydroxylase
2552.5- Other adrenogenital syndromes with salt loss
2552.6- Other adrenogenital syndromes without mention of salt loss
2552.7- Precocious puberty with adrenocortical hyperfunction
2552.8- Virilism
2552.9- Other and unspecified adrenogenital disorders

Cortico adrenal insufficiency
2554.0- Addison's disease
2554.3- Hypoaldosteronism

Other types of dwarfism not elsewhere classified

2598.0- Progeria
2598.1- Cerebral gigantism
2598.2- Leprechaunism
2598.3- Werner's syndrome
2598.9- Other types of dwarfism

Disorders of amino-acid transport and metabolism

Excludes : abnormal findings without manifest disease (791-796)
disorders of purine and pyrimidine metabolism (2772)
gout (274-)

Disturbances of amino-acid transport
2700.0- Cystinosis
2700.1- Cystinuria
2700.2- Fanconi (-de Toni) (-Debre) syndrome
2700.3- Hartnup disease
2700.8- Other specified disturbances of amino-acid transport
2700.9- Unspecified disturbances of amino-acid transport

Phenylketonuria / Hyperphenylalaninaemia
2701.-- Phenylketonuria / Hyperphenylalaninaemia

Other disturbances of aromatic amino-acid metabolism
2702.0- Albinism
2702.1- Hypertyrosinaemia
2702.2- Tyrosinosis
2702.3- Tyrosinuria
2702.4- Alkaptonuria
2702.8- Other specified disturbances of aromatic amino-acid metabolism

2702.9- Unspecified disturbances of aromatic amino-acid metabolism

Disturbances of branched-chain amino-acid metabolism
2703.0- Leucinosis
2703.1- Maple-syrup-urine disease
2703.8- Other specified disturbances of branched-chain amino-acid metabolism
2703.9- Unspecified disturbances of branched-chain amino-acid metabolism

Disturbances of sulphur-bearing amino-acid metabolism
2704.0- Homocystinuria
2704.8- Other specified disturbances of sulphur-bearing amino-acid metabolism
2704.9- Unspecified disturbances of sulphur-bearing amino-acid metabolism

Disturbances of histidine metabolism
2705.0- Histidinaemia
2705.8- Other specified disturbances of histidine metabolism
2705.9- Unspecified disturbances of histidine metabolism

Disorders of urea cycle metabolism
2706.0- Arginosuccinic aciduria
2706.1- Hyperammonaemia
2706.8- Other specified disorders of urea cycle metabolism
2706.9- Unspecified disorders of urea cycle metabolism

Other disturbances of straight-chain amino-acid metabolism
2707.0- Hyperglycinaemia
2707.1- Methylmalonic acidaemia
2707.2- Acidaemia pipecolic
2707.8- Other specified disturbances of straight-chain amino-acid metabolism
2707.9- Unspecified disturbances of straight-chain amino-acid metabolism

Other disorders of amino-acid transport and metabolism
2708.0- Ethanolaminuria
2708.1- Hydroxyprolinaemia
2708.2- Hyperprolinaemia
2708.8- Other specified disorders of amino-acid transport and metabolism
2708.9- Unspecified disorders of amino-acid transport and metabolism

Unspecified disorders of amino-acid transport and metabolism
2709.-- Unspecified disorders of amino-acid transport and metabolism

Disorders of carbohydrate transport and metabolism

Glycogenosis
2710.0- Hepatorenal glycogenosis / Glucose-6-phosphatase deficiency / Glycogenosis, type I / Von Gierke's syndrome
2710.1- Generalised glycogenosis / Glycogenosis, type 2 / Pompe's disease
2710.2- Glycogenosis of liver and muscle / Deficiency of debrancher enzyme / Glycogenosis, type 3
2710.3- Glycogenosis associated with hepatic cirrhosis / Glycogenosis, type 4
2710.4- Glycogenosis, type 5 / McArdle's syndrome / Myophosphorylase deficiency
2710.8- Other specified types of glycogenosis
2710.9- Unspecified types of glycogenosis

Galactosaemia
2711.0- Galactose-I-phosphate uridyl transferase deficiency
2711.2- Deficiency of galactokinase
2711.8- Other specified disorders of galactose metabolism

2711.9- Unspecified disorders of galactose metabolism

Hereditary fructose intolerance
2712.-- Hereditary fructose intolerance

Intestinal disaccharidase deficiencies and disaccharide malabsorption
2713.0- Primary lactose intolerance
2713.1- Secondary lactose intolerance
2713.2- Glucose intolerance
2713.3- Glucose-galactose intolerance
2713.4- Sucrose intolerance
2713.5- Sucrose-isomaltose intolerance
2713.6- Disaccharidase deficiencies
2713.7- Sucrosuria
2713.8- Other specified disaccharide intolerance
2713.9- Unspecified disaccharide intolerance

Renal glycosuria
2714.-- Renal glycosuria

Other disorders of carbohydrate transport and metabolism
2718.0- Oxalosis
2718.1- Oxaluria, primary
2718.2- Other and unspecified oxaluria
2718.3- Aspartylglucosaminuria
2718.4- Fucosidosis
2718.8- Other specified disorders of carbohydrate transport and metabolism

Unspecified disorders of carbohydrate transport and metabolism
2719.-- Unspecified disorders of carbohydrate transport and metabolism

Disorders of lipoid metabolism

Pure hypercholesterolaemia
2720.0- Familial hypercholesterolaemia
2720.1- Hyper-beta-lipoproteinaemia
2720.2- Hyperlipidaemia, group A
2720.3- Low-density-lipoid-type (LDL) hyperlipoproteinaemia
2720.9- Other pure hypercholesterolaemia

Pure hyperglyceridaemia
2721.-- Pure hyperglyceridaemia

Mixed hyperlipidaemia
2722.-- Mixed hyperlipidaemia

Hyperchylomicronaemia
2723.-- Hyperchylomicronaemia

Other and unspecified hyperlipidaemia
2724.-- Other and unspecified hyperlipidaemia

Lipoprotein deficiencies
2725.-- Lipoprotein deficiencies

Lipodystrophy
2726.-- Lipodystrophy : complete or partial

Lipidoses
2727.-- Lipidoses

Other disorders of lipoid metabolism
2728.0- Lipase deficiency
2728.1- Steatosis
2728.2- Lipomatosis
2728.8- Other disorders of lipoid metabolism

Disorders of plasma protein metabolism
2730.-- Polyclonal hypergamma-globulinaemia
2732.-- Other paraproteinaemia
2733.-- Macroglobulibaemia
2738.-- Other disorders of plasma protein metabolism
2739.-- Unspecified disorders of plasma protein metabolism

<u>Disorders of mineral metabolism</u>

Disorders of iron metabolism
2750.0- Haemosiderosis, primary
2750.2- Haemochromatosis / Von Recklinghausen-Appelbaum disease
2750.8- Other specified disorders of iron metabolism
2750.9- Unspecified disorders of iron metabolism

Disorders of copper metabolism
2751.-- Disorders of copper metabolism/Wilson's disease

Disorders of magnesium metabolism
2752.0- Hypomagnesaemic tetany
2752.1- Other hypomagnesaemia
2752.2- Hypermagnesaemia

Disorders of phosphorus metabolism
2753.0- Vitamin-D-resistant rickets
2753.1- Hypophosphatasia rickets
2753.2- Hypophosphatasia, other
2753.3- Hypophosphataemia
2753.4- Hyperphosphatasia

Disorders of calcium metabolism
2754.0- Nephrocalcinosis
2754.1- Other calcinosis
2754.2- Pseudohypoparathyroidism
2754.4- Other hypocalcaemia
2754.5- Hypercalcaemia, familial benign
2754.6- Hypercalcaemia, idiopathic
2754.7- Other hypercalcaemia
2754.8- Hypercalciuria
2754.9- Other and unspecified disorders of calcium metabolism

Other disorders of mineral metabolism
2758.-- Other specified disorders of mineral metabolism

Unspecified disorders of mineral metabolism
2759.-- Unspecified disorders of mineral metabolism

2762.1- Acidemia metabolic NEC

Other and unspecified disorders of metabolism

Cystic fibrosis
2770.-- Cystic fibrosis

Disorders of porphyrin metabolism
2771.-- Disorders of porphyrin metabolism

Other disorders of purine and pyrimidine metabolism
2772.-- Other disorders of purine and pyrimidine metabolism

Amyloidosis
2773.0- Familial
2773.1- Nephropathic
2773.8- Other specified amyloid disease
2773.9- Unspecified amyloid disease

Disorders of bilirubin excretion
Excludes: hyperbilirubinaemia specific to the perinatal period (see code 774-)
2774.0- Crigler-Najjar syndrome
2774.1- Dubin-Johnson syndrome
2774.2- Gilbert's syndrome
2774.3- Rotor's syndrome
2774.4- Hyperbilirubinaemia, congenital other specified
2774.9- Hyperbilirubinaemia congenital unspecified

Mucopolysaccharidosis
2775.0- Mannosidosis
2775.1- Mucopolysaccharidosis, type I
2775.2- Mucopolysaccharidosis, type II
2775.3- Mucopolysaccharidosis, type III
2775.4- Mucopolysaccharidosis, type IV
2775.5- Pyknodysostosis
2775.6- Mucopolysaccharidosis, type VI
2775.7- Mucopolysaccharidosis, type VII
2775.8- Multiple sulphatases deficiency
2775.98 Other specified mucopolysaccharidosis
2775.99 Unspecified mucopolysaccharidosis

Other deficiencies of circulating enzymes
2776.0- Angio-oedema
2776.1- Alpha 1 -antitrypsin hepatitis
2776.2- Other alpha 1-antitrypsin deficiency
2776.8- Other specified deficiencies of circulating enzymes
2776.9- Unspecified deficiencies of circulating enzymes

Other specified disorders of metabolism
Excludes: Letterer-Siwe disease (see code 2025)
Histiocytosis X, acute (see code 2025)
2778.0- Enterokinase deficiency
2778.1- Trypsinogen deficiency
2778.2- Acatalasia
2778.3- Eosinophilic granuloma
2778.4- Histiocytosis X, chronic
2778.5- Histiocytosis X, unspecified
2778.6- Histiocytosis, unspecified
2778.7- Hand-Schuller-Christian syndrome

2778.8- Other specified disorders of metabolism

Unspecified disorders of metabolism
2779.-- Unspecified disorders of metabolism

Disorders involving the immune mechanism

Deficiency of humoral immunity
2790.0- Agammaglobulinaemia, congenital sex-linked
2790.1- Agammaglobulinaemia, X-linked
2790.2- IgA deficiency (selective)
2790.3- Agammaglobulinaemia, other and unspecified
2790.4- Hypogammaglobulinaemia
2790.5- Dysimmunoglobulinaemia
2790.8- Other deficiency of humoral immunity
2790.9- Unspecified deficiency of humoral immunity

Deficiency of cell-mediated immunity
2791.0- Di George sequence
2791.1- Wiskott-Aldrich syndrome
2791.2- Lymphocyte deficiency
2791.9- Other deficiency of cell-mediated immunity

Combined immune deficiency
2792.0- Swiss type agammaglobulinaemia
2792.1- Thymic dysplasia with immune deficiency
2792.8- Other combined immune deficiency
2792.9- Unspecified combined immune deficiency

Unspecified immune deficiency
2793.-- Unspecified immune deficiency

Autoimmune disease not elsewhere classified
2794.-- Autoimmune disease not elsewhere classified

Other disorders of the immune mechanism
2798.-- Other disorders of the immune mechanism

Unspecified disorders of the immune mechanism
2799.-- Unspecified disorders of the immune mechanism

IV. DISEASES OF BLOOD AND BLOOD-FORMING ORGANS

Hereditary haemolytic anaemias

Hereditary spherocytosis
2820.-- Hereditary spherocytosis/Acholioric jaundice(?)

Hereditary elliptocytosis
2821.-- Hereditary elliptocytosis

Anaemia due to disorders of glutathione metabolism
2822.0- Glucose-6-phosphate dehydrogenase deficiency anaemia
2822.1- Enzyme deficiency drug induced anaemia
2822.2- Favism
2822.8- Other specified anaemia due to disorders of glutathione metabolism
2822.9- Unspecified anaemia due to disorders of glutathione metabolism

Other haemolytic anaemia due to enzyme deficiency
2823.-- Other haemolytic anaemia due to enzyme deficiency

Thalassaemias
2824.0- Alpha thalassaemia
2824.1- Alpha thalassaemia trait
2824.2- Beta thalassaemia trait
2824.3- Beta intermedia thalassaemia
2824.4- Beta major thalassaemia
2824.5- Beta minor thalassaemia
2824.6- Sickle-cell thalassaemia
2824.7- Minor thalassaemia, other and unspecified
2824.8- Major thalassaemia, other and unspecified
2824.9- Thalassaemia unspecified

Sickle-cell trait
2825.-- Sickle-cell trait

Sickle-cell anaemia
2826.-- Sickle-cell anaemia

Other haemoglobinopathies
2827.0- Hb-C disease
2827.1- Hb-D disease
2827.2- Hb-E disease
2827.3- Hereditary persistence of fetal haemoglobin
2827.4- Hb-H disease
2827.5- Heinz body anaemia, congenital
2827.9- Other haemoglobinopathies

Other specified hereditary haemolytic anaemias
2828.-- Other specified hereditary haemolytic anaemias

Unspecified hereditary haemolytic anaemies
2829.-- Unspecified hereditary haemolytic anaemies

Aplastic anemia

2840.0- Constitutional aplastic anaemia without mention of malformation
2840.1- Constitutional aplastic anaemia with malformation

2840.2- Red-cell aplasia and hypoplasia, constitutional
2840.8- Other specified constitutional aplastic anaemia
2840.9- Unspecified constitutional aplastic anaemia

Coagulation defects

2860.-- Congenital factor VIII disorder/Haemophilia A
2861.-- Congenital factor IX disorder / Christmas disease/Haemophilia B
2862.-- Congenital factor XI disorder

Congenital deficiency of other clotting factors
2863.0- Afibrin ogenaemia, congenital
2863.8- Other specified congenital deficiency of clotting factors
2863.9- Unspecified congenital deficiency of clotting factors
2864.-- von Willebrand's disease
2866.-- Defibrination syndrome

Purpura and other haemorrhagic conditions

Qualitative platelet defects

2871.1- Glanzmann's syndrome
2871.8- Other specified qualitative platelet defects
2871.9- Unspecified qualitative platelet defects

Primary thrombocytopenia

2873.1- Congenital thrombocytopenic purpura
2873.2- Thrombocytopenia with t radius/TAR syndrome
2873.8- Other primary thrombocytopenia

Diseases of white blood cells

2880.0- Congenital neutropenia / Agranulocytosis congenitale
2880.4- Cyclical neutropenia
2881.-- Functional disorders of neutrophil polymorphonuclear cells
2882.-- Genetic anomaly of leucocytes
2888.8- Other specified anomalies of white blood cells
2888.9- Unspecified anomalies of white blood cells

Other diseases of blood and blood-forming organs

2896.-- Familial polycythaemia
2897.0- Methaemoglobinaemia congenital
2899.-- Other diseases of blood and blood-forming organs

V. MENTAL DISORDERS

Exclude all codes in this chapter

VI. DISEASES OF THE NERVOUS SYSTEM AND SENSE ORGANS

Spinocerebellar disease

3340.-- Friedreich's ataxia
3341.-- Hereditary spastic paraplegia
3342.-- Primary cerebellar degeneration
3343.-- Other cerebellar ataxia
3344.-- Cerebellar ataxia in diseases classified elsewhere
3348.-- Other specified spinocerebellar disease
3349.-- Unspecified spinocerebellar disease

Anterior horn cell disease

3350.-- Werdnig-Hoffmann disease/Infantile spinal muscular atrophy
3351.-- Spinal muscular atrophy / Kugelberg-Welander disease
3352.-- Motor neurone disease
3358.-- Other specified anterior horn cell disease
3359.-- Unspecified anterior horn cell disease
Excludes : syringomyelia (see code 7425.5)

Disorders of the autonomic nervous system, unspecified

3379.-- Disorders of the autonomic nervous system, unspecified
Excludes : familial dysautonomia (Riley-Day syndrome) (see code 7428)

Other Disorders of the Central Nervous System (see codes340-349)
Excludes: Arachnoïd cyst (see code 7424.---)
Moebius syndrome (see code 7428.--)
3431.-- Congenital hemiplegia

Myoneural disorders

Myasthenia gravis
3580.0- Neonatal persistent myasthenia
3580.1- Adult or juvenile myasthenia
3580.9- Unspecified myasthenia gravis

3581.-- Myasthenic syndromes in diseases classified elsewhere

3588.1- Amyotonia congenita
3588.2- Congenital benign hypotonia / Oppenheim syndrome
3588.3- Congenital pseudoparalysis atonica
3588.4- Floppy infant syndrome
3588.9- Other myoneural disorders
Excludes : Infantile spinal muscular atrophy (see code 3350)
Werdnig-Hoffmann syndrome (see code3350)
3589.-- Unspecified myoneural disorders

Muscular dystrophies and other myopathies

Congenital hereditary muscular dystrophy
3590.0- Central core disease
3590.1- Myotubular myopathy
3590.2- Nemaline body disease
3590.3- Benign congenital myopathy
3590.8- Other specified congenital hereditary muscular dystrophy

3590.9- Unspecified congenital hereditary muscular dystrophy

Hereditary progressive muscular dystrophy
3591.0- Duchenne (pseudohypertrophic) dystrophy
3591.1- Becker dystrophy
3591.2- Facioscapulohumeral dystrophy
3591.3- Limb-girdle dystrophy
3591.4- Ocular dystrophy
3591.5- Oculopharyngeal dystrophy
3591.8- Other specified hereditary progressive muscular dystrophy
3591.9- Unspecified hereditary progressive muscular dystrophy

Myotonic disorders / Steinert disease
3592.0- Myotonia congenita
3592.1- Paramyotonia congenita
3592.2- Infantile myotonia
3592.8- Other specified myotonic disorder
3592.9- Unspecified myotonic disorder
Excludes : Myotonic dystrophy / Steinert's neonatal form (see code ~~7592.81~~)

Disorders of eye and adnexa

Hereditary retinal dystrophies
3627.0- Retinitis pigmentosa
3627.1- Batten's disease
3627.2- Leber's optic atrophy
3627.3- Lipofuscinosis NOS
3627.4- Hereditary retinal dystrophy in cerebroretinal lipodoses
3627.5- Tapetoretinal dystrophy
3627.8- Other specified hereditary retinal dystrophies
3627.9- Unspecified hereditary retinal dystrophies

Congenital nystagmus
3631.1- Congenital nystagmus

Deafness

3890.-- Conductive deafness
3891.-- Sensorineural deafness
3892.-- Mixed conductive and sensorineural deafness
3897.-- Deaf mutism, not elsewhere classified
3898.-- Other specified forms of deafness
3899.-- Unspecified deafness

VII. DISEASES OF THE CIRCULATORY SYSTEM

Cardiomyopathy

4250.-- Endomyocardial fibrosis
4251.-- Hypertrophic obstructive cardiomyopathy
4252.-- Obscure cardiomyopathy of Africa
4253.-- Endocardial fibroelastosis
4254.1- Cardiovascular collagenosis
4254.2- Familial cardiomyopathy
4254.9- Other primary cardiomyopathies

Conduction disorders

4260.-- Atrioventricular block, complete
4261.-- Atrioventricular block, other and unspecified
4262.-- Left bundle branch hemiblock
4263.-- Other left bundle branch block
4264.-- Right bundle branch block
4265.-- Bundle branch block, unspecified
4266.-- Sino atrial block
4268.-- Other heart block

Anomalous atrioventricular excitation
4267.1- Wolff-Parkinson-White syndrome
4267.8- Other anomalous atrioventricular excitation

Other conduction disorders
4268.1- Atrioventricular dissociation
4268.2- Interference dissociation
4268.3- Jervell-Lange-Neilsen syndrome
4268.4- Lange-Neilson syndrome
4268.5- Lown-Ganong-Levine syndrome
4268.8- Other specified conduction disorders
4269.-- Unspecified conduction disorders

Cardiac dysrrhythmias

Excludes : postoperative dysrrhythmia (see code 9971)

Paroxysmal supraventricular tachycardia
4270.1- Paroxysmal tachycardia atrial
4270.2- Paroxysmal tachycardia atrioventricular
4270.3- Paroxysmal tachycardia junctional
4270.4- Paroxysmal tachycardia nodal

Paroxysmal ventricular tachycardia
4271.-- Paroxysmal ventricular tachycardia

Paroxysmal tachycardia, unspecified
4272.-- Paroxysmal tachycardia, unspecified

Atrial fibrillation and flutter
4273.1- Atrial fibrillation
4273.2- Atrial flutter

Ventricular fibrillation and flutter
4274.1- Ventricular fibrillation
4274.2- Ventricular flutter

Other cardiac dysrhythmias
4278.-- Other cardiac dysrhythmias

Unspecified cardiac dysrhythmias
4279.-- Unspecified cardiac dysrhythmias

VIII. DISEASES OF THE RESPIRATORY SYSTEM

Exclude all codes in this chapter

IX. DISEASES OF THE DIGESTIVE SYSTEM

Disorders of tooth development and eruption

5200.-- Anodontia
5201.-- Supernumerary teeth
5202.-- Abnormalities of size and form of tooth
5203.-- Mottled teeth
5204.-- Disturbances of tooth formation
5205.1- Amelogenesis imperfecta
5205.8- Hereditary disturbances in tooth structure, not elsewhere classified

Disturbances in tooth eruption
5206.1- Impacted teeth
5206.2- Natal teeth
5206.3- Neonatal teeth

Major anomalies of jaw size

5240.1- Mandibular hyperplasia - mandible OUT
5240.2- Maxillary hyperplasia - maxillary
5240.3- Mandibular hypoplasia - mandible
5240.4- Maxillary hypoplasia - maxillary
5240.5- Micrognathism = retrognathism ?
5240.6- Unilateral micrognathism
5268.-- Unilateral condylar hyperplasia
Excludes : Pierre Robin syndrome (see code 7560)

Defects of abdominal wall

550-.-- Inguinal hernia
5530.-- Femoral hernia
5531.-- Umbilical hernia

X. DISEASES OF THE GENITOURINARY SYSTEM

Exclude all conditions in this chapter - see chapter XIV for :
Congenital nephrotic syndrome (see code 7533.6)
Vesicoureteric reflux (see code 7532.3)
Ureterocele (see code 7534.3)

XI. COMPLICATIONS OF PREGNANCY, CHILDBIRTH AND THE PUERPERIUM

Exclude all codes in this chapter

XII. DISEASES OF THE SKIN AND SUBCUTANEOUS TISSUE

Exclude all codes in this chapter

XIII. DISEASES OF THE MUSCULOSKELETAL SYSTEM AND CONNECTIVE TISSUE

Exclude all codes in this chapter

XIV. CONGENITAL ANOMALIES

CONGENITAL ANOMALIES OF THE NERVOUS SYSTEM

Anencephaly / Exencephaly

Excludes: absence of head (acephaly) in amorphous, acardiac twins (see code 7623.01)

7400.2- Anencephaly, unspecified / acrania
7400.3- Anencephaly, incomplete / hemianencephaly / meroacrania / meroanencephaly, hemicrania. *(Partial absence of brain and cranial vault, the defect not extending through the foramen magnum)*
7400.4- Anencephaly, complete / holoanencephaly / holoacrania *(extending through the foramen magnum.)*

Craniorachischisis / Total or partial craniorachischisis

7401.1- Craniorachischisis, type 1 *(total neural dysraphism) (Anencephaly with exposure of the entire or greater part of the spinal cord)*
7401.2- Craniorachischisis, type 2 *(Anencephaly with dysraphism extending down to the cervical region only.)*
7401.9- Craniorachischisis, unspecified.

Iniencephaly

7402.1- Iniencephaly, open.
7402.2- Iniencephaly, closed.
7402.9- Iniencephaly, unspecified.

Spina bifida / Spina bifida cystica / spina bifida aperta / meningocele / myelomeningocele / meningomyelocele / myelocele / spinal rachischisis / myeloschisis / myelocystocele

Excludes : Spina bifida occulta (see code 7561.01) ← = uncomplicated

NOTE Code hydrocephaly separately, if present. The ICD/BPA code 7410 is not used.

Meningocele (without involvement of neural tissue)

7411.1- Meningocele cervical or cervico-thoracic
7411.2- Meningocele thoracic (dorsal)
7411.3- Meningocele lumbar
7411.4- Meningocele sacral
7411.5- Meningocele thoraco-lumbar
7411.6- Meningocele thoraco-lumbo-sacral
7411.7- Meningocele lumbo-sacral
7411.8- Meningocele other specified
7411.9- Meningocele unspecified

Myelomeningocele * *(spina bifida with involvement of neural tissue, can be open or closed.)*

Includes : meningomyelocele / myelocele / myelocystocele / myeloschisis / spina bifida aperta.

7412.1- Myelomeningocele cervical or cervico-thoracic
7412.2- Myelomeningocele thoracic (dorsal)
7412.3- Myelomeningocele lumbar
7412.4- Myelomeningocele sacral
7412.5- Myelomeningocele thoraco-lumbar
7412.6- Myelomeningocele thoraco-lumbo-sacral
7412.7- Myelomeningocele lumbo-sacral
7412.8- Myelomeningocele other specified
7412.9- Myelomeningocele NOS

Spina bifida * *(spina bifida cystica can be open or closed)*

NOTE This code is to be used when the involvement of neural tissue is not specified.

7419.1- Spina bifida cervical or cervico-thoracic
7419.2- Spina bifida thoracic (dorsal)
7419.3- Spina bifida lumbar
7419.4- Spina bifida sacral
7419.5- Spina bifida thoraco-lumbar
7419.6- Spina bifida thoraco-lumbo-sacral
7419.7- Spina bifida lumbo-sacral
7419.8- Spina bifida other specified
7419.9- Spina bifida unspecified

Cephaloceles

Encephalocele (cystic extension of meninges outside the cranium containing brain tissue.) / encephalomeningocele / encephalomeningomyelocele / meningoencephalocele.

7420.11 Encephalocele occipital
7420.12 Encephalocele frontal
7420.13 Encephalocele parietal
7420.14 Encephalocele orbital
7420.15 Encephalocele nasal
7420.16 Encephalocele nasopharyngeal
7420.17 Encephalocele nasofrontal
7420.18 Encephalocele other specified
7420.19 Encephalocele unspecified site

Cranial meningocele (cystic expansion of meninges outside the cranium without brain tissue.) / cranium bifidum cysticum / cerebral meningocele.

7420.21 Meningocele occipital
7420.22 Meningocele frontal
7420.23 Meningocele parietal
7420.24 Meningocele orbital
7420.25 Meningocele nasal
7420.26 Meningocele nasopharyngeal

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* NOTE Code for 6th digit
1 Open (not covered with skin) or
2 Closed
9 Not known

7420.27 Meningocele nasofrontal
7420.28 Meningocele other specified
7420.29 Meningocele unspecified site

Cephalocele, unspecified
7420.91 Cephalocele occipital
7420.92 Cephalocele frontal
7420.93 Cephalocele parietal
7420.94 Cephalocele orbital
7420.95 Cephalocele nasal
7420.96 Cephalocele nasopharyngeal
7420.97 Cephalocele nasofrontal
7420.98 Cephalocele other specified
7420.99 Cephalocele unspecified site

Microcephaly

7421.01 Microcephaly
7421.11 Microcephalic primordial dwarfism syndrome

Reduction deformities of brain

Anomalies of cerebrum
7422.01 Anomalies of cerebrum

Anomalies of corpus callosum
7422.11 Total absence corpus callosum
7422.12 Partial absence corpus callosum
7422.18 Other specified anomalies of corpus callosum
7422.19 Unspecified anomalies of corpus callosum

Anomalies of hypothalamus
7422.21 Anomalies of hypothalamus

Anomalies of cerebellum
7422.31 Agenesis of cerebellum
7422.32 Hypoplasia of cerebellum
7422.38 Other specified anomalies of cerebellum
7422.39 Unspecified anomalies of cerebellum

Agyria or lissencephaly
7422.41 Miller-Dieker syndrome / Lissencephaly syndrome / Agyria []
7422.5- Microgyria, micropolygyria, polygyria

Pachygyria ?

Arhinencephaly /Holoprosencephaly (absence of the first cranial (olfactory) nerve & tract with or without single undivided cerebral ventricle)*
7422.61 Holoprosencephaly with no or mild visible external facial malformations (e.g. mild hypotelorism)
7422.62 Holoprosencephaly with severe hypotelorism, nose anomalies or cleft lip.
7422.63 Cebocephaly hypotelorism, single nostril
7422.64 Ethmocephaly: agenesis of nose, proboscis attached to interorbital space
7422.65 Cyclopia (fused eyes)
7422.67 Holoprosencephaly sequence
7422.68 Other specified type of holoprosencephaly
7422.69 Unspecified type of holoprosencephaly

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* NOTE Oral clefts or nose anomalies should also be coded additionally, if present

7422.8- Other specified reduction deformities of brain
7422.9- Unspecified reduction deformities of brain

Congenital hydrocephaly *

7423.10 Congenital hydrocephaly
7423.11 Hydrocephaly Dandy-Walker "Syndrome"
7423.12 Hydrocephaly due to atresia of foramina of Magendie & Luschka
7423.13 Hydrocephaly due to atresia of foramina of Monro
7423.14 Hydrocephaly with Arnold Chiari malformation
7423.15 Hydrocephaly due to blockage of aqueduct of Sylvius
7423.16 External hydrocephaly
7423.18 Hydrocephaly other specified
7423.19 Hydrocephaly unspecified

Other specified anomalies of brain

Enlarged brain / Megalencephaly / Macrocephaly
7424.01 Partial / unilateral / asymmetrical enlarged brain
7424.02 Symmetrical enlarged brain
7424.09 Unspecified enlarged brain
7424.1- Porencephaly or porencephalic cyst (not due to neonatal event)
7424.2- Multiple cerebral cysts.
7424.3- Arachnoid cyst
7424.4- Hydranencephaly
7424.5- Arnold Chiari Malformation without hydrocephaly
Excludes : Arnold Chiari Malformation with hydrocephaly (see code 7423.14)
7424.6- Intracranial calcifications.
7424.8- Other specified anomalies of brain

Other anomalies of spinal cord

7425.1- Hypoplasia & dysplasia of spinal cord.
Atelomyelia, myelodysplasia.
7425.2- Diastematomyelia.
7425.3- Other cauda equina anomalies.
7425.4- Hydromyelia, hydrorachischisis
7425.5- Syringomyelia
7425.8- Other specified anomalies of spinal cord

Other specified anomalies of nervous system

7428.01 Jaw-Winking or Marcus-Gunn phenomenon []
7428.1- Familial dysautonomia / Riley-Day syndrome []
7428.21 Möbius (Moebius) syndrome / nuclear agenesis, congenital facial diplegia []
7428.3- Optic atrophy.
7428.4- Septo-optic dysplasia
7428.8- Other specified anomalies of nervous system

* NOTE Code spina bifida separately, if present
Do not code to ICD 7410
Hydrocephaly with neonatal causes, e.g. intraventricular haemorrhage which is not a congenital anomaly is excluded.

Unspecified anomalies of brain, spinal cord and nervous system

7429.01 Unspecified anomalies of brain
7429.1- Unspecified anomalies of spinal cord
7429.9- Unspecified anomalies of nervous system

CONGENITAL ANOMALIES OF EYE

Anophthalmos-Microphthalmos

7430.1- Anophthalmos/Agenesis of eye
7430.2- Cryptophthalmos
7430.22 Cryptophthalmos Fraser Syndrome []
7431.01 Microphthalmos / Aplasia of eye / Dysplasia of eye / Hypoplasia of eye / Rudimentary eye
7431.02 Lenz microphthalmos []

Buphthalmos

7432.01 Buphthalmos
7432.11 Congenital glaucoma
7432.12 Hydrophthalmos
7432.13 Melnick Needles osteodysplasty []
7432.19 Enlarged eye NOS
7432.21 Enlarged cornea / Megalocornea
7432.22 Keratoglobus, congenital

Congenital cataract and lens anomalies

7433.01 Absence of lens / Congenital aphakia
7433.1- Spherical lens/ Spherophakia
7433.2- Cataract
7433.3- Displaced lens
7433.4- Coloboma of lens
7433.8- Other specified lens anomalies
7433.9- Unspecified lens anomalies

Coloboma and other anomalies of anterior segments

7434.01 Corneal opacity
7434.1- Other corneal anomalies
Excludes : megalocornea (see code 7432.21)
7434.21 Absence of iris / Aniridia
7434.22 Anridia Wilms Tumour association []
7434.3- Coloboma of iris
7434.41 Polycoria
7434.42 Ectopic pupil
7434.48 Other anomalies of iris
7434.5- Blue sclera
7434.6- Rieger's syndrome / Iridogoniodysgenesis with somatic anomalies []
7434.8- Other specified anomalies of sclera or anterior part of eye
7434.9- Unspecified anomalies of anterior part of eye

Congenital anomalies of posterior segment

7435.01 Specified anomalies of vitreous humour
~~7435.08 Specified anomalies of retina~~ 743518 Other Specified anomalies of retina
7435.1- Congenital retinal aneurysm
7435.2- Coloboma of optic disc
7435.3- Specified anomalies of choroid
7435.4- Specified anomalies of optic disc
7435.8- Other specified anomalies of posterior segment
7435.9- Unspecified anomalies of posterior segment

Congenital anomalies of eyelids, lacrimal system and orbit

7436.01 Blepharoptosis / Congenital ptosis
7436.02 Blepharophimosis-ptosis syndrome []
7436.1- Ectropion
7436.2- Entropion
7436.30 Epicanthic folds/narrow palpebral fissures
7436.31 Fused eyelids : Symblepharon
7436.32 Absence of cilia / absence of eyelashes
7436.38 Other anomalies of eyelids
7436.41 Absence or agenesis of lacrimal apparatus
7436.42 Absence of punctum lacrimale
7436.5- Stenosis or stricture of lacrimal duct
7436.6- Other anomalies of lacrimal apparatus
7436.7- Anomalies of orbit
7436.81 Anti-mongoloid slant
7436.82 Mongoloïd slant

Other specified anomalies of eye

7438.-- Other specified anomalies of eye
Excludes : Ocular albinism (see code 2702)
congenital nystagmus (see code 3795)
retinitis pigmentosa (see code 3627)

Unspecified anomalies of eye

7439.-- Congenital anomaly NOS of eye (any part) / deformity NOS of eye (any part)

CONGENITAL ANOMALIES OF EAR, FACE AND NECK

Anomalies of ear causing impairment of hearing*

7440.0- Absence or stricture of auditory canal
7440.1- Absence of auricle / Absence of ear NOS

Anomaly of middle ear
7440.21 Fusion of ossicles
7440.22 Absence of ossicles
7440.23 Ankylosis of ossicles
7440.28 Other anomalies of middle ear

Anomaly of inner ear
7440.30 Congenital anomaly of membranous labyrinth
7440.31 Congenital anomaly of the organ of Corti
7440.38 Other congenital anomaly of the inner ear
7440.39 Unspecified congenital anomaly of the inner ear

7440.9- Unspecified anomaly of ear with impairment of hearing

*NOTE Congenital deafness due to nervous problems or NOS (see 389- -)

Accessory auricle

7441.01 Accessory auricle / Polyotia
7441.1- Preauricular appendage, tag or lobule
7441.2- Other appendage, tag or lobule

Other specified anomalies of ear

7442.01 Macrotia
7442.1- Microtia
7442.2- Bat ear
7442.31 Simple ears
7442.33 Pointed ear
7442.34 Vulcan ears
7442.38 Other misshapen ears
7442.41 Misplaced ears
7442.42 Low set ears
7442.5- Absence or anomaly of Eustachian tube
7442.6 Darwin's tubercle
7442.8- Other specified anomalies of ear

Unspecified anomalies of ear

7443.9- Congenital anomaly / deformity of ear (any part) NOS

First branchial arch anomalies, cleft, cyst or fistula, preauricular sinus

7444.01 Branchial cleft associated with first branchial arch
7444.02 Branchial sinus associated with first branchial arch
7444.03 Internal branchial fistula associated with first branchial arch
7444.04 Branchial cyst associated with first branchial arch
7444.1- Preauricular sinus or cyst
7444.21 Melnick-Fraser / Branchio-oto-renal / BOR syndrome []

7444.8- Other branchial cleft anomalies (first arch)
Excludes : anomalies associated with second, third and fourth branchial arches (see code 7502.8)

Webbing of neck

7445.-- Pterygium colli / Webbing of neck

Other specified anomalies of face and neck

7448.01 Macrostomia
7448.1- Microstomia
7448.2- Macrocheilia
7448.3- Microcheilia
7448.4- Short/restricted neck
7448.8- Other specified anomalies of face and neck

Unspecified anomalies of face and neck

7449.09 Congenital anomaly of neck NOS
7449.19 Congenital anomaly of face NOS
Excludes : Potter's facies (see 75401)

CONGENITAL ANOMALIES OF HEART*

Malformations of cardiac chambers and connections

Common arterial truncus
7450.01 Persistent truncus arteriosus type I
7450.02 Persistent truncus arteriosus type II
7450.03 Persistent truncus arteriosus type III
7450.09 Persistent truncus arteriosus unknown type
NOTE See 7473.13 for truncus arteriosus type IV
7450.1- Aortic septal defect / Aorto-pulmonary window

Discordant ventriculoarterial connection
7451.01 Dextrotransposition of aorta
7451.02 Dextrotransposition in situs solitus / complete transposition of great vessels
7451.03 Dextro transposition in situs inversus
7451.11 Double outlet right ventricle
7451.12 Taussig-Bing syndrome

Discordant atrioventricular connection, atrial isomerism

7451.21 Corrected transposition of great vessels / ventricular inversion with T.G.A.
7451.22 Levotransposition in situs inversus
7451.23 Levotransposition in situs solitus
7451.24 Ventricular inversion
7451.3- Miror image atrial arrangement (with asplenia or polysplenia)
7451.8- Other specified malformations of cardiac chambers and connections
7451.9- Unspecified malformations of cardiac chambers and connections
d transposition of GV, unspecified
Tetralogy of Fallot
7452.0- Fallot's tetralogy
7452.1- Fallot's pentalogy

Single ventricle
7453.31 Double inlet left ventricle with TGA
7453.32 Double inlet left ventricle with normally related arteries
7453.33 Double inlet right ventricle with TGA
7453.34 Double inlet right ventricle with normally related arteries
7453.35 Single ventricle with TGA
7453.36 Single ventricle with normally related arteries
7453.37 Cor triloculare biatriatum
7453.39 Single ventricle unspecified

Malformations of cardiac septa

Ventricular septal defect
7454.0- Roger's disease
7454.1- Ventricular septal defect, Eisenmenger's syndrome
7454.2- Gerbode defect
7454.3- Ventricular septal defect supracristal
7454.4- Ventricular septal defect subpulmonary
7454.5- Ventricular septal defect muscular
7454.6- Ventricular septal defect perimembranous
7454.7- Ventricular septal defect multiple

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* For rhythm disturbances see 4260 to 4279

7454.8- Other type of ventricular septal defect
7454.9- Unspecified type of ventricular septal defect

Atrial septal defect (ostium secundum type) ASD II
7455.0- Foramen ovale patent / non-closure of foramen ovale
7455.1- Ostium secundum defect
7455.2- Lutembacher's syndrome
7455.8- Other specified atrial septal defect
7455.9- Unspecified atrial septal defect

Endocardial cushion defect complete AVSD 7456-
7456.0- Ostium primum defect
7456.1- Single common atrium defect
7456.2- Common atrioventricular canal type ventricular septal defect
7456.3- Common atrioventricular canal
7456.8- Other specified atrioventricular septal defects
7456.9- Unspecified atrioventricular septal defects

7457.-- Cor biloculare / cor triloculare biventriculare
7458.-- Other specified type of malformations of the septa
7459.-- Unspecified defect of septal closure

Other congenital anomalies of heart cong. malformations of pulmonary and tricuspid valves

Anomalies of pulmonary and tricuspid valves
7460.01 Pulmonary valve atresia with intact ventricular septum
7460.02 Pulmonary valve atresia with ventricular septal defect and normal pulmonary trunk and branches
7460.11 Pulmonary stenosis, valvar
Exclude : pulmonary infundibular stenosis (see code 7468.31)
Truncus arteriosus type IV (see code 7473.13)
7460.19 Pulmonary stenosis, non specified
7460.2- Pulmonary valve insuffisiency (congenital) / regurgitation
7460.8- Other specified anomalies of pulmonary valve
7460.9- Unspecified anomalies of pulmonary valve

7461.11 Tricuspid atresia
7461.12 Tricuspid stenosis (congenital)
7461.13 Tricuspid insufficiency (congenital)
7461.14 Straddling of tricuspid valve
7461.16 Dysplasia of tricuspid valve

7462.-- Ebstein's anomaly
7462.8- Other specified anomalies of tricuspid valve
7462.9- Unspecified anomalies of tricuspid valve

Cong. *Anomalies of aortic and mitral valves*
7463.1- Aortic valve atresia (exclude if part of hypoplastic left heart syndrome)
7463.2- Aortic stenosis valvar
Excludes : supraaortic stenosis (see code 7472.2)

7464.1- Aortic valve insufficiency (congenital)
7464.2- Bicuspid aortic valve
7464.8- Other specified anomalies of aortic valve
7464.9- Unspecified anomalies of aortic valve

7465.1- Mitral atresia (exclude if part of hypoplastic left heart syndrome)
7465.2- Mitral stenosis (congenital)

7466.1- Mitral insufficiency (congenital)
7466.8- Other specified anomalies of mitral valve
7466.9- Unspecified anomalies of mitral valve

7467.1- Hypoplastic left heart syndrome with aortic valvar atresia
7467.2- Hypoplastic left heart syndrome with aortic valvar hypoplasia
7467.3- Hypoplastic left heart syndrome with hypoplastic aorta
7467.4- Hypoplastic left heart syndrome with mitral valve atresia
7467.5- Hypoplastic left heart syndrome with mitral valve hypoplasia
7467.9- Hypoplastic left heart syndrome NOS

Other malformations and anomalies of the heart
(exclude endocardial fibroelastosis, malformations of coronary arteries)

7468.0- Dextrocardia
Exclude : dextrocardia with situs inversus, miror image atrial arrangement
7468.1- Levocardia
7468.2- Cor triatriatum
7468.31 Pulmonary infundibular stenosis
7468.32 Subaortic stenosis dynamic (hypertrophic)
7468.33 Subaortic stenosis fixed (fibrous shelf or tunnel)
7468.4- Trilogy of Fallot
7468.5- Malformations of pericardium
7468.6- Malformations of myocardium
7468.81 Ectopia cordis
7468.88 Other specified anomalies of heart
7468.92 Malposition of the heart
7468.93 Congenital diverticulum of left ventricle
7468.94 Uhl's (parchment right ventricle)
Exclude : dextrocardia with situs inversus)

Unspecified anomalies of heart
7469.0- Congenital anomalies of the heart
7469.1- Congenital anomalies of heart (valves) / Anomalous bands of heart
7469.2- Congenital heart disease NOS- acyanotic
7469.3- Cyanotic congenital heart disease NOS / Blue baby
7469.9- Anomaly of heart unspecified

Anomalies of great arteries

7470 ~~7513~~.-- Patent ductus arteriosus
7470 1, aneurysm of ductus arteriosus.
Coarctation of aorta
7471.0- Preductal
7471.1- Postductal
7471.9- Unspecified type of coarctation of aorta

Other anomalies of aorta
7472.01 Interruption of aorta distal to left subclavian artery
7472.02 Interruption of aorta distal to common carotid
7472.03 Interruption of aorta distal to innominate artery
7472.09 Interruption of aorta NOS
7472.1- Hypoplasia of aorta (tubular hypoplasia)
7472.2- Supraaortic stenosis / Supravalvular aortic stenosis
7472.3- Persistent right aortic arch
7472.4- Aneurysm (congenital) of sinus of Valsalva

? et titre page précédent ?

BPA = .0 .1 .2 .3 .4

• 0 = heart valves
• 1 = and bands of H.
• 2 acyanotic ...

Pourquoi changer .0 et .1 ?
⇒ remettre cf BPA

7472.51 Double aortic arch
7472.52 Left aortic arch with anomalous origin of right subclavian artery
7472.53 Right aortic arch with left ligamentum arteriosum
7472.59 Vascular ring of aorta NOS
7472.6- Overriding aorta
7472.7- Congenital aneurysm of aorta / congenital dilatation of aorta
7472.8- Other specified anomalies of aorta
7472.9- Unspecified anomalies of aorta

artère pulmonaire ?

Anomalies of pulmonary artery
7473.11 Atresia of pulmonary artery
7473.12 Absence of pulmonary trunk
7473.13 Persistent truncus arteriosus type IV
7473.14 Absence of right pulmonary artery
7473.15 Absence of left pulmonary artery
7473.21 Stenosis of left artery isolated
7473.22 Stenosis of left artery multiple
7473.23 Stenosis of right artery isolated
7473.24 Stenosis of right artery multiple
7473.25 Stenosis of pulmonary artery unspecified
7473.33 Aneurysm of pulmonary artery
7473.34 Pulmonary arteriovenous aneurysm
7473.35 Anomalous origin of left pulmonary artery
7473.36 Anomalous origin of right pulmonary artery
7473.38 Other specified anomalies of pulmonary artery
7473.39 Unspecified anomalies of pulmonary artery

erreur ces 3 03 (digit 5e) ne devraient pas être là !

Anomalies of great veins

7474.0- Congenital stenosis of vena cava (inferior and/or superior)
7474.1- Persistent left superior vena cava
7474.21 Total anomalous pulmonary vein connection infradiaphragmatic
7474.22 Total anomalous pulmonary vein connection supradiaphragmatic
7474.3- Partial anomalous pulmonary venous connection
7474.38 Unspecified anomalous pulmonary venous connection
7474.41 Anomalous portal venous connection
7474.42 Portal vein - hepatic artery fistula

Other specified malformations of great veins
7474.81 Absence of vena cava superior/inferior
7474.82 Azygos continuation of inferior vena cava
7474.83 Persistent left posterior cardinal vein
7474.84 Scimitar syndrome
7474.99 Unspecified anomalies of great veins

Malformations of peripheral vascular system

Exclude : anomalies of cerebral vessels, pulmonary artery, congenital retinal aneurysm, hemangioma, lymphangioma)
7475.-- Absence or hypoplasia of umbilical artery / Single umbilical artery
7476.0- Congenital renal artery stenosis
7476.1- Other specified anomalies of renal artery
7476.8- Unspecified anomalies of renal artery

?

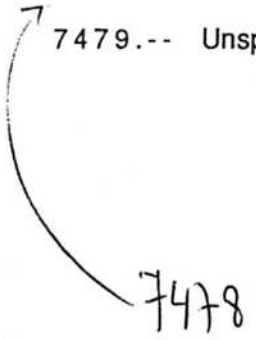
7478 cerebral aneurysm où le mettre ?

Anomalies d'autres vaisseaux que les vaisseaux rénaux ?

Other specified anomalies of circulatory system

Exclude : congenital aneurysm : coronary , peripheral , pulmonary, retinal; ruptured : cerebral arteriovenous aneurysm, congenital cerebral aneurysm)

7479.-- Unspecified anomalies of circulatory system



CONGENITAL ANOMALIES OF RESPIRATORY SYSTEM

Anomalies of nose

7480.01 Bilateral posterior choanal atresia
7480.02 Bilateral anterior choanal atresia
7480.09 Bilateral choanal atresia, site unspecified
7480.11 Unilateral posterior choanal atresia
7480.12 Unilateral anterior choanal atresia
7480.19 Unilateral choanal atresia, site unspecified
7480.21 CHARGE association []
7480.29 Choanal atresia, site unspecified, multiplicity unspecified
7480.51 Bilateral posterior choanal stenosis
7480.52 Bilateral anterior choanal stenosis
7480.59 Bilateral choanal stenosis, site unspecified
7480.61 Unilateral posterior choanal stenosis
7480.62 Unilateral anterior choanal stenosis
7480.69 Unilateral choanal stenosis, site unspecified
7480.79 Choanal stenosis, site unspecified, multiplicity unspecified
7480.99 ~~Choanal atresia/stenosis, unspecified NOS~~ unspecified and of choanae.

Other anomalies of nose

Excludes : arhinencephaly (see code 7422.6-)
7481.01 Complete nasal aplasia with nasal cavity
7481.02 Complete nasal aplasia without nasal cavity
7481.03 Unilateral nasal aplasia with nasal cavity
7481.04 Unilateral nasal aplasia without nasal cavity
7481.1- Accessory nose
7481.21 Fissured, notched or cleft nose median
7481.22 Fissured, notched of cleft nose lateral
7481.3- Nasal sinus wall anomalies
7481.4- Perforated nasal septum
7481.81 Johnson Blizzard syndrome []
7481.82 Marshall syndrome []
7481.83 Langer Geidion syndrome / Trichorhinophalangeal type II syndrome []
7481.88 Other specified anomalies of nose
7481.9- Unspecified anomalies of nose

Anomalies of larynx

7482.0- Web of larynx : glottic
7482.1- Web of larynx : subglottic
7482.3- Web of larynx : NOS

Other anomalies of larynx, trachea and bronchus

7483.01 Congenital atresia of larynx
7483.02 Congenital cyst of larynx
7483.03 Congenital papilloma of larynx
7483.04 Congenital laryngomalacia
7483.05 Congenital subglottic hemangioma
7483.06 Congenital subglottic lymphangioma
7483.07 Laryngocele
7483.1- Congenital subglottic stenosis
7483.2- Tracheomalacia
7483.31 Absence of trachea
7483.32 Stenosis of trachea

7483.33 Diverticulum of trachea
Excludes : Tracheooesophageal fistula (see code 7503.12)
7483.34 Duplication of trachea
Excludes : Laryngotracheooesophageal cleft (see code 7503.22)
7483.41 Bronchial atresia
7483.42 Bronchial stenosis
7483.43 Supernumerary bronchus
7483.51 Isolated bronchomalacia
7483.52 Other anomalies of bronchus
Excludes : bronchobiliary fistula (see code 7516.71)
7483.61 Congenital laryngeal stridor NOS
7483.62 Absence of vocal cords
7483.63 Duplication of vocal cords
7483.64 Vocal cord palsy
7483.65 Sulcus glottitis
7483.66 Malformations of laryngeal cartilages
7483.8- Other specified anomalies of larynx, trachea or bronchus
7483.9- Unspecified anomalies of larynx, trachea or bronchus

Congenital cystic lung

7484.0- Single cyst of lung / Lung cyst
7484.1- Multiple lung cysts / Polycystic lung / Cystic lung
7484.21 Honeycomb lung
7484.22 Congenital lobar emphysema
7484.23 Congenital pneumatocele
7484.8- Other specified types of cystic lung
7484.9- Unspecified type of cystic lung

Agenesis, hypoplasia and dysplasia of lung

7485.01 Agenesis of lung *(complete absence of lungs, bronchi and vascular structures)*
7485.02 Aplasia of lung *(absence of pulmonary and vascular structure)*
7485.1- Hypoplasia of lung (*abnormal development)*
7485.2- Sequestration of lung tissue
7485.8- Other specified dysplasia of lung
7485.9- Unspecified dysplasia of lung

Other anomalies of lung

7486.01 Hamartoma of lung
7486.02 Other ectopic tissues in lung
7486.1- Bronchiectasis, congenital
7486.21 Supernumerary lung
7486.22 Accessory lobe of lung
7486.23 Supernumerary tissues (lobation) of the lung
7486.24 Azygos lobe
7486.8- Other specified anomalies of lung
7486.9- Unspecified anomalies of lung
Excludes : Pulmonary arterio-venous fistula (see code 7473.34)

Other specified anomalies of respiratory system

7488.0- Anomaly of pleura
7488.11 Bronchogenic cyst of the mediastinum
7488.12 Other congenital cyst of mediastinum
7488.8- Other specified anomalies of respiratory system

Unspecified anomalies of respiratory system

7489.19 Absence of respiratory organ NOS
7489.29 Anomaly of respiratory system NOS

CLEFT PALATE AND CLEFT LIP

Cleft of the secondary palate / cleft lying posterior to the incisive foramen = palate only

7490.01 Cleft hard palate unilateral, right
7490.02 Cleft hard palate unilateral, left
7490.03 Cleft hard palate unilateral, unspecified side
7490.1- Cleft hard palate bilateral
7490.2- Cleft hard palate central
7490.3- Cleft hard palate NOS
7490.41 Cleft soft palate unilateral / cleft velum palate unilateral, right
7490.42 Cleft soft palate unilateral / cleft velum palate unilateral, left
7490.43 Cleft soft palate unilateral unspecified side / cleft velum palate unilateral, unspecified side
7490.5- Cleft soft palate bilateral
7490.6- Cleft soft palate central
7490.71 Submucous cleft
7490.72 Cleft soft palate NOS
7490.8- Cleft uvula
7490.91 Cleft palate total *(soft palate and hard palate)* unilateral, right
7490.92 Cleft palate total *(soft palate and hard palate)* unilateral, left
7490.93 Cleft palate total *(soft palate and hard palate)* unilateral, unspecified side
7490.94 Cleft palate total bilateral
7490.95 Cleft palate total median
7490.99 Cleft palate NOS

Cleft of primary palate / Cleft lip and premaxilla / Cleft including and lying anterior to the incisive foramen (include dental ridge/gum) = lip only

7491.01 Cleft of primary palate unilateral subtotal *(lip only)*, right
7491.02 Cleft of primary palate unilateral subtotal *(lip only)*, left
7491.03 Cleft of primary palate unilateral subtotal *(lip only)*, unspecified side
7491.04 Cleft of primary palate unilateral total, right
7491.05 Cleft of primary palate unilateral total, left
7491.06 Cleft of primary palate unilateral total, unspecified side
7491.11 Cleft of primary palate bilateral subtotal *(lip only)*
7491.12 Cleft of primary palate bilateral total
7491.21 Cleft of primary palate central subtotal *(lip only)*
7491.22 Cleft of primary palate central total
7491.98 Other specified cleft of primary palate
7491.99 Cleft of primary palate NOS

Cleft of secondary and primary palate = both lip & palate

7492.01 Cleft of primary and secondary palate unilateral total, right
7492.02 Cleft of primary and secondary palate unilateral total, left
7492.0' Cleft of primary and secondary palate unilateral total, unspecified side
7492.04 Cleft of primary and secondary palate unilateral subtotal, right
7492.05 Cleft of primary and secondary palate unilateral subtotal, left
7492.06 Cleft of primary and secondary palate unilateral subtotal, unspecified side
7492.11 Cleft of primary and secondary palate bilateral total
7492.12 Cleft of primary and secondary palate bilateral subtotal
7492.21 Cleft of primary and secondary palate central total
7492.22 Cleft of primary and secondary palate central subtotal
7492.91 Van der Woude syndrome []
7492.99 Cleft of primary and secondary palate NOS

Unusual cleft

7493.-- Unusual cleft such as mandibular clefts, cleft of the lower lip, naso-ocular cleft, oro-ocular clefts

749

OTHER CONGENITAL ANOMALIES OF UPPER ALIMENTARY TRACT

Tongue tie

7500.-- Ankyloglossia

Other anomalies of tongue

7501.0- Aglossia / Absence of tongue
7501.1- Hypoplasia of tongue / Microglossia
7501.2- Macroglossia
7501.3- Dislocation or displacement of tongue
7501.4- Cleft tongue
7501.5- Adhesions tongue, congenital *(to gum or roof of mouth)*
7501.8- Other specified anomalies of tongue
7501.9- Unspecified anomalies of tongue

Other specified anomalies of mouth and pharynx

7502.01 Pharyngeal pouch
7502.1- Other pharyngeal anomalies
7502.2- Ranula
7502.31 Parotid aplasia or hypoplasia
7502.32 Diverticulum or ectasia of parotid duct system
7502.33 Anomalies of submaxillary glands
7502.34 Anomalies of sublingual glands
7502.35 Other specified anomalies of salivary glands or ducts
7502.4- High arched palate
7502.5- Other anomalies of palate
7502.6- Lip fistulae or pits
7502.7- Other lip anomalies
Excludes : Cleft lip (see section 749)
7502.81 Branchial cyst anomalies associated with 2nd branchial arch
(Anomalies associated with first branchial arch (see code 7444.0))
7502.82 Branchial cyst anomalies associated with 3rd branchial arch
7502.83 Branchial cyst anomalies associated with 4th branchial arch
7502.84 Branchial fistula anomalies associated with 2nd branchial arch
7502.85 Branchial fistula anomalies associated with 3rd branchial arch
7502.86 Branchial fistula anomalies associated with 4th branchial arch
7502.91 Other specified anomalies of mouth and pharynx
7502.92 Fordyce disease (mouth)
7502.93 Congenital cyst of the mouth
7502.94 Laryngo-tracheo-oesophageal diastema / fissure
7502.99 Other unspecified anomaly of mouth and pharynx

Tracheo-oesophageal fistula, oesophageal atresia and stenosis

7503.0- Oesophageal atresia without fistula (Gross type A)
7503.11 Atresia of the upper part of oesophagus with tracheo-oesophageal fistula (Gross type C)
7503.12 Atresia of the lower oesophagus with tracheo-oesophageal fistula (Gross type B)
7503.13 Atresia of oesophagus with both segments communicating with the trachea (Gross type D)
7503.21 Tracheo-oesophageal fistula without oesophageal atresia (Gross type E)
7503.22 Laryngo-tracheo-oesophageal cleft
7503.31 Broncho-oesophageal fistula with oesophageal atresia
7503.32 Broncho-oesophageal fistula without oesophageal atresia

7503.33 Broncho-oesophageal fistula with tracheal atresia (uni or bilateral)
7503.34 Broncho-oesophageal fistula without tracheal atresia (uni or bilateral)

Stenosis or stricture of oesophagus
7503.41 Oesophageal diaphragm / oesophageal web
7503.42 Oesophageal fibromuscular stenosis
7503.43 Oesophageal stenosis due to inclusion of ectopic or hamartomatous tissue

7503.8- Other anomaly of oesophagus (stenosis)
7503.9- Unspecified type of oesophageal atresia

Other specified anomalies of oesophagus

7504.0- Congenital dilatation of oesophagus / Giant oesophagus
7504.1- Displacement of oesophagus
7504.2- Diverticulum of oesophagus / Oesophageal pouch
7504.3- Duplication of oesophagus
7504.8- Other specified anomaly of oesophagus
7504.9- Unspecified anomaly of oesophagus

Congenital hypertrophic pyloric stenosis

7505.1- Congenital hypertrophic pyloric stenosis
7505.8- Other congenital pyloric obstruction

Other anomalies of the pylorus

7506.01 Congenital hiatus hernia / Displacement of cardia through oesophageal hiatus / Partial thoracic stomach
Excludes : congenital diaphragmatic hernia (see code 7566.1)

Other specified anomalies of the stomach

7507.01 Microgastria
7507.02 Congenital atresia of the stomach
7507.03 Congenital gastric obstruction *(mucosal diaphragm)*
7507.1- Megalogastria
7507.2- Congenital achalasia of the cardia
7507.31 Isolated situs inversus of the stomach
7507.32 Other displacement or transposition of the stomach
7507.41 Diverticulum of stomach
7507.42 Enterogenous cyst of the stomach
7507.5- Duplication of stomach
7507.8- Other specified anomaly of stomach
7507.9- Unspecified anomaly of stomach

Other specified anomalies of upper alimentary tract

7508.-- Other specified anomalies of upper alimentary tract

Unspecified anomalies of unspecified part of upper alimentary tract

7509.-- Unspecified anomalies of unspecified part of upper alimentary tract

OTHER CONGENITAL ANOMALIES OF DIGESTIVE SYSTEM

Meckel's diverticulum

7510.0- Persistent omphalomesenteric duct / Persistent vitelline duct
7510.1- Meckel's diverticulum
7510.2- Isolated Meckel bridle
7510.3- Omphalomesenteric cyst (independent of digestive tract)

Atresia and stenosis of small intestine

7511.01 Absence of duodenum
7511.02 Atresia of duodenum
7511.03 Stenosis of duodenum
7511.04 Duodenal diaphragm
7511.11 Absence of jejunum
7511.12 Atresia of jejunum, isolated
7511.13 Atresia of jejunum, multiple
7511.14 Stenosis of jejunum
7511.21 Absence of ileum
7511.22 Atresia of ileum
7511.23 Stenosis of ileum
7511.91 Absence of small intestine NOS
7511.92 Atresia of small intestine NOS
7511.93 Stenosis of small intestine NOS

Atresia and stenosis of large intestine, rectum and anal canal*

7512.01 Atresia of the colon / Colic agenesis (= absence of various length of the colon)
7512.02 Stenosis of the colon
7512.03 Stenosis, atresia or absence of appendix
7512.12 Rectal atresia (with normal anal canal and anus) / Anorectal agenesis (supra levator) with fistula (rectovesical or rectourethral or rectovaginal)
7512.21 High anorectal agenesis (supra levator) without fistula
7512.31 Anal agenesis (intermediate anomaly) with fistula (rectovulvar or
7512.32 rectovaginal low or rectovestibular)
7512.41 Anal agenesis (intermediate anomaly) without fistula
7512.42 Complete covered anus, at normal anal site (low anomaly) without fistula
7512.43 Complete covered anus at normal anal site with perineal fistula (males only)
Excludes: low anomaly with fistula in female (see code 7515.43 and other)
7512.44 Anal stenosis
7512.45 Imperforate anal membrane

Hirschsprung's disease and other congenital functional disorders of colon

7513.0- Total intestinal aganglionosis
7513.11 Total colonic aganglionosis with aganglionosis in the small intestine (variable length)
7513.12 Total colonic aganglionosis with normal small intestine
7513.2- Aganglionosis of a colonic segment (variable length)

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* For other low anomalies see code numbers 7515.3, 7515.4

7513.3- Hirschsprung's disease NOS
7513.4- Congenital megacolon (without aganglionosis) / Congenital macrocolon (not aganglionic)

Anomalies of intestinal fixation and mesenteric anomalies

7514.01 Abnormal position of intestine
7514.02 Reversed rotation of intestine
7514.03 Volvulus due to abnormal rotation or fixation
7514.11 Common mesentery
7514.12 Mesenteric cyst
7514.18 Other anomalies of mesentery
7514.2- Congenital adhesions or bands of omentum and peritoneum
7514.8- Other anomalies of intestinal fixation
7514.9- Unspecified anomalies of intestinal fixation

Other anomalies of intestine

7515.01 Duplication of anus
7515.02 Duplication of appendix
7515.03 Duplication of caecum
7515.03 Duplication of colon
7515.04 Duplication of small intestine
7515.05 Congenital intestinal diverticulum
7515.1- Transposition of appendix, colon and intestine
7515.2- Microcolon
7515.31 Anterior perineal anus (female)
7515.33 Ectopic anus
7515.43 Anovestibular fistula (female)
7515.5- Persistent cloaca
7515.81 Short bowel syndrome
7515.88 Other specified anomaly of intestine
7515.99 Unspecified anomaly of intestine

Anomalies of gallbladder, bile ducts and liver

7516.01 Absence or agenesis of liver (total)
7516.02 Absence or agenesis of liver (part)
7516.03 Abnormal lobation of liver
7516.04 Atresia of the cystic duct
7516.11 Intrahepatic congenital cyst
7516.12 Polycystic disease of the liver
7516.13 Congenital hepatic fibrosis
7516.2- Other anomalies of liver
Excludes : metabolic disease of liver
7516.3- Agenesis or hypoplasia of gallbladder
7516.41 Duplication of gallbladder
7516.42 Abnormal position of gallbladder
7516.48 Other anomalies of gallbladder
7516.51 Congenital atresia of the extrahepatic bile ducts
7516.52 Congenital hypoplasia of the intrahepatic bile ducts
7516.53 Alagille syndrome
7516.54 Congenital atresia of bile ducts NOS
7516.6- Choledochal cysts
7516.71 Tracheobiliary fistula
7516.72 Other anomalies of hepatic or bile ducts
7516.8- Other specified anomalies of biliary tract
7516.9- Unspecified anomalies of biliary tract or bile ducts

Anomalies of pancreas

7517.01 Aplasia of the whole pancreas
7517.02 Aplasia of the central pancreas
7517.03 Aplasia of the dorsal pancreas
7517.04 Hypoplasia of the exocrine pancreas / Lipomatosis of the pancreas
7517.05 Pancreas divided
7517.06 Stenosis and other anomalies of excretory duct of the pancreas
7517.1- Accessory pancreas
7517.2- Annular pancreas
7517.3- Ectopic pancreas
7517.41 Isolated pancreatic cyst
7517.42 Congenital cystosis of pancreas
7517.8- Other specified anomalies of the pancreas
7517.9- Unspecified anomalies of pancreas

Other specified anomalies of digestive system

7518.0- Absence (complete) (partial) of alimentary tract NOS
7518.1- Duplication of alimentary tract
7518.2- Ectopic digestive organs NOS
7518.8- Other specified anomaly of digestive organs not elsewhere classified (= NEC)

Unspecified anomalies of digestive system

7519.-- Congenital anomaly NOS of digestive system

CONGENITAL ANOMALIES OF THE GENITAL ORGANS

Excludes : testicular feminization syndrome (see code 2578)
syndromes associated with anomalies in the number and form of chromosomes (see code 758--)

Anomalies of ovaries

7520.01 Bilateral absence of the ovaries / agenesis of ovaries
7520.02 Unilateral absence of the ovary
7520.1- Streak ovary
7520.2- Accessory ovary
7520.3- Isolated cyst of ovary
7520.4- Multicystic ovary
7520.81 Dysplasia of ovary
7520.82 Pedicle torsion of ovary
7520.9- Unspecified anomalies of ovaries

Anomalies of fallopian tubes and broad ligament

7521.01 Absence of fallopian tube / oviduct, one or both sides
7521.02 Atresia of fallopian tube/oviduct, one or both sides
7521.03 Absence of broad ligament, one or both sides
7521.04 Atresia of broad ligament, one or both sides
7521.05 Absence of round ligament, one or both sides
7521.11 Epoophoron cyst
7521.12 Cyst of Gartner's duct
7521.13 Anomaly of Gartner's duct
7521.21 Fimbrial cyst / hydatid of Morgagni
7521.22 Para-ovarian cyst
7521.23 Cyst of mesenteric remnant
7521.81 Accessory broad ligament
7521.82 Accessory fallopian tube
7521.88 Other specified anomalies of the broad ligament
7521.99 Unspecified anomalies of the fallopian tubes, oviduct or broad ligament

Anomalies of uterus

7522.11 Double uterus / didelphic uterus with double cervix and vagina
7522.12 Accessory uterus with extra cervix and vagina
7523.01 Absence / agenesis of uterus
7523.11 Displaced uterus
7523.13 Prolapsed uterus
7523.21 Fistula of uterus with intestine
7523.22 Fistula of uterus with rectum
7523.23 Fistula of uterus with digestive tract / utero intestinal fistula
7523.24 Fistula of uterus with urinary bladder / utero vesical fistula
7523.81 Single horned / unicornate uterus
7523.82 Partially divided / bicornuate / septate uterus (not vagina)
7523.83 Aplasia of round ligament
7523.88 Other specified anomalies of uterus
7523.99 Unspecified anomalies of uterus

Anomalies of cervix, vagina and external genitalia

7524.01 Absence / agenesis of cervix
7524.02 Atresia / stricture of cervix
7524.03 Imperforate cervix

7524.04 Rudimentary cervix
7524.11 Absence of vagina / complete atresia of vagina
7524.12 Partial atresia of vagina
7524.13 Murcs association
7524.21 Congenital rectovaginal fistula
7524.31 Imperforate hymen
7524.32 Hydrometrocolpos with post-axial polysyndactyly syndrome [] / Kaufman McKusick syndrome
7524.41 Absence of vulva
7524.42 Fusion of vulva
7524.48 Other anomalies of vulva
7524.51 Absence of clitoris
7524.58 Other anomaly of clitoris
7524.61 Embryonal cyst of vagina / persistent canal of Nuck
7524.71 Perineal cyst
7524.88 Other specified anomalies of cervix, vagina or external female genitalia
7524.99 Unspecified anomalies of cervix, vagina or external female genitalia

Anomalies of testicles

7525.01 Undescended testicle/ cryptorchidism unilateral
7525.11 Undescended testicles / cryptorchidism bilateral
7525.21 Undescended testicles / cryptorchidism NOS
7525.31 Ectopic testicle unilateral
7525.32 Ectopic testicle bilateral

7525.39 Ectopic testicle unspecified

Anomalies of penis

7526.01 Hypospadias opening on lower side of glans penis / glandular hypospadias . 1er d°.
7526.02 Hypospadias opening on body of penis / penile
7526.03 Hypospadias opening at the base of scrotum / penoscrotal
7526.04 Hypospadias opening behind the scrotum / perineal
7526.09 Hypospadias, position of opening not stated / anaspadia / paraspadia
7526.11 Epispadias / urethra opening on upper side of penis
7526.21 Congenital chordee / lateral curved penis

Indeterminate sex and pseudo-hermaphroditism*

7527.01 True hermaphroditism
7527.02 Ovotestis
7527.03 Pseudo-hermaphroditism in male
7527.11 Pseudo-hermaphroditism in female
7527.12 Testicular feminization syndrome
7527.31 Pseudo-hermaphroditism NOS
7527.41 Gonadal dysgenesis
7527.51 Gynandrism
7527.99 Ambiguous genitalia / Indeterminate sex NOS

Other anomalies of the male genital tract

7528.01 Absence of both testes / aplasia of testes
7528.02 Single testis / monorchism / absence of testis unilateral
7528.11 Hypoplasia testis
7528.12 Hypoplasia scrotum

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* Adrenortical disorders in female; gonadal disorders in male or associated with chromosomal anomaly are excluded.

7528.13 Hypoplasia scrotum and testis
7528.21 Polyorchism
Excludes : congenital hydrocele (see code 7786.--)
7528.28 Other anomalies of testis or scrotum
7528.31 Atresia of vas deferens / seminal ducts / spermatic cord / ejaculatory duct
7528.41 Absence of prostate / aplasia
7528.48 Other anomalies of vas deferens and prostate
7528.51 Absence / aplasia of penis
7528.52 Displaced penis / retroscrotal penis
7528.53 Hypoplastic / micro penis
7528.54 Absence of foreskin
7528.55 Hooded penis
7528.56 Double penis
7528.57 Cyst on penis
7528.68 Other specified anomalies of penis
7528.71 Cysts of embryonal remains in male / persistent Wolffian duct*

7528.78 Other specified anomalies of male genital tract

7529.99 Unspecified anomalies of genital organs

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* NOTE Hydatid of Morgagni is in females only (see code 7521.21)

CONGENITAL ANOMALIES OF THE URINARY SYSTEM

Renal agenesis and dysgenesis

7530.01 Bilateral renal agenesis / absence / aplasia
7530.02 Bilateral renal hypoplasia / dysplasia
7530.03 Oligohydramnios sequence / Potter sequence*
7530.11 Unilateral renal agenesis / absence / aplasia
7530.12 Unilateral renal hypoplasia / dysplasia
7530.9- Renal agenesis , unspecified / absence / aplasia

Cystic kidney disease

7531.01 Simple / single / solitary / renal cyst or cysts
7531.11 Polycystic kidneys, infantile type / sponge kidney, polycystic kidney of the newborn/ Potter type I
7531.12 Autosomal recessive polycystic kidney disease / congenital hepatic fibrosis / Carolli disease) []
7531.21 Polycystic kidneys, adult type / adult polycystic kidney disease, Potter type III
7531.22 Autosomal dominant polycystic kidney disease []
7531.3- Polycystic kidney NOS / unspecified polycystic kidney disease
7531.41 Medullary cystic disease / medullary type polycystic kidneys / cystic disease of the renal medulla / nephronophthisis, juvenile type
7531.42 Autosomal recessive polycystic kidneys, medullary type []
7531.51 Medullary cystic disease / medullary type polycystic kidneys / cystic disease of the renal medulla, adult type
7531.52 Autosomal dominant polycystic kidneys, medullary type []
7531.61 Multicystic renal dysplasia, enlarged kidney type / multicystic kidney / multilocular cysts / Potter type IIa
7531.62 Multicystic renal dysplasia, reduced kidney type / multicystic kidney / multilocular cysts / Potter type IIb
7531.7- Polycystic kidneys due to urethral obstruction / cortical cysts / Potter type IV
7531.81 Cysts of Bowman's capsule / glomerular cysts
7531.88 Other specified polycystic kidney disease
7531.89 Medullary sponge kidney, unspecified

Obstructive defects of renal pelvis and ureter

7532.01 Congenital hydronephrosis
7532.1- Atresia, stricture or stenosis of ureter or ureteropelvic junction
7532.21 Megaureter / hydro-ureter / dilated ureter
7532.22 Retrocaval ureter
7532.3- Vesicoureteral reflux grade I or II / dysfunction or insufficiency of the vesicoureteral junction
7532.4- Vesicoureteral reflux grade III or IV
7532.9- Upper urinary tract obstruction NOS

Other specified anomalies of kidney

7533.0- Accessory / supernumerary kidney
7533.1- Double or triple kidney and pelvis
7533.2- Lobulated, fused or horseshoe kidney
7533.3- Ectopic kidney / renal ectopia

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* NOTE The pathological cause of the sequence should be specified

7533.4- Enlarged, hyperplastic or giant kidney
7533.5- Congenital renal calculi
7533.61 Congenital glomerular disease
7533.62 Congenital nephritis,
7533.63 Congenital nephrosis / congenital nephrotic syndrome
7533.8- Other specified anomalies of kidney and renal pelvis

Other specified anomalies of ureter

7534.01 Absence of ureter
7534.1- Duplication of ureter / accessory ureter / bifid ureter
7534.2- Ectopic ureter
7534.3- Ureterocele
7534.8- Other specified anomalies of ureter

Exstrophy of bladder or cloaca

7535.01 Exstrophy of urinary bladder sequence
7535.03 Extroversion of bladder
7535.1- Exstrophy of cloaca sequence / ectrophia splanchnica / OEIS complex
7535.2- Superior vesical fissure

Atresia and other congenital stenosis of urethra and bladder neck

7536.01 Posterior urethral valves / Urethral valvular obstruction
7536.1- Atresia, obstruction or stenosis of bladder neck
7536.2- Obstruction, obliteration, atresia or stenosis of urethra
7536.3- Obstruction, atresia or stenosis of urinary meatus
7536.4- Early urethral obstruction sequence
7536.9- Unspecified obstruction of the lower urinary tract

Anomalies of urachus

7537.0- Patent urachus
7537.1- Cyst of urachus
7537.8- Other anomaly of urachus
7537.9- Unspecified anomaly of urachus

Other specified anomalies of bladder and urethra

7538.01 Absence / agenesis of bladder or urethra
7538.1- Ectopic bladder
7538.2- Diverticulum or hernia of bladder
7538.3- Prolapse of bladder (mucosa)
7538.4- Double urethra or urinary meatus
7538.5- Ectopic urethra or urethral orifice
7538.61 Rectovesical fistula
7538.62 Other congenital digestive-urinary tract fistulae
7538.7- Urethral fistula NOS
7538.81 Neurogenic bladder dysfunction
7538.88 Other specified anomalies of bladder and urethra

Megacime forte rule.

Unspecified anomalies of urinary system

7539.09 Unspecified anomaly of kidney
7539.19 Unspecified anomaly of ureter
7539.29 Unspecified anomaly of bladder
7539.39 Unspecified anomaly of urethra
7539.99 Unspecified anomaly of urinary system

CERTAIN CONGENITAL MUSCULOSKELETAL DEFORMITIES

Of skull, face and jaw

Excludes : dentofacial anomalies, syphilitic saddle nose

7540.01 Asymmetry of face
7540.02 Cayler syndrome / Asymmetric crying facies []
7540.1- Compression of face / Potter's facies
7540.21 Congenital deviation of nasal septum
7540.22 Bent nose, deviated nose
7540.23 Beak like nose / downcurved nose
7540.24 Broad nasal bridge
7540.25 Broad nasal tip
7540.26 Other abnormalities of the bridge of the nose
7540.3- Dolichocephaly
Excludes : dolichocephaly due to craniosynostosis (see code 7560.05)
7540.41 Depression in skull
7540.42 Flat occiput
7540.51 Plagiocephaly, oblique head deformation
Excludes : plagiocephaly due to craniosynostosis (see code 7560.06)
7540.61 Brachycephaly
Excludes : acrobrachycephaly (see 7556.02)
7540.62 Ridge metopic suture
7540.64 Mandibular asymmetry
7540.65 Asymmetric head
7540.68 Other specified anomalies of skull or jaw
7540.69 Anomalies of skull or jaw, unspecified

754052
754053
754054
754055
754089
754099

Of sternocleidomastoid muscle

7541.1- Congenital sternocleidomastoid torticollis
7541.2- Congenital contracture of sternocleidomastoid muscle
7541.3- Congenital sternocleidomastoid hematoma
7541.4- Congenital sternocleidomastoid tumour

Of spine

7542.01 Congenital postural scoliosis, cervical
7542.02 Congenital postural scoliosis, dorsal
7542.03 Congenital postural scoliosis, lumbar
7542.04 Congenital postural scoliosis, dorso-lumbar
7542.05 Congenital postural scoliosis, all spine
7542.11 Congenital postural lordosis, cervical
7542.12 Congenital postural lordosis, dorsal
7542.13 Congenital postural lordosis, lumbar
7542.14 Congenital postural lordosis, dorso-lumbar
7542.15 Congenital postural lordosis, all spine
7542.2- Congenital postural curvature of spine, NOS

Congenital dislocation of hip

7543.01 Congenital dislocation of hip, bilateral
7543.02 Congenital dislocation of left hip
7543.03 Congenital dislocation of right hip
7543.04 Congenital dislocation of hip, NOS

Congenital dysplasia of hips? 7543 —?

Unstable hip
7543.11 Preluxation of hip
7543.12 Subluxation of hip
7543.13 Predislocation status of hip at birth
7543.2- Clicking hip
7543.99 Unstable hip NOS

Congenital genu recurvatum and bowing of long bones of leg

7544.01 Bowing, femur
7544.11 Bowing, tibia
7544.12 Bowing, fibula
7544.13 Bowing, tibia and fibula
7544.2- Bow legs NOS
7544.3- Genu recurvatum
7544.4- Dislocation of knee, congenital

Varus deformities of feet

7545.0- Talipes equinovarus / club foot varus
7545.1- Talipes calcaneovarus / dorsiflexed foot
7545.21 Metatarsus varus
7545.22 Metatarsus adductus
7545.3- Complex varus deformities
7545.9- Unspecified varus deformities of feet

Valgus deformities of feet

7546.01 Talipes equinovalgus / club foot valgus
7546.02 Talipes calcaneovalgus
7546.03 Metatarsus valgus
7546.1- Congenital pes planus
7546.9- Unspecified valgus deformities of feet

Other deformities of feet

7547.01 Pes equinus
7547.02 Pes cavus
7547.1- Claw foot
7547.2- Short achilles tendon
7547.31 Club foot NOS, talipes NOS
7547.8- Other specified deformities of foot NEC
7547.9- Congenital deformities of foot NOS

Other specified deformities of thorax and limbs

7548.0- Pectus carinatum / pigeon chest
7548.11 Pectus excavatum / funnel chest
7548.12 Narrow thorax
7548.13 Bell shaped chest
7548.14 Shield chest
7548.2- Other anomalies of chest wall

Other specified deformities of upper limbs

7548.3- Dislocation of elbow
7548.41 Club hand
7548.42 Club fingers
7548.43 Radial deviation of hand
7548.44 Ulnar deviation of hand
7548.5- Spade-like hand
7548.6- Overlapping fingers
7548.8- Other specified

Other congenital anomalies of limbs

Polydactyly, hand
7550.01 Polydactyly post-axial / accessory fingers, bilateral
7550.02 Polydactyly post-axial / accessory fingers, left
7550.03 Polydactyly post-axial / accessory fingers, right
7550.04 Polydactyly post-axial / accessory fingers, unknown
7550.05 Long fingers
7550.11 Polydactyly pre-axial / accessory thumbs, bilateral
7550.12 Polydactyly pre-axial / accessory thumb, left
7550.13 Polydactyly pre-axial / accessory thumb, right
7550.14 Polydactyly pre-axial / accessory thumb, unknown
7550.15 Triphalangeal thumb / finger like thumb
7550.16 Highly inserted thumb
7550.17 Broad thumb
7550.18 Bifid thumb

Polydactyly, foot
7550.21 Polydactyly post-axial / accessory toes, bilateral
7550.22 Polydactyly post-axial / accessory toes, left
7550.23 Polydactyly post-axial / accessory toes, right
7550.24 Polydactyly post-axial / accessory toes, unknown
7550.25 Polydactyly pre-axial / big toe, bilateral
7550.26 Polydactyly pre-axial / big toe, left
7550.27 Polydactyly pre-axial / big toe, right
7550.28 Polydactyly pre-axial / big toe, unknown
7550.31 Polydactyly axial
7550.32 Polydactyly multiple
7550.33 Polydactyly with duplication of structure of limb (e.g. double humerus or structure)
7550.34 Broad great toe

*Polysyndactyly hand or foot**
7550.41 Polysyndactyly fingers, bilateral
7550.42 Polysyndactyly fingers, left
7550.43 Polysyndactyly fingers, right
7550.44 Polysyndactyly fingers, unknown
7550.45 Polysyndactyly toes, bilateral
7550.46 Polysyndactyly toes, left
7550.47 Polysyndactyly toes, right
7550.48 Polysyndactyly toes, unknown
7550.5- Wide groove between 1st and 2d toes
7550.9- Accessory digits NOS

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* NOTE Both extra and fused digits

Syndactyly, hand

7551.01 Fused fingers, isolated skin syndactyly
7551.02 Fused fingers associated with absence of fingers
7551.03 Osseous syndactyly of fingers
7551.04 Symphalangism of fingers / synostosis
7551.1- Webbed fingers

7551.09 Fused fingers NOS

Syndactyly, foot

7551.21 Fused toes, isolated skin syndactyly
7551.22 Fused toes, associated with absence of toes
7551.23 Syndactyly 2nd and 3rd toes
7551.24 Osseous syndactyly of toes
7551.25 Synostosis of toes
7551.3- Webbed toes
7551.9- Unspecified syndactyly of foot

7551.29 Fused toes NOS

ICD-BPA 9

Q71

Reduction deformities upper limb

Terminal transverse defects and proximal intercalary defects

7552.01 Complete absence of upper limb / amelia of upper limb, bilateral
7552.02 Complete absence of upper limb / amelia of upper limb, left
7552.03 Complete absence of upper limb / amelia of upper limb, right
7552.04 Complete absence of upper limb / amelia of upper limb, unknown

7552.11 Absence of upper arm and forearm with hand present, phocomelia of upper limb, bilateral
7552.12 Absence of upper arm and forearm with hand present / phocomelia, left
7552.13 Absence of upper arm and forearm with hand present / phocomelia, right
7552.14 Absence of upper arm and forearm with hand present / phocomelia, unknown

7552.21 Absence of forearm only, bilateral
7552.22 Absence of forearm only, left
7552.23 Absence of forearm only, right
7552.24 Absence of forearm only, unknown

7552.31 Absence of forearm and hand / hemimelia of upper limb, bilateral
7552.32 Absence of forearm and hand / hemimelia of upper limb, left
7552.33 Absence of forearm and hand / hemimelia of upper limb, right
7552.34 Absence of forearm and hand / hemimelia of upper limb, unknown

7552.41 Absence of hand / acheiria, bilateral
7552.42 Absence of hand, left
7552.43 Absence of hand, right
7552.44 Absence of hand, unknown
7552.45 Absence of fingers / adactylia, bilateral
7552.46 Absence of fingers, left
7552.47 Absence of fingers, right
7552.48 Absence of fingers, unknown
Excludes : Lobster claw hand (see code 7555.1)

7552.51 Total absence of one or more phalanx / ectrodactyly of upper limb, bilateral
7552.52 Total absence of one or more phalanx, left hand
7552.53 Total absence of one or more phalanx, right hand
7552.54 Total absence of one or more phalanx, unknown hand
7552.58 Short arm NOS

7552.0 abs of upper limb

7552.1 abs of upper arm

7552.2 abs of forearm

7552.3 abs of forearm and hand

7552.4 abs of hand and/or fingers

7552.5 shortening of arm

Longitudinal defects upper limbs, arm radial side

6
- 7552.61 Pre-axial absence or severe hypoplasia of radius, thumb and first metacarpal, bilateral
- 7552.62 Pre-axial absence or severe hypoplasia of radius, thumb and first metacarpal, left
- 7552.63 Pre-axial absence or severe hypoplasia of radius, thumb and first metacarpal, right
- 7552.64 Pre-axial absence or severe hypoplasia of radius, thumb and first metacarpal, unknown
- 7552.65 Absence of thumb
- 7552.68 TAR syndrome / Radial aplasia-thrombocytopenia syndrome []

Longitudinal defects upper limbs, arm ulnar side

4
- 7552.71 Post-axial absence or severe hypoplasia of ulna, 5th finger and 5th metacarpal, bilateral
- 7552.72 Post-axial absence or severe hypoplasia of ulna, 5th finger and 5th metacarpal, left
- 7552.73 Post-axial absence or severe hypoplasia of ulna, 5th finger and 5th metacarpal, right
- 7552.74 Post-axial absence or severe hypoplasia of ulna, 5th finger and 5th metacarpal, unknown

OK : intégrée au EURAS + inclus au SPRAS 75527

- 7552.78 Other longitudinal reduction deformity of arm

inclus : hypoplasie tarse.

- 7552.8- Other specified reduction deformity of arm
- 7552.9- Unspecified reduction deformity of arm / Amputation of upper limb NOS

Reduction deformities of lower limb

Q72

Terminal transverse defects and proximal intercalary defects

- 7553.01 Complete absence of lower limb / amelia of lower limb, bilateral
- 7553.02 Complete absence of lower limb / amelia of lower limb, left
- 7553.03 Complete absence of lower limb / amelia of lower limb, right
- 7553.04 Complete absence of lower limb / amelia of lower limb, unknown
- 7553.11 Absence of thigh and lower leg with foot present / phocomelia of lower limb / bilateral
- 7553.12 Absence of thigh and lower leg with foot present / phocomelia of lower limb, left
- 7553.13 Absence of thigh and lower leg with foot present / phocomelia of lower limb, right
- 7553.14 Absence of thigh and lower leg with foot present / phocomelia of lower limb, unknown
- 7553.15 Isolated femoral agenesis / Absence of thigh with lower leg and foot present / bilateral
- 7553.16 Isolated femoral agenesis / Absence of thigh with lower leg and foot present / left
- 7553.17 Isolated femoral agenesis / Absence of thigh with lower leg and foot present / right
- 7553.18 Isolated femoral agenesis / Absence of thigh with lower leg and foot present / unknown
- 7553.21 Absence of lower leg only, bilateral
- 7553.22 Absence of lower leg only, left
- 7553.23 Absence of lower leg only, right
- 7553.24 Absence of lower leg only, unknown
- 7553.31 Absence of lower leg and foot / hemimelia of lower limb / bilateral
- 7553.32 Absence of lower leg and foot / hemimelia of lower limb / left
- 7553.33 Absence of lower leg and foot / hemimelia of lower limb / right
- 7553.34 Absence of lower leg and foot / hemimelia of lower limb / unknown
- 7553.41 Absence of foot, bilateral
- 7553.42 Absence of foot, left

7553.43 Absence of foot, right
7553.44 Absence of foot, unknown
7553.45 Absence of toes / adactylia / bilateral
7553.46 Absence of toes / adactylia / left
7553.47 Absence of toes / adactylia / right
7553.48 Absence of toes / adactylia / unknown
Excludes : Claw foot (see code 7547.1)
7553.51 Total absence of one or more phalanx /ectrodactyly of lower limb / bilateral
7553.52 Total absence of one or more phalanx, left foot
7553.53 Total absence of one or more phalanx, right foot
7553.54 Total absence of one or more phalanx, unknown foot
7553.58 Short leg NOS

long. reduct. def of femur ?

Longitudinal defects, leg tibial side
7553.61 Pre-axial absence or severe hypoplasia of tibia, big toe, first metatarsal, bilateral
7553.62 Pre-axial absence or severe hypoplasia of tibia, big toe, first metatarsal, left
7553.63 Pre-axial absence or severe hypoplasia of tibia, big toe, first metatarsal, right
7553.64 Pre-axial absence or severe hypoplasia of tibia, big toe, first metatarsal, unknown
7553.65 Absence of big toe

Longitudinal defects, leg fibular side
7553.71 Post-axial absence or severe hypoplasia of fibula, 5th toe, 5th metatarsal, bilateral
7553.72 Post-axial absence or severe hypoplasia of fibula, 5th toe, 5th metatarsal, left
7553.73 Post-axial absence or severe hypoplasia of fibula, 5th toe, 5th metatarsal, right
7553.74 Post-axial absence or severe hypoplasia of fibula, 5th toe, 5th metatarsal, unknown
7553.78 Other longitudinal reduction deformity of leg ← ? ?
7553.8- Other longitudinal defects ←
7553.9- Unspecified longitudinal defects / Amputation of lower limb NOS

Reduction deformities, unspecified limb

Q73

7554.01 Absence of unspecified limb / Amelia of unspecified limb
7554.1- Phocomelia of unspecified limb
7554.2- Amputation of unspecified limb
7554.3- Longitudinal reduction deformity of unspecified limb
7554.4- Absent digits of unspecified limb
7554.8- Other specified reduction deformity, unspecified limb
7554.9- Unspecified reduction deformity of unspecified limb

Other anomalies of upper limb, including shoulder girdle

Anomalies of fingers and hands
7555.01 Acrocephalosyndactyly type I / Apert []
7555.02 Acrocephalosyndactyly type II / Carpenter []
7555.03 Acrocephalosyndactyly type III / Saethre-Chotzen []
7555.04 Acrocephalosyndactyly type V / Pfeiffer syndrome / Noack syndrome []
7555.05 Macrodactylia
7555.06 Camptodactyly / Flexion deformities of finger
7555.07 Clinodactyly

7555.08 AASE syndrome

Anomalies of hand
7555.11 Lobster hand / split hand, bilateral
7555.12 Typical split / lobster hand, left
7555.13 Typical split / lobster hand, right
7555.14 Typical split / lobster hand, unknown
7555.15 Monodactyly, bilateral
7555.16 Monodactyly, left
7555.17 Monodactyly, right
7555.18 Monodactyly, unknown

Anomalies of wrist
7555.21 Accessory carpal bones
7555.22 Madelung's deformity

Anomalies of forearm
7555.31 Radio-ulnar dysostosis
7555.32 Radio-ulnar synostosis

Anomalies of elbow and upper arm
7555.41 Cubitus valgus

Anomalies of shoulder
7555.51 Cleidocranial dysostosis / Cleidocranial dysplasia []
7555.52 Sprengel's deformity
7555.53 Absent scapulae
7555.54 Webbed shoulder

7555.68 Other anomalies of whole arm

7555.81 Shortening of arm / micromelia
7555.82 Shortening of fingers / brachydactyly / bilateral
7555.83 Shortening of fingers / brachydactyly / unilateral, left
7555.84 Shortening of fingers / brachydactyly / unilateral, right
7555.85 Shortening of fingers / brachydactyly / unilateral, unknown

7555.9- Unspecified anomalies of upper limb

Other anomalies of lower limb, including pelvic girdle

Includes : complex anomalies involving all or part of lower limb

Anomalies of toes
7556.01 Hallux valgus
7556.02 Hallux varus
7556.03 Widespaced 1st and 2nd toes

one of toes 7556.09

Anomalies of foot
7556.11 Rocker bottom foot
7556.12 Typical split/ lobster claw, bilateral
7556.13 Typical split / lobster claw, left
7556.14 Typical split / lobster claw, right
7556.15 Typical split / lobster claw, unilateral, unknown
7556.16 Monodactyly bilateral
7556.17 Monodactyly, foot left
7556.18 Monodactyly, foot right
7556.19 Monodactyly, foot unilateral, unknown

scaphoid < Hand ?

Anomalies of ankle
7556.21 Accessory tarsal bones
7556.22 Astragaloscaphoid synostosis
7556.31 Angulation of tibia

Anomalies of knee
7556.41 Genu valgum
7556.42 Genu varum
7556.43 Absent patella
7556.44 Rudimentary patella
7556.51 Anomalies of upper leg / anteversion of femur

Anomalies of hip
7556.61 Coxa valga
7556.62 Coxa vara

Anomalies of pelvis
7556.71 Fusion of sacroiliac joint
7556.78 Other anomalies of pelvis

Other anomalies of lower limb
7556.81 Shortening of leg / micromelia
7556.82 Shortening of toes, bilateral
7556.83 Shortening of toes, unilateral, left
7556.84 Shortening of toes, unilateral, right
7556.85 Shortening of toes, unilateral, unknown
89 oth specified anomalies of lower limb
7556.9- Unspecified anomalies of lower limb

Other specified anomalies of unspecified limb
7558.01 Arthrogryposis multiplex congenita / fetal akinesia sequence / congenital contractures
7558.02 Distal arthrogryposis, contractures of fingers and toes
7558.03 Distal arthrogryposis, contractures of fingers
7558.04 Distal arthrogryposis, contractures of toes
7558.11 Larsen's syndrome []
7558.8- Other specified anomalies of unspecified limb

Unspecified anomalies of unspecified limb
7559.-- Unspecified anomalies of unspecified limb

OTHER CONGENITAL MUSCULOSKELETAL ANOMALIES

Anomalies of skull and face bones

Excludes : dentofacial anomalies and skull defects associated with anomalies of brain, such as : anencephalus, encephalocele, hydrocephalus, microcephaly

7560.01 Oxycephaly / Turricephaly / Acrocephaly / Fusion of sagittal and coronal sutures
7560.02 Acrobrachycephaly / Fusion of the coronal sutures
7560.03 Trigonocephaly / Arrest of development of the frontal bone
7560.04 C syndrome / Opitz trigonocephaly syndrome []
7560.05 Scaphocephaly / Leptocephaly / Bathmocephaly / Dolichocephaly due to craniostenosis / Fusion of sagittal suture
7560.06 Plagiocephaly / Closure of half of the coronal suture
Excludes : plagiocephaly not due to craniosynostosis (see code 7540.51)
7560.07 Cloverleaf skull / Kleeblattschadel
7560.08 Craniosynostosis NOS
7560.11 Craniofacial dysostosis / Crouzon disease []
7560.21 Hypertelorism
7560.22 Hypotelorism
7560.31 Pierre Robin sequence []
7560.32 Stickler syndrome / Arthro-ophthalmopathy []
7560.41 Mandibulofacial dysostosis / Treacher-Collins syndrome / Franceschetti []
7560.51 Oculomandibular dysostosis / Hallerman-Streiff syndrome / François dyscephaly
7560.61 Goldenhar syndrome / oculo-auriculo-vertebral first and second arch syndrome/ hemifacial microsomia
7560.7- Localised skull defects
7560.8- Other specified anomalies of skull and face
7560.9- Unspecified anomalies of skull and face

Anomalies of spine

7561.01 Spina bifida occulta uncomplicated
7561.11 Klippel-Feil sequence / syndrome []
7561.12 Wildervanck syndrome / cervico-oculo-acoustic syndrome
7561.21 Kyphosis
7561.22 Kyphoscoliosis
7561.23 Gibbus
7561.3- Congenital spondylolisthesis
7561.4- Anomalies of cervical vertebrae
7561.51 Agenesis of thoracic vertebrae
7561.52 Hypoplasia of thoracic vertebrae
7561.53 Thoracic hemivertebrae
7561.54 Thoracic supernumerary vertebrae
7561.58 Other anomalies of thoracic vertebrae
7561.61 Hypoplasia of lumbar vertebrae
7561.62 Agenesia of lumbar vertebrae
7561.63 Lumbar hemivertebrae
7561.64 Lumbar supernumerary vertebrae
7561.68 Others anomalies of lumbar vertebrae
7561.71 Sacrococcygeal agenesia
7561.72 Sacrococcygeal hypoplasia
7561.73 Sacrococcygeal hemivertebrae
7561.74 Sacrococcygeal supernumerary vertebrae
7561.75 Sacral dimple
7561.76 Sacral sinus
7561.78 Other sacrococcygeal anomalies

7561.8- Other specified anomalies of spine
7561.9- Unspecified anomalies of spine

Other anomalies of ribs and sternum

7562.-- Supernumerary rib in cervical region

7563.01 Absence of ribs
7563.1- Mis-shapen ribs
7563.2- Fused ribs
7563.3- Extra ribs
7563.4- Other anomalies of ribs
7563.51 Absence of sternum / asternia
7563.52 Absence of part of sternum
7563.61 Mis-shapen sternum
7563.62 Bifid sternum
7563.63 Cleft of sternum
Excludes : pectus excuvatum (see code 7548.11), carinatum (see code 7548.0)
7563.8- Other anomalies of sternum
7563.9- Anomalies of thoracic cage, unspecified

Chondrodystrophy, osteochondrodystrophy, chondrodysplasia, osteochondrodysplasia

7564.01 Asphyxiating thoracic dysplasia / Jeune []
7564.02 Hypochondrogenesis
7564.03 Fibrochondrogenesis
7564.04 Atelosteogenesis
7564.05 Short rib syndromes (with or without polydactyly),
7564.1- Ollier disease / Osteochondromatosis / Enchondromatosis / Dyschondroplasia
Excludes : Campomelic dysplasia (see code 7598.65)
7564.2- Chondrodysplasia with haemangioma / Maffuci syndrome / Kast syndrome
7564.31 Achondroplasia / Achondroplastic dwarfism []
7564.32 Dyssegmental dysplasia / Dyssegmental dwarfism
7564.33 Hypochondroplasia []
7564.34 Acromesomelic dwarfism []
7564.35 Mesomelic dysplasia / type Nievergelt / type Langer / homozygous dyschondrosteosis / type Robinow / type Rheinardt,

7564.41 Diastrophic dwarfism / Diastrophic dysplasia / Nanism []
7564.42 Metatropic dwarfism / Metatropic dysplasia []
7564.43 Kniest dysplasia []
7564.44 Acrodystosis
7564.45 Thanatophoric dysplasia
7564.46 Thanatophoric dysplasia with clover-leaf skull
7564.51 Cartilage-hair hypoplasia
7564.52 Pseudo-achondroplasia / Metaphyseal chondrodysplasia Mc Kusick type []
7564.61 Spondylo-epiphyseal dysplasia congenita, autosomal dominant form []
7564.62 Spondylo-thoracic dysplasia / Tarcho Levin [] JARCHO et JS TARCHO ! idée incert.
7564.71 Exostosis
7564.72 Diaphyseal aclasia
7564.8- Other specified chondrodystrophies
7564.9- Unspecified chondrodystrophies

Osteodystrophies

Includes : osteopsathyrosis, fragilitas ossium
7565.01 Osteogenesis imperfecta type I / Blue sclera / Losbstein disease []
7565.02 Osteogenesis imperfecta type II / Congenita NOS / Vrolic syndrome []
7565.03 Osteogenesis imperfecta type III / / Neonatal lethal []
7565.04 Osteogenesis imperfecta type IV / Normal sclera []
7565.11 Polyostotic fibrous dysplasia / Albright-Mc Cune-Sternberg syndrome []
7565.21 Chondroectodermal dysplasia / Ellis van Creveld []
7565.31 Infantile cortical hyperostosis / Caffey's syndrome []
7565.41 Osteopetrosis / Albers-Schonberg syndrome / Marble bones []
7565.51 Progressive diaphyseal dysplasia / Engelmann's syndrome
7565.61 Osteopoikilosis
7565.62 Spondylothoracic dysplasia / Jarcho Levin []
7565.63 Marden Walker []
7565.71 Multiple ephiphyseal dysplasia / Conradi Hünermann syndrome / Chondrodysplasia punctata []
7565.72 Chondro-dysplasia punctata rhizomelic syndrome []
7565.81 Dyschondrosteosis / Mesomelic dwarfism []
7565.88 Other specified osteodystrophies
7565.9 Unspecified osteodystrophies

Anomalies of diaphragm

7566.0- Absence of diaphragm

Congenital diaphragmatic hernia
7566.11 Left hemidiaphragmatic defect
7566.12 Right hemidiaphragmatic defect
7566.13 Hernia through the foramen of Morgagni
7566.14 Hernia through the foramen of Bochdalek
7566.15 Postero-lateral diaphragmatic hernia
7566.18 Atypical hernia of diaphragm
7566.19 Unspecified type of diaphragmatic hernia
7566.2- Eventration of diaphragm
7566.8- Other specified anomaly of diaphragm
7566.9- Unspecified anomaly of diaphragm

Anomalies of abdominal wall

7567.01 Exomphalos / omphalocele
7567.02 Abdominal muscle, aplasia or severe hypoplasia
7567.03 Diastasis recti
7567.11 Laparoschisis / gastroschisis
Excludes : umbilical hernia (see code 5531)
7567.21 Prune belly syndrome
7567.2- Other specified anomaly of abdominal wall
7567.9- Unspecified anomaly of abdominal wall

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

7568.01 Poland syndrome []
7568.1- Other absent muscle
7568.2- Absent tendon
7568.31 Nail-patella syndrome / Hereditary osteo-onychodysplasia []
7568.41 Amyotrophia congenita

7568.51 Ehlers-Danlos syndrome Dominant []
7568.52 Ehlers-Danlos syndrome Recessive []
7568.53 Ehlers-Danlos syndrome unspecified
7568.81 Schwartz-Jampel syndrome []
7568.82 Popliteal web syndrome / Popliteal pterygium []
7568.88 Other specified anomalies of muscle, tendon fascia and connective tissue

Unspecified anomalies of musculoskeletal system

7569.01 Unspecified anomalies of muscle
7569.11 Unspecified anomalies of tendon
7569.21 Unspecified anomalies of bone
7569.31 Unspecified anomalies of cartilage
7569.41 Unspecified anomalies of connective tissue
7569.99 Unspecified anomalies of musculoskeletal system NOS

CONGENITAL ANOMALIES OF THE INTEGUMENT

Hereditary oedema of legs / Hereditary trophoedema / Hereditary Lymphoedema

7570.1- Nonne-Milroy's disease []
7570.2- Meige disease []
7570.3- Other specified hereditary oedema of legs
7570.4- Unspecified hereditary oedema of legs

Ichthyosis congenita

7571.0- Harlequin fetus / Ichthyosis Fetalis / Keratum Malignum / Ichthyosis Congenita []
7571.11 Collodion baby (condition seen in Epidermolytic hyperkeratosis)
7571.12 Ichthyosiform erythroderma, (an X-linked Ichthyosis) []
7571.2- Sjogren-Larsson syndrome []
7571.3- Normokinetic Ichthyosis
includes Ichthyosis Vulgaris [] and X-linked Ichthyosis
7571.8- Other specified type of Ichthyosis
7571.9- Unspecified Ichthyosis

Dermatoglyphic anomalies

7572.1- Unilateral palmar simian crease
7572.2- Bilateral palmar simian creases
7572.3- Absent or hypoplastic palmar creases
7572.4- Deep palmar and/or plantar furrows
7572.5- Abnormal patterns on the fingertips
7572.8- Other specified anomalies of dermatoglyphics
7572.9- Unspecified anomalies of dermatoglyphics

Other specified anomalies of skin

7573.0- Specified syndromes, NEC, involving skin anomalies
7573.1- Skin tags
7573.2- Urticaria pigmentosa / Mast cell disease / Mastocytosis []

Epidermolysis Bullosa
7573.31 Epidermolysis bullosa simplex
(includes Koebner type, Ogna type, with mottled pigmentation) []
7573.32 Epidermolysis bullosa letalis []
includes epidermolysis bullosa letalis with pyloric atresia []
7573.33 Epidermolysis bullosa dystrophica []
includes Bart type, Hallopen-Siemens type, inversa, neurotrophica, Pasini type []
7573.38 Other specified type of epidermolysis bullosa
7573.39 Unspecified type of epidermolysis bullosa

Ectodermal dysplasia
Excludes : Ellis-Van-Creveld syndrome []
7573.41 Anhidrotic ectodermal dysplasia includes Christ-Siemens syndrome [], autosomal recessive type, with cleft lip and palate []
7573.42 Hypohidrotic ectodermal dysplasia [] includes type with hypothyroidism and ciliary dyskynesia []
7573.43 Hidrotic ectodermal dysplasia []
7573.48 Other specified type of ectodermal dysplasia

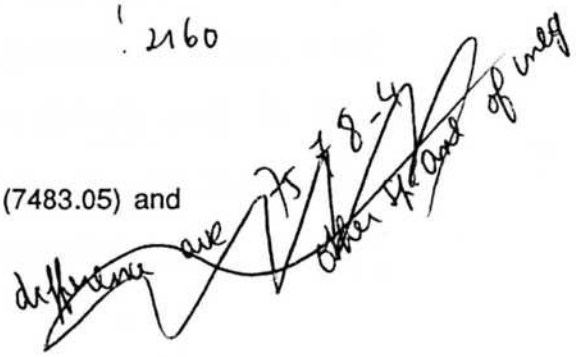
7573.49 Unspecified type of ectodermal dysplasia

Incontinentia Pigmenti
7573.51 Incontinentia pigmenti / Bloch-Sulzberger syndrome []
7573.52 Incontinentia pigmenti achromions / Hypomelanosis of Ito []
7573.60 Xeroderma pigmentosum
includes recessive form [] and dominant form []

Cutis Laxa Hyperelastica / Ehlers-Danlos syndrome
(see code 7568.5)

Birthmarks * Note : Isolated naevi smaller than 4 cm^2 should not be notified
7573.80 *Naevus, capillary / cavernous / flammeus / port wine / strawberry / vascular / haemangioma of skin/ angioma of skin
7573.81 *Naevus, pigmented
7573.82 *Mongolian blue spot
7573.83 *Naevus, other
7573.84 Cystic hygroma of neck
7573.85 Cystic hygroma, other
7573.86 Lymphangioma
Excludes : congenital subglottic haemangioma (7483.05) and lymphangioma (7483.06)

7573.98 Other specified anomalies of skin
7573.99 Unspecified anomalies of skin



Specified anomalies of hair

Excludes : Kinky Hair syndrome / Menkes syndrome (see code 7598.73)
Congenital Alopecia
7574.01 Congenital Universal Alopecia (absence of hair follicles) / Hypotrichosis
7574.02 Localised Alopecia
Excludes: Cartilage-hair hypoplasia (Metaphyseal Chondrodysplasia) (see code 7564.51)
7574.03 Alopecia Areata
7574.08 Other type of alopecia
7574.09 Unspecified type of alopecia
7574.1- Beaded hair / Monilethrix / Isolated beaded hair []
7574.2- Twisted hair / Pili torti / Isolated twisted hair []
7574.31 Taenzer's hair
7574.4- Woolly hair
7574.5- Brittle hair
7574.6- Trichorrhexis nodosa
7574.71 Abnormal hair patterning
7574.72 Widow's peak
7574.73 Frontal upsweep
7574.74 Abnormal hair whorls
7574.8- Other specified anomalies of hair
7574.9- Unspecified anomalies of hair

Specified anomalies of nails

7575.0- Congenital anonychia / Absent nails /Onychoatrophy
7575.1- Enlarged nails / Hypertrophic nails / Onychauxis / Pachyonychia
7575.2- Congenital koilonychia / Spoon nails
7575.3- Congenital leukonychia / White nails

7575.4- Club nails
Excludes: Nail-Patella syndrome (Hereditary Osteo-Onychodysplasia) (see code7568.31)
7575.8- Other specified anomaly of nails
7575.9- Unspecified anomaly of nails

Specified anomalies of breast

7576.0- Absent breast with absent nipple
7576.1- Hypoplastic breast with hypoplastic nipple
7576.2- Accessory (ectopic) breast with nipple
7576.3- Absent nipple
7576.4- Small nipple
7576.5- Accessory nipple / ectopic
7576.8- Other anomalies of breast

Other specified anomaly of integument

7578.8- Other specified anomalies of the integument

Unspecified anomaly of integument

7579.9- Unspecified anomalies of the integument NOS

CHROMOSOMAL ANOMALIES

Down syndrome

7580.01 Trisomy 21 ("free" or "standard" trisomy 21)
7580.02 Trisomy 21, mosaic
7580.11 Trisomy G NOS (no banding)
7580.12 Trisomy G NOS, mosaic
7580.21 Translocation trisomy 21 (with banding) "de novo"
7580.22 Translocation trisomy 21(with banding) inherited
7580.31 Translocation trisomy G (no banding)
7580.41 Down syndrome on clinical grounds (chromosomal analysis not available)
7580.91 Down syndrome NOS

Patau's syndrome (trisomy 13) and other total trisomies of the D group

7581.01 Trisomy 13 (classical Patau's syndrome)
7581.02 Trisomy 13, mosaic
7581.11 Trisomy D NOS (no banding)
7581.12 Trisomy D NOS, mosaic
7581.13 Trisomy 14
7581.14 Trisomy 15
7581.21 Translocation trisomy 13
7581.31 Translocation trisomy D NOS (no banding)
7581.32 Translocation trisomy 14
7581.41 Patau's syndrome on clinical grounds only (chromosomal analysis not available)
7581.91 Patau's syndrome NOS

Edward's syndrome and other total trisomies of E group

7582.01 Trisomy 18 (Edward's syndrome)
7582.02 Trisomy 18, mosaic
7582.03 Trisomy 16
7582.04 Trisomy 17
7582.11 Trisomy E NOS
7582.12 Trisomy 18 on clinical grounds
7582.21 Translocation trisomy 18
7582.31 Translocation trisomy E
7582.91 Edward's syndrome NOS

Autosomal deletion syndromes

Group E (Y excluded)
7583.01 Partial deletion of chromosome 21
7583.02 Partial deletion of chromosome 22
7583.03 Ring (21)
7583.04 Ring (22)

7583.08 Partial deletion of chromosome G NOS
7583.09 Ring G NOS

Chromosome 5
7583.11 Partial deletion of short arm of chromosome 5 "cri du chat"
7583.12 "Cri du chat" syndrome on clinical grounds
7583.13 Partial deletion of long arm of chromosome 5

Chromosome 4
7583.21 Partial deletion of short arm of chromosome 4 "Wolf-Hirschhorn"
7583.22 "Wolf-Hirschhorn" syndrome on clinical ground
7583.23 Partial deletion of long arm of chromosome 4

7583.28 Partial deletion of short arm of chromosome B NOS
7583.29 Partial deletion of long arm of chromosome B NOS

Group D
7583.31 Deletion of long arm of chromosome 13
7583.32 Ring (13)
7583.33 Deletion of long arm of chromosome 14
7583.34 Ring (14)
7583.35 Deletion of long arm of chromosome 15
7583.36 Ring (15)
7583.38 Deletion of long arm of chromosome D NOS
7583.39 Ring (D) NOS

Group E long arms
7583.41 Deletion of long arm of chromosome 18 (18 q) 187-
7583.42 Ring (18)

7583.48 Deletion of long arm of chromosome E
7583.49 Ring (E) NOS

Group E short arms
7583.51 Deletion of short arm of chromosome 18 (18 p) 181-
7583.52 Isochromosome of short arm of chromsome 18
7583.53 Deletion of short arm of chromosome 17
7583.59 Deletion of short arm of chromosome E NOS
7583.6- Monosomy G Mosaicism

Other specified deletion / Loss of autosomal material NEC
7583.71 Deletion of short arm of chromosome 1
7583.72 Deletion of long arm of chromosome 1
7583.73 Deletion of short arm of chromosome 2
7583.74 Deletion of long arm of chromosome 2
7583.75 Deletion of short arm of chromosome 3
7583.76 Deletion of long arm of chromosome 3
7583.77 Deletion of short arm of chromosome 6
7583.78 Deletion of long arm of chromosome 6
7583.79 Deletion of short arm of chromosome 7
7583.81 Deletion of long arm of chromosome 7
7583.82 Deletion of short arm of chromosome 8
7583.83 Deletion of long arm of chromosome 8
7583.84 Deletion of short arm of chromosome 9
7583.85 Deletion of long arm of chromosome 9
7583.86 Deletion of short arm of chromosome 10
7583.87 Deletion of long arm of chromosome 10
7583.88 Deletion of short arm of chromosome 11
7583.89 Deletion of long arm of chromosome 11
7583.91 Deletion of short arm of chromosome 12
7583.92 Deletion of long arm of chromosome 12
7583.93 Deletion of short arm of chromosome 19
7583.94 Deletion of long arm of chromosome 19
7583.95 Deletion of short arm of chromosome 20
7583.96 Deletion of long arm of chromosome 21
7583.98 Deletion of short arm of chromosome C NOS

7583.99 Deletion of long arm of chromosome C NOS

Balanced autosomal translocations in normal individual

7584.1- Balanced autosomal translocation

7584.2- Balanced X autosomal translocation

7584.31 Pericentric inversion chromosome group A
7584.32 Pericentric inversion chromosome group B
7584.33 Pericentric inversion chromosome group C
7584.34 Pericentric inversion chromosome group D
7584.35 Pericentric inversion chromosome group E
7584.36 Pericentric inversion chromosome group F
7584.37 Pericentric inversion chromosome group G
7584.39 Pericentric inversion chromosome NOS

7584.41 Paracentric inversion chromosome group A
7584.42 Paracentric inversion chromosome group B
7584.43 Paracentric inversion chromosome group C
7584.44 Paracentric inversion chromosome group D
7584.45 Paracentric inversion chromosome group E
7584.46 Paracentric inversion chromosome group F
7584.47 Paracentric inversion chromosome group G

Other conditions due to autosomal anomalies

7585.01 Trisomy 8
7585.02 Trisomy 8, mosaic

Other trisomy C syndromes
7585.11 Trisomy 6
7585.12 Trisomy 7
7585.13 Trisomy 8
7585.14 Trisomy 9
7585.15 Trisomy 10
7585.16 Trisomy 11
7585.17 Trisomy 12
7585.19 Trisomy C NOS

Other total trisomy syndromes
(C, D, E, 21 & sex chromosomes excluded)
7585.21 Total trisomy A group
7585.22 Total trisomy B group
7585.23 Total trisomy F group
7585.24 Trisomy 22
7585.29 Total trisomy NOS

Partial trisomy syndromes
7585.31 Partial trisomy 1 NOS
7585.32 Partial trisomy 2NOS
7585.33 Partial trisomy 3 NOS
7585.34 Partial trisomy 4 NOS
7585.35 Partial trisomy 5 NOS
7585.36 Partial trisomy 6 NOS
7585.37 Partial trisomy 7 NOS
7585.38 Partial trisomy 8 NOS
7585.39 Partial trisomy 9 NOS

7585.41 Partial trisomy 10 NOS
7585.42 Partial trisomy 11 NOS
7585.43 Partial trisomy 12 NOS
7585.44 Partial trisomy 13 NOS
7585.45 Partial trisomy 14 NOS
7585.46 Partial trisomy 15 NOS
7585.47 Partial trisomy 16 NOS
7585.48 Partial trisomy 17 NOS
7585.49 Partial trisomy 18 NOS
7585.51 Partial trisomy 19 NOS
7585.52 Partial trisomy 20 NOS
7585.53 Partial trisomy 21 NOS
7585.54 Partial trisomy 22 NOS
7585.55 Additional marker chromosome
7585.59 Partial trisomy NOS

Other specified anomalies of autosomes NOS
7585.81 Triploidy G1. XXX
7585.82 Triploidy G1. XXY
7585.83 Triploidy, mosaic
7585.84 Tetraploidy
7585.85 Tetraploidy, mosaic

Unspecified anomalies of unspecified antosome
7585.9- Unspecified anomalies of unspecified antosome

Aberrations of the X chromosome in absence of a Y chromosome

7586.0- Classical Turner's syndrome, karyotype 45 X
7586.11 Mosaic 45 X / 46 XX
7586.12 Partial X deletion (short arms)
7586.13 Partial X deletion (long arms)
7586.14 Isochromosome X
7586.15 Ring chromosome X
7586.2- Turner's phenotype, karyotype normal / Bonnevie Ullrich syndrome
7586.31 47 XXX syndrome / trisomy X
7586.32 48 XXXX syndrome / tetrasomy X
7586.33 49 XXXXX syndrome / pentasomy X
7586.34 Mosaic 46 XX / 47 XXX
7586.91 Turner's phenotype, karyotype not done or unspecified
7586.92 Fetal Turner's syndrome

Klinefelter's syndrome

7587.0- Klinefelter's phenotype, karyotype 47 XXY
7587.11 Mosaic 46 XY / 47 XXY
7587.12 Male XX
7587.13 48 XXXY
7587.14 49 XXXXY
7587.15 47 XXY 47 + Y Y ?
7587.21 Partial Y deletion (short arms)
7587.22 Partial Y deletion (long arms)
7587.23 Isochromosome Y (short arms)
7587.24 Isochromosome Y (long arms)
7587.25 Ring chromosome Y
7587.26 Pericentric inversion Y
7587.9- Klinefelter's syndrome NOS

Mosaic XO/XX
XY/XXY
758.81
758.82
758.83
75888 other spec. cond. due to SA chrom. anal.

Other conditions due to sex chromosome anomalies

7588.0- Mosaic XO / XY
7588.61 Triploidy 69 XXX
7588.62 Triploidy 69 XXY
7588.81 XY, female phenotype
7588.9- Unspecified conditions due to sex chromosome anomalies

Conditions due to anomaly of unspecified chromosomes

7589.0- Mosaicism NOS
7589.1- Additional chromosome(s) NOS
7589.2- Deletion of chromosome(s) NOS
7589.3- Duplication of chromosome(s) NOS
7589.8- Other specified conditions due to anomaly of unspecified chromosomes
7589.9- Unspecified conditions due to anomaly of unspecified chromosomes

CONGENITAL ANOMALIES OF ENDOCRINE GLANDS

Anomalies of spleen

7590.01 Asplenia / Ivemark syndrome/ Right atrial isomerism
7590.02 Absence of spleen
7590.1- Hypoplasia of spleen
7590.2- Hyperplasia of spleen
7590.3- Mis shapen spleen
7590.41 Accessory spleen
7590.42 Polysplenia syndrome / Left atrial isomerism
7590.5- Ectopic spleen / wandering spleen
7590.81 Cyst of the spleen
7590.82 Hemangioma of the spleen
7590.83 Other specified anomaly
7590.9- Unspecified anomaly of spleen

Anomalies of adrenal gland

Excludes : Congenital adrenal hyperplasia (see code 2552)
Cushing syndrome (see code 2550)
7591.01 Absence / Aplasia of adrenal gland unilateral
7591.02 Absence / Aplasia of adrenal gland bilateral
7591.1- Hypoplasia of adrenal gland
7591.2- Accessory adrenal gland
7591.3- Ectopic adrenal gland
7591.8- Other specified anomaly of adrenal gland
7591.9- Unspecified anomaly of adrenal gland

Anomalies of other endocrine glands

Excludes : Metabolic disorders
7592.01 Absence of pituitary gland
7592.02 Hypoplasia of pituitary gland
7592.03 Other anomalies of pituitary gland
7592.04 Unspecified anomalies of pituitary gland
7592.11 Congenital athyrosis / absence of thyroid
7592.12 Sublingual ectopic thyroid
7592.14 Congenital hyperthyroidism
Excludes : Goitrous cretinism familial type (see code 2439.1-)
Pendred's syndrome (see code 2439.0-)
7592.18 Other anomalies of thyroid gland
7592.19 Unspecified anomalies of thyroid gland
7592.21 Thyroglossal duct anomalies
7592.22 Thyroglossal cyst
7592.31 Absence of parathyroid gland
7592.38 Other specified anomalies of parathyroid gland
7592.39 Unspecified anomalies of parathyroid gland
7592.41 Total agenesis of the thymus / Absence of thymus
7592.42 Hypoplasia of the thymus
7592.43 Abnormal position of the thymus
7592.48 Other specified anomalies of the thymus
7592.49 Unspecified anomalies of thymus
7592.8- Other specified anomalies of endocrine glands
7592.9- Unspecified anomalies of endocrine glands

OTHER CONGENITAL ANOMALIES

Situs inversus

Excludes : Asplenia syndrome (see code7590.01)
Polysplenia syndrome (see code7590.42)

7593.01 Situs inversus (dextrocardia and trilobed lung, liver and gallbladder on the left side)
7593.02 Dextrocardia without abdominal anomalies
7593.1- Situs inversus with levocardia
7593.2- Situs inversus thoracis / reversed thoracic contents
7593.3- Situs inversus abdominis / reversed abdominal contents
7593.41 Kartagener's syndrome (triad)
7593.8- Other specified situs inversus
7593.9- Unspecified situs inversus

Conjoined twins

7594.0- Dicephalus / two heads
7594.1- Craniopagus / head-to-head fusion
7594.2- Thoracopagus / chest-to-chest fusion
7594.3- Xiphopagus / xiphoid-pelvis fusion
7594.4- Pygopagus / pelvis-to-pelvis fusion
7594.5- Cephalothoracopagus
7594.6- Duplicatas anterior /complete fusion of lower parts of bodies but upper parts separate (excluding dicephalus)
7594.7- Duplicatas posterior /upper parts of bodies completely merged but lower parts still separate
7594.80 Other specified side-to-side fusion
7594.81 Unequal conjoined twins
7594.9- Unspecified side-to-side fusion

Phakomatoses

7595.01 Tuberous sclerosis / Bourneville's disease / Epiloia []
7595.08 Other type of phakomatosis

Hamartoses, not elsewhere classified

7596.01 Peutz-Jegher's syndrome

7596.12 Sturge-Weber syndrome []
7596.21 Von Hippel-Lindau syndrome []
7596.31 Gardner's syndrome []
7596.8- Other specified hamartoses
7596.9- Unspecified hamartoses

MAIN SYNDROMES AND MULTIPLE CONGENITAL ANOMALIES NOT ELSEWHERE CLASSIFIED

Malformation affecting facial appearance

7598.01 -
7598.02 Oro-facio-digital S. type I / OFD I / Papillon-Leage S.[]
7598.03 Oro-facio-digital S. type II / OFD II/Mohr syndrome []
7598.06 Robinow syndrome. / Robinow dwarfism / Fetal face syndrome []
7598.07 Whistling face syndrome/ Freeman-Sheldon syndrome/ Cranio-carpo-tarsal syndrome/ Windmill vane hand syndrome[]
7598.08 Williams syndrome

Malformation affecting facial appearance (cont.)

7598.11 Nager syndrome / Acrofacial dysostosis []
7598.12 Ectrodactyly-ectodermal dysplasia-clefting syndrome/ EEC syndrome
7598.13 Trichorhinophalangeal syndrome type I []
7598.14 Roberts syndrome []
7598.15 Cherubism []
7598.16 G-syndrome / Opitz-Frias syndrome / Hypertelorism-hypospadias with dysphagia syndrome []
7598.17 Opitz (H-H) syndrome / Hypertelorism-hypospadias syndrome/ BBB syndrome []
7598.18 Aarskog syndrome / Aarskog Scott syndrome / Facio-digito-genital dysplasia []

Congenital malformation with short stature

7598.21 Cornelia de Lange syndrome / Brachmann-de Lange syndrome []
7598.22 Bloom syndrome []
7598.23 Russell-Silver syndrome / Silver syndrome
7598.24 Dubowitz syndrome []
7598.26 Cockayne syndrome []
7598.27 Seckel syndrome / Bird-headed dwarfism / Nanocephalic dwarfism []
7598.28 Smith-Lemli-Opitz type I syndrome []

Congenital malformation with overgrowth

7598.32 Sotos syndrome / Cerebral gigantism
7598.33 Marshall-Smith syndrome
7598.34 Weaver syndrome

Congenital malformation syndromes involving limbs

7598.41 Rubinstein-Taybi syndrome
7598.42 Holt-Oram syndrome / Cardiac limb syndrome / Heart hand syndrome []
7598.43 Klippel-Trenaunay syndrome []
7598.44 Sirenomelia / Caudal regression syndrome []
7598.45 Oto-palato-digital syndrome / Taybi syndrome []
7598.46 Aglossia adactylia syndrome / Hanhart syndrome / Hypoglossia-hypodactylia syndrome
7598.47 Achondrogenesis type II / Grebe syndrome []
7598.48 Multiple pterygium syndrome / Escobar syndrome []
7598.49 Smith-Lemli-Opitz type II syndrome []

7598 52 Brotes Syndrome

Congenital malformation syndromes with skeletal changes

7598.61 Marfan syndrome [] / Arachnodactyly syndrome
7598.62 Arachnodactyly, contractual, Beals type / Beals Hecht syndrome []
7598.63 Cerebrocostomandibular []
7598.64 COFS / Cerebro-oculo-facio-skeletal syndrome[] / Pena Shokeir II syndrome []
7598.65 Campomelic dysplasia or dwarfism []
7598.66 Coffin-Lowry syndrome[]
7598.67 Coffin-Siris syndrome[]
7598.68 Cranio metaphyseal dysplasia
7598.69 Noonan syndrome []

Congenital malformation syndromes with metabolic disturbances

7598.71 Biemond II syndrome []
7598.72 Prader-Willi syndrome []
7598.73 Menkès syndrome / Kinky hair disease []
7598.74 Beckwith Wiedemann syndrome / Wiedemann-Beckwith syndrome / Exomphalos macroglossia gigantism syndrome/ EMG syndrome []
7598.75 Zellweger syndrome / Cerebro-hepato-renal syndrome []
7598.76 Ataxia-telangiectasia / Louis Barr syndrome []
7598.77 Alport syndrome / Hereditary nephropathy and deafness []

Other syndromes or associations

7598.81 Meckel-Grüber syndrome []
7598.82 Pena-Shokeir Type I syndrome []
7598.84 Angelmann syndrome / Happy Puppet syndrome[]
7598.8- Other specified syndrome or association
7598.95 VATER Association / VACTERL Association
7598.98 Multiple congenital anomalies NEC
7598.99 Multiple congenital anomalies NOS

pas de code de remplissage

Congenital anomaly unspecified

7599.9- Congenital anomaly NOS

XV. CERTAIN CONDITIONS ORIGINATING IN THE PERINATAL PERIOD*

Noxious influences transmitted via placenta

Anticonvulsant drugs
7607.01 Dilantin syndrome
7607.02 Fetal trimethadione
7607.03 Fetal valproate
7607.08 Other fetal anticonvulsant syndrome

Anticoagulant drugs
7607.11 Fetal warfarin syndrome
7607.18 Other anticoagulant syndrome

Antibiotic and chemotherapeutic agents
7607.21 Aminopterin embryopathy
7607.28 Other chemotherapeutic embryopathy

Uterine depressant drugs
7607.38 Other uterine depressant drugs

Hypoglycaemic agents
7607.41 Infant of diabetic mother

Endocrine agents
7607.58 Other endocrine agents

Alcohol
7607.61 Fetal alcohol syndrome

7608.01 Fetal diabetes syndrome/Fetal methylmercury
7608.02 Maternal PKU fetal effect
7608.03 Fetal retinoic acid syndrome
7608.88 Other noxious substance transmitted via placenta
7609.-- Unspecified noxious substance transmitted via placenta

Oligohydramnios sequence

7612.01 Oligohydramnios sequence

Placental transfusion syndrome

Twin to twin transfusion (monozygotic twins). Code only if malformations are present.
7623.01 Artery - Artery transfusion / Acardia / Acephaly
7623.02 Artery - venous transfusion
7623.03 Death of the co-twin (vanishing twin)

7628.01 Amniotic band syndrome/Amniotic band disruption complex / Early amniorupture sequence / ADAM complex

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* Use code 771 for congenital infection in chapter XV, exclude code 7602.

Infections specific to the perinatal period

7710.11 Fetal rubella/Rubella embryopathy

7711.0- Fetal cytomegalovirus infection

7712.11 Fetal toxoplasmosis
7712.21 Fetal herpes infection
7712.91 Fetal varicella
7712.31 Fetal listeriosis
7712.4- Congenital tuberculosis
7712.5- Congenital syphillis
7712.8- Other specified congenital infection
7712.9- Other unspecified congenital infection

Other specific anomalies originating in prenatal period

7780.1- Hydrops fetalis not due to isoimmunisation
7780.2- Hydrothorax
7780.3- Ascites

7786.-- Congenital hydrocele

779.82 hypotonia, cong ?

Index to the Coding Manual

The index is intended as a guide to the Coding Manual, which is much more detailed. The index is not intended to be used for coding. If it is used for this purpose, errors will certainly occur.

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2552.4-	17 Alpha hydroxylase defective synthesis
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2552.0-	21 Beta Hydroxylase defective synthesis
2552.2-	3 Beta Hydroxysteroid dehydrogenase defective synthesis
7587.15	47 XXY
7587.13	48 XXXY
7587.14	49 XXXXY
7598.18	Aarskog syndrome
7598.18	Aarskog Scott syndrome
7555.08	Aase syndrome
7567.-	Abdominal wall anomaly
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7623.01	Acephaly
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7564.31	Achondroplasia
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2762.--	Acidaemia metabolic, NEC
2707.2-	Acidaemia pipecolic
7400.2-	Acrania
7560.02	Acrobrachycephaly
7555.0	Acrocephalosyndactyly
7564.44	Acrodysostosis
7598.11	Acrofacial dysostosis
7564.34	Acromesomelic dwarfism
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7628.01	ADAM complex
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7598.77	Alport syndrome
7527.99	Ambiguous genitalia
7553	Amelia limb lower
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7554.01	Amelia limb, unspecified
5205.1-	Amelogenesis imperfecta
270	Amino-acid transport and metabolism, disorders
7607.21	Aminopterin embryopathy
7628.01	Amniotic band disruption complex
7553.-	Amputation limb lower
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7555.	Arm, other anomalies
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7461.11	Atresia of tricuspid valve
7532.1-	Atresia of ureter
7532.1-	Atresia of ureteropelvic junction
7536.2-	Atresia of urethra
7536.3-	Atresia of urinary meatus

7524.1	Atresia of vagina
7528.31	Atresia of vas deferens
7590.	Atrial isomerism
7456.0	Atrial septal defect, ostium primum
7455.-	Atrial septal defect, other
426.--	Atrioventricular block
7456.-	Atrioventricular septal defects
7428.3-	Atrophy optic
2794.--	Autoimmune disease, NEC
7474.82	Azygos continuation of inferior vena cava
7486.24	Azygos lobe
7514.2-	Bands of omentum and peritoneum, congenital
7442.2-	Bat ear
7560.05	Bathmocephaly
3627.1-	Batten's hereditary retinal dystrophy
7598.17	BBB syndrome
7574.1-	Beaded hair
7540.23	Beak like nose
7598.62	Beals Hecht syndrome
3591.1-	Becker dystrophy
7598.74	Beckwith Wiedemann
7548.13	Bell shaped chest
7540.22	Bent nose, deviated nose
7523.82	Bicornuate uterus
7464.2-	Bicuspid aortic valve
7598.71	Biemond syndrome
7563.62	Bifid sternum
7550.18	Bifid thumb
7534.1-	Bifid ureter
7516.-	Biliary tract anomalies
7598.27	Bird-headed dwarfism
7573.81	Birthmarks NOS
7535.01	Bladder, exstrophy
7538.88	Bladder, other specified anomaly
7539.29	Bladder, unspecified anomaly
7436.02	Blepharophimosis-ptosis syndrome
7436.01	Blepharoptosis
7573.51	Bloch-Sulzberger syndrome
7598.22	Bloom syndrome
7434.5-	Blue sclera
7586.2-	Bonnevie Ullrich's syndrome
7444.21	BOR syndrome
7595.01	Bourneville's disease
7544.2-	Bow legs, NOS
7544.01	Bowing, femur
7544.13	Bowing, tibia / fibula
7540.61	Brachycephaly
7555.8	Brachydactyly
7598.21	Brachmann-de Lange syndrome
7424.-	Brain anomalies
7429.01	Brain, unspecified anomaly
7422.9-	Brain, unspecified reduction deformities

2703.- Branched-chain amino-acid metabolism, disturbance
7444.- Branchial cleft anomalies (first arch)
7444.21 Branchio-oto-renal syndrome
7576.- Breast anomalies
7574.5- Brittle hair
7521. Broad ligament anomalies
7486.1- Bronchiectasis, congenital
7483.51 Bronchomalacia, isolated
7483. Bronchus anomalies
426.-- Bundle branch block
7432.01 Buphthalmos
7560.04 C syndrome
7565.31 Caffey's syndrome
7555.06 Camptodactyly
7598.65 Campomelic dysplasia
7598.65 Campomelic dwarfism
2719.-- Carbohydrate transport and metabolism, unspecified disorders
425.-- Cardiomyopathy
7531.12 Carolli disease
7555.02 Carpenter syndrome
7564.51 Cartilage-hair hypoplasia syndrome
7433.2- Cataract
7425.- Cauda equina anomalies
7598.44 Caudal regression syndrome
7540.02 Cayler syndrome
7422.63 Cebocephaly
3590.0- Central core disease
7420.9 Cephalocele
7422.3 Cerebellum, anomalies
7598.32 Cerebral gigantism
7598.75 Cerebro-hepato-renal syndrome
7598.64 Cerebro-oculo-facio-skeletal syndrome
7598.63 Cerebrocostomandibular syndrome
7561.4- Cervical vertebrae, anomaly ????
7561.12 Cervico-oculo-acoustic syndrome
7524. Cervix, anomalies
7480.21 CHARGE association
7598.15 Cherubism
7480. Choanal atresia
7564.2- Chondrodysplasia with haemangioma
7565.7 Chondrodysplasia punctata
7564.- Chondrodystrophy
7526.21 Chordee, congenital
7435.3- Choroid, specified anomaly
2861.-- Christmas disease
758 Chromosome anomalies
7580. Down syndrome
7581. Patau's sundrome
7582. Edward's syndrome
7583 Autosomal deletions
7584. Autosomal translocations, balanced
7585. Autosomal anomalies, other

7586.	X chromosome aberrations in absence of Y
7587.	Klinefelter's syndrome
7588.	Sex chromosome anomalies, other
7589.	Unspecified chromosome anomalies
7547.1-	Claw foot
7498.5	Cleft, facial, unusual
7491.	Cleft lip
7492.	Cleft lip and palate
7490.	Cleft palate
7563.63	Cleft sternum
7555.51	Cleidocranial dysostosis / dysplasia
7543.2-	Clicking hip
7555.07	Clinodactyly
7524.5	Clitoris, anomalies
7560.07	Cloverleaf skull
7548.42	Club fingers
7547.31	Club foot NOS
7546.	Club foot valgus
7545.-	Club foot varus
7548.41	Club hand
7575.4-	Club nail(s)
7471.-	Coarctation of aorta
7598.26	Cockayne syndrome
7598.66	Coffin-Lowry syndrome
7598.67	Coffin-Siris syndrome
7598.64	COFS syndrome
7571.11	Collodion baby
7434.3-	Coloboma iris
7433.4-	Coloboma lens
7435.2-	Coloboma optic disc
2792.-	Combined immune deficiency
7456.3-	Common atrioventricular canal
7456.2-	Common atrioventricular canal type VSD
7540.1-	Compression of face
426-.	Conduction disorders, cardiac
5268.--	Condylar hyperplasia, unilateral
7594.	Conjoined twins
7568.	Connective tissue, specified anomalies
7569.41	Connective tissue, unspecified anomaly
7565.71	Conradi Hünermann syndrome
7598.61	Contractural arachnodactyly
7541.2-	Contracture of sternocleidomastoid muscle, congenital
7558.	Contracture of toe(s) and finger(s)
7457.--	Cor biloculare
7468.2-	Cor triatriatum
7453.37	Cor triloculare biatriatum
7457.	Cor triloculare biventricular
7432.21	Cornea enlarged
7434.01	Cornea, opacities
7434.1-	Corneal anomaly, other
7598.21	Cornelia de Lange syndrome
7422.1	Corpus callosum, anomalies

7512.4	Covered anus
7556.61	Coxa valga
7556.62	Coxa vara
7598.68	Cranio metaphyseal dysplasia syndrome dominant
7598.68	Cranio metaphyseal dysplasia syndrome recessive
7598.07	Cranio-carpo-tarsal syndrome
7401.-	Craniorachischisis
7560.08	Craniosynostosis, NOS
7583.1	Cri du chat syndrome
2774.0-	Crigler-Najjar syndrome
7560.11	Crouzon disease
7430.2-	Cryptophthalmos
7430.22	Cryptophthalmos Fraser Syndrome
7525.	Cryptorchidism
7555.41	Cubitus valgus
7542.2-	Curvature postural of spine, congenital, NOS
2550.--	Cushing's syndrome
2880.4-	Cyclical neutropenia
7422.65	Cyclopia (fused eyes)
7424.3-	Cyst arachnoid
7531.81	Cyst Bowman's capsule
7424.	Cyst, brain
7444.04	Cyst branchial associated with 1st branchial arch
7502.81	Cyst branchial associated with 2nd branchial arch
7502.82	Cyst branchial associated with 3rd branchial arch
7502.83	Cyst branchial associated with 4th branchial arch
7424.	Cyst cerebral
7488.11	Cyst bronchogenic of the mediastinum
7516.6-	Cyst choledochal
7521.21	Cyst cimbrial
7531.7-	Cyst cortical
7528.71	Cyst embryonal remains in male
7521.12	Cyst Gartner's duct
7516.11	Cyst intrahepatic congenital
7531.	Cyst kidney
7483.02	Cyst larynx, congenital
7484.0-	Cyst lung
7488.12	Cyst mediastinum, other congenital
7514.12	Cyst mesenteric
7521.23	Cyst mesenteric remnant
7502.93	Cyst mouth, congenital
7531.81	Cyst of the kidney, glomerular
7510.3-	Cyst omphalomesenteric
7520.3-	Cyst ovary isolated
7517.41	Cyst pancreatic isolated
7521.22	Cyst para-ovarian
7528.57	Cyst penis
7524.71	Cyst perineal
7424.1-	Cyst porencephalic (not due to neonatal event)
7444.1-	Cyst preauricular
2538.0-	Cyst Rathke's pouch
7531.	Cyst renal

7592.22	Cyst thyroglossal
7537.1-	Cyst urachus
7531.	Cystic disease of the kidney
2770.--	Cystic fibrosis of pancreas
7573.8	Cystic hygroma
7484.1-	Cystic lung
2700.0-	Cystinosis
2700.1-	Cystinuria
7517.42	Cystosis pancreas, congenital
7442.6-	Darwin's tubercle
2866.--	Defibrination syndrome
7583.	Deletion, autosomal
7586.13	Deletion X (long arms)
7586.12	Deletion X (short arms)
7587.22	Deletion Y (long arms)
7587.21	Deletion Y (short arms)
7572.-	Dermatoglyphics, anomalies
7540.21	Deviation of nasal septum, congenital
7451.0	Dextro transposition
7468.0-	Dextrocardia, isolated
7593.	Dextrocardia, other
2791.0-	Di George sequence
2535.-	Diabetes insipidus
250.--	Diabetes mellitus
7566.-	Diaphragm anomalies
7564.72	Diaphyseal aclasia
7565.51	Diaphyseal dysplasia, progressive
7567.03	Diastasis recti
7502.94	Diastema laryngo-tracheo-oesophageal
7425.2-	Diastematomyelia
7564.41	Diastrophic dwarfism / dysplasia
7522.11	Didelphic uterus with double cervix and vagina
7607.01	Dilantin syndrome
7504.0-	Dilatation of oesophagus, congenital
2713.6-	Disaccharidase deficiencies
2713.-	Disaccharide intolerance
7548.3-	Dislocation of elbow, congenital
7543.0	Dislocation of hip, congenital
7544.4-	Dislocation of knee, congenital
7501.3-	Dislocation of tongue
7538.2-	Diverticulum of bladder
7515.05	Diverticulum of intestine, congenital
7468.93	Diverticulum of left ventricle, congenital
7504.2-	Diverticulum of oesophagus
7502.32	Diverticulum of parotid duct system
7507.41	Diverticulum of stomach
7483.33	Diverticulum of trachea
7540.3-	Dolichocephaly
7560.05	Dolichocephaly due to craniostenosis
7545.1-	Dorsiflexed foot
7472.51	Double aortic arch
7453.3	Double inlet left ventricle

7453.3	Double inlet right ventricle
7533.1-	Double kidney and pelvis
7451.11	Double outlet right ventricle
7528.56	Double penis
7538.4-	Double urethra
7538.4-	Double urinary meatus
7522.11	Double uterus
7580.	Down syndrome
2774.1-	Dubin-Johnson syndrome
7598.24	Dubowitz syndrome
3591.0-	Duchenne dystrophy
7511.04	Diaphragma duodenal
7518.1-	Duplication of alimentary tract
7515.01	Duplication of anus
7515.02	Duplication of appendix
7515.03	Duplication of caecum
7515.03	Duplication of colon
7516.41	Duplication of gallbladder
7504.3-	Duplication of oesophagus
7515.04	Duplication of small intestine
7507.5-	Duplication of stomach
7483.34	Duplication of trachea
7534.1-	Duplication of ureter
7483.63	Duplication of vocal cords
7564.31	Dwarfism, achondroplastic
2598.9-	Dwarfism, NOS
7428.1-	Dysautonomia familial
7564.1-	Dyschondroplasia
7565.81	Dyschondrosteosis
2790.5-	Dysimmunoglobulinaemia
7560.11	Dysostosis craniofacial
7555.31	Dysostosis radio-ulnar
7431.01	Dysplasia of eye
7530.	Dysplasia of kidney
7485.-	Dysplasia of lung
7520.81	Dysplasia of ovary
7425.1-	Dysplasia of spinal cord
7461.16	Dysplasia of tricuspid valve
7530.	Dysplasia renal
427.--	Dysrrhythmias, cardiac
7564.32	Dyssegmental dwarfism / dysplasia
359-.	Dystrophy muscular
744-.	Ear, congenital anomalies
7628.01	Early amniorupture sequence
7536.4-	Early urethral obstruction sequence
7462.--	Ebstein's anomaly
7502.32	Ectasia of parotid duct system
7573.4	Ectodermal dysplasia
7468.81	Ectopia cordis
7591.3-	Ectopic adrenal gland
7515.33	Ectopic anus
7538.1-	Ectopic bladder

7518.2-	Ectopic digestive organs NOS
7468.81	Ectopic heart
7533.3-	Ectopic kidney
7576.5-	Ectopic nipple
7517.3-	Ectopic pancreas
7434.42	Ectopic pupil
7590.5-	Ectopic spleen
7525.3	Ectopic testicle
7486.02	Ectopic tissues in lung
7534.2-	Ectopic ureter
7538.5-	Ectopic urethra
7553.	Ectrodactyly lower limb
7552.	Ectrodactyly of upper limb
7598.12	Ectrodactyly-ectodermal dysplasia-clefting (EEC) syndrome
7535.1-	Ectrophia splanchnica
7436.1-	Ectropion
7582.	Edward's syndrome
7598.12	EEC syndrome
7568.5	Ehlers-Danlos syndrome
2821.--	Elliptocytosis hereditary
7565.21	Ellis van Creveld syndrome
7607.28	Embryopathy chemotherapeutic other
7598.74	EMG syndrome
7484.22	Emphysema lobar, congenital
7420.1	Encephalocele
7564.1-	Enchondromatosis
4253.--	Endocardial fibroelastosis
7592.-	Endocrine glands anomalies
4250.--	Endomyocardial fibrosis
7565.51	Engelmann's syndrome
7507.42	Enterogenous cyst of the stomach
2778.0-	Enterokinase deficiency
7436.2-	Entropion
2778.3-	Eosinophilic granuloma
7436.32	Epicanthic folds
7573.3	Epidermolysis bullosa
7595.01	Epiloia
7526.11	Epispadias
7598.48	Escobar syndrome
2708.0-	Ethanolaminuria
7422.64	Ethmocephaly
7442.5-	Eustachian tube, anomaly
7566.2-	Eventration of diaphragm
7567.01	Exomphalos
7598.74	Exomphalos macroglossia gigantism syndrome
7564.71	Exostosis
7535.1-	Exstrophy of cloaca sequence
7535.01	Exstrophy of urinary bladder sequence
7535.03	Extroversion of bladder
743.--	Eye, congenital anomalies
7449.19	Face, congenital anomaly NOS
7598.18	Facio-digito-genital dysplasia syndrome

7560.61	Facio-auriculo-vertebral first and second arch syndrome
2860.--	Factor VIII disorder, congenital
2861.--	Factor IX disorder, congenital
2862.--	Factor XI disorder, congenital
7521.	Fallopian tubes, anomalies
7452.1-	Fallot's pentalogy
7452.0-	Fallot's tetralogy
7468.4-	Fallot's trilogy
2700.2-	Fanconi (-de Toni) (-Debre) syndrome
7568.	Fascia, anomalies
2822.2-	Favism
7558.01	Fetal akinesia sequence
7607.61	Fetal alcohol syndrome
7607.0	Fetal anticonvulsant syndrome
7711.0-	Fetal cytomegalovirus infection
7608.01	Fetal diabetes syndrome
7712.21	Fetal herpes infection
7712.31	Fetal listeriosis
7608.01	Fetal methylmercury syndrome
7608.03	Fetal retinoic acid syndrome
7710.11	Fetal rubella
7712.11	Fetal toxoplasmosis
7607.02	Fetal trimethadione
7586.92	Fetal Turner's syndrome
7607.03	Fetal valproate syndrome
7712.91	Fetal varicella
7607.11	Fetal warfarin syndrome
4273.1-	Fibrillation atrial
4274.1-	Fibrillation ventricular
7564.03	Fibrochondrogenesis
7550.15	Finger like thumb
7550.05	Finger, long
7502.94	Fissure laryngo-tracheo-oesophageal
7535.2-	Fissure vesical, superior
7515.43	Fistula anovestibular (female)
7444.03	Fistula branchial Internal associated with first branchial arch
7502.84	Fistula branchial, 2nd branchial arch
7502.85	Fistula branchial, 3rd branchial arch
7502.86	Fistula branchial, 4th branchial arch
7503.3	Fistula broncho-oesophageal
7523.23	Fistula of uterus with digestive tract
7523.21	Fistula of uterus with intestine
7523.22	Fistula of uterus with rectum
7523.24	Fistula of uterus with urinary bladder
7524.21	Fistula rectovaginal, congenital
7538.61	Fistula rectovesical
7503.21	Fistula tracheo-oesophageal without oesophageal atresia (Gross type E)
7516.71	Fistula tracheobiliary
7538.7-	Fistula urethra, NOS
7523.23	Fistula utero intestinal
7523.24	Fistula utero-vesical
7502.6-	Fistula lip

7555.06	Flexion deformities of finger
4273.2-	Flutter atrial
4274.2-	Flutter ventricular
7455.0-	Foramen ovale patent
7502.92	Fordyce disease (mouth)
7560.41	Franceschetti syndrome
7560.51	François dyscephalic syndrome
7598.07	Freeman-Sheldon syndrome
7574.73	Frontal upsweep
2712.--	Fructose intolerance, hereditary
2718.4-	Fucosidosis
7548.11	Funnel chest, congenital
7598.16	G-syndrome
2711.9-	Galactose metabolism, unspecified disorders
2711.0-	Galactose-l-phosphate uridyl transferase deficiency
7516.42	Gallbladder, abnormal position
7516.3	Gallbladder, agenesis or hypoplasia
7516.4	Gallbladder, other anomalies
7596.31	Gardner's syndrome
7521.13	Gartner's duct anomaly
7567.11	Gastroschisis
7529.99	Genital organs, unspecified anomaly
752	Genital tract
7544.3-	Genu recurvatum
7556.41	Genu valgum
7556.42	Genu varum
7454.2-	Gerbode defect
7504.0-	Giant oesophagus
7561.23	Gibbus
2530.	Gigantism
2774.2-	Gilbert's syndrome
2871.1-	Glanzmann's syndrome
7432.11	Glaucoma, congenital
7533.61	Glomerular disease, congenital
2713.2-	Glucose intolerance
2822.0-	Glucose-6-phosphate dehydrogenase deficiency anaemia
2713.3-	Glucose-galactose intolerance
2710.-	Glycogenosis
2439.1-	Goitrous cretin
7560.61	Goldenhar syndrome
7527.41	Gonadal dysgenesis
7474.	Great veins, anomaly
7598.47	Grebe syndrome.
7527.51	Gynandrism
7573.8	Haemangioma of skin
7590.82	Haemangioma of spleen
7483.05	Haemangioma, subglottic
2750.2-	Haemochromatosis
2827.-	Haemoglobinopathies, other than thalassaemia and sickle-cell
2750.0-	Haemosiderosis, primary
7574.-	Hair, anomaly
7560.51	Hallerman-Streiff syndrome

7556.01	Hallux valgus
7556.02	Hallux varus
7486.01	Hamartomas of lung
7596.-	Hamartoses
7598.42	Hand heart syndrome
2778.7-	Hand-Schuller-Christian disease
7598.46	Hanhart syndrome
7598.84	Happy Puppet syndrome
7571.0-	Harlequin fetus
2700.3-	Hartnup disease
2827.0-	Hb-C disease
2827.1-	Hb-D disease
2827.2-	Hb-E disease
2827.4-	Hb-H disease
7468.87	Heart block, congenital
7598.42	Heart-limb syndrome
2827.5-	Heinz body anaemia, congenital
7400.3-	Hemianencephaly
7400.3-	Hemicrania
7566.1	Hemidiaphragmatic defect
7560.61	Hemifacial microsomia
7553.3	Hemimelia of lower limb
7552.3	Hemimelia of upper limb
3431.--	Hemiplegia, congenital
7561.	Hemivertebrae
7516.	Hepatic ducts, anomaly
7516.13	Hepatic fibrosis, congenital
7527.01	Hermaphroditism true
7538.2-	Hernia bladder
7566.1	Hernia diaphragmatic
5530.--	Hernia femoral
550-.--	Hernia inguinal
7566.14	Hernia through the foramen of Bochdalek
7566.13	Hernia through the foramen of Morgagni
5531.--	Hernia umbilical
7506.01	Hiatus hernia, congenital
7573.43	Hidrotic ectodermal dysplasia
7502.4-	High arched palate
7513.-	Hirschsprung's disease
2705.-	Histidine metabolism disturbance
2778.-	Histiocytosis X
7400.4-	Holoacrania
7400.4-	Holoanencephaly
7422.6	Holoprosencephaly
7598.42	Holt-Oram syndrome
2704.0-	Homocystinuria
7484.21	Honeycomb lung
7533.2-	Horseshoe kidney
7521.21	Hydatid of Morgagni
7424.4-	Hydranencephaly
7532.21	Hydro-ureter
7786.--	Hydrocele, congenital

7423.1	Hydrocephaly
7524.3	Hydrometrocolpos
7425.4-	Hydromyelia
7532.01	Hydronephrosis, congenital
7432.12	Hydrophthalmos
7780.1-	Hydrops fetalis not due to isoimmunization
7425.4-	Hydrorachis
7780.2-	Hydrothorax
2708.1-	Hydroxyprolinaemia
2720.1-	Hyper-beta-lipoproteinaemia
2706.1-	Hyperammonaemia
2774.-	Hyperbilirubinaemia congenital
2754.-	Hypercalcaemia / hypercalciuria
2720.-	Hypercholesterolaemia
2723.--	Hyperchylomicronaemia
2721.--	Hyperglyceridaemia pure
2707.0-	Hyperglycinaemia
251.--	Hyperinsulinism
272.--	Hyperlipidaemia
2720.3-	Hyperlipoproteinaemia low-density-lipoid-type (LDL)
2752.2-	Hypermagnesaemia
7565.31	Hyperostosis cortical infantile
2520.--	Hyperparathyroidism
2701.--	Hyperphenylalaninaemia
2753.4-	Hyperphosphatasia
7590.2-	Hyperplasia spleen
7533.4-	Hyperplastic kidney
2708.2-	Hyperprolinaemia
7560.21	Hypertelorism
7598.17	Hypertelorism-hypospadias
7598.16	Hypertelorism-hypospadias with dysplasia
7592.14	Hyperthyroidism congenital
2702.1-	Hypertyrosinaemia
2554.3-	Hypoaldosteronism
2754.-	Hypocalcaemia
7564.02	Hypochondrogenesis
7564.33	Hypochondroplasia
2790.4-	Hypogammaglobulinaemia
7598.46	Hypoglosso-hypodactylia syndrome
2512.--	Hypoglycaemia, unspecified
7573.42	Hypohidrotic ectodermal dysplasia
2752.-	Hypomagnesaemia
7573.52	Hypomelanosis of Ito
2521.--	Hypoparathyroidism
2753.3-	Hypophosphataemia
2753.-	Hypophosphatasia
7567.02	Hypoplasia abdominal muscle
7591.1-	Hypoplasia adrenal gland
7472.1-	Hypoplasia aorta (tubular Hypoplasia)
7422.32	Hypoplasia cerebellum
7517.04	Hypoplasia exocrine pancreas
7431.01	Hypoplasia eye

7516.3-	Hypoplasia gallbladder
7516.52	Hypoplasia intrahepatic bile ducts, congenital
7530.	Hypoplasia kidney
7561.61	Hypoplasia lumbar vertebrae
7485.1-	Hypoplasia lung
7425.1-	Hypoplasia spinal cord.
7502.31	Hypoplasia parotid
7592.02	Hypoplasia pituitary gland
2840.2-	Hypoplasia red-cell, constitutional
7530.	Hypoplasia renal
7561.72	Hypoplasia sacrococcygeal
7528.12	Hypoplasia scrotum / testis
7590.1-	Hypoplasia spleen
7561.52	Hypoplasia thoracic vertebrae
7592.42	Hypoplasia thymus
7501.1-	Hypoplasia tongue
7475.--	Hypoplasia umbilical artery
7576.1-	Hypoplastic breast with Hypoplastic nipple
7467.-	Hypoplastic left heart syndrome
7572.3-	Hypoplastic palmar creases
7528.53	Hypoplastic penis
7526.	Hypospadias
7560.22	Hypotelorism
7422.21	Hypothalamus anomaly
7592.11	Hypothyroidism congenital
7574.01	Hypotrichosis congenital
7571.-	Ichthyosis congenita
279.--	Immunodeficiency, unspecified
7512.45	Imperforate anal membrane
7524.03	Imperforate cervix
7524.31	Imperforate hymen
7573.51	Incontinentia pigmenti
7573.52	Incontinentia pigmenti achromions
7527.	Indeterminate sex
7402.-	Iniencephaly
7440.3	Inner ear, congenital anomaly
4268.2-	Interference dissociation conduction disorder
7472.0	Interruption of aorta
7514.-	Intestinal fixation anomaly
7424.6-	Intracranial calcification
7434.6-	Iridogoniodysgenesis with somatic anomaly
7434.	Iris anomaly
	Isochromosome - see Chromosomes
7590.01	Ivemark association
7565.62	Jarcho Levin syndrome
7428.01	Jaw-winking phenomenon
4268.3-	Jervell-Lange-Neilsen syndrome
7564.01	Jeune disease
7481.81	Johnson Blizzard syndrome
7593.41	Kartagener's syndrome (triad)
7564.2-	Kast syndrome
7524.32	Kaufman Mc Kusick syndrome

7432.22	Keratoglobus, congenital
7571.0-	Keratum malignum
7533.1-	Kidney and pelvis, double / triple
7533.3	Kidney ectopic
7533.2-	Kidney fused / lobulated
7598.73	Kinky hair
7560.07	Kleeblattschadel
7587.-	Klinefelter's syndrome
7561.11	Klippel-Feil sequence / syndrome
7598.43	Klippel-Trenaunay syndrome
7564.43	Kniest dysplasia
7575.2-	Koilonychia, congenital
3351.--	Kugelberg-Welander disease
7561.22	Kyphoscoliosis
7561.21	Kyphosis
7440.31	Labyrinth membranous, congenital anomaly
7436.-	Lacrimal apparatus anomaly
2713.-	Lactose intolerance
4268.4-	Lange-Neilson syndrome
7481.83	Langer-Geidion syndrome
7567.11	Laparoschisis
7558.11	Larsen's syndrome
7482.	Larynx, web
7483.	Larynx, other anomalies
3627.2-	Leber's dystrophy
7553.	Leg, reduction deformities
7556.	Leg, other anomalies
7433.-	Lens anomaly
7431.02	Lenz microphthalmos
2598.2-	Leprechaunism
7560.05	Leptocephaly
2703.0-	Leucinosis
282.--	Leucocytes diseases
7575.3-	Leukonychia, congenital
7468.1-	Levocardia
7451.2	Levotransposition
3591.3-	Limb-girdle dystrophy
7502.-	Lip anomaly
2728.0-	Lipase deficiency
2727.--	Lipidoses
2726.--	Lipodystrophy
3627.3-	Lipofuscinosis NOS
2728.2-	Lipomatosis
7517.04	Lipomatosis pancreas
2725.--	Lipoprotein deficiencies
7422.41	Lissencephaly syndrome
7516.-	Liver anomaly
7556.	Lobster claw foot
7555.1	Lobster claw hand
7565.01	Lobstein disease
7598.76	Louis Bar syndrome
7553.	Lower limb, reduction deformities

744242 low set ears

7556.	Lower limb, other anomalies
4268.5-	Lown-Ganong-Levine syndrome
7486.-	Lung anomaly
7455.2-	Lutembacher's syndrome
7573.86	Lymphangioma
7483.06	Lymphangioma subglottic, congenital
7448.2-	Macrocheilia
7513.4-	Macrocolon (not aganglionic), congenital
7555.05	Macrodactylia
2733.--	Macroglobulibaemia
7501.2-	Macroglossia
7448.01	Macrostomia
7442.01	Macrotia
7555.22	Madelung's deformity
7564.2-	Maffuci syndrome
7540.64	Mandibular asymmetry
5240.-	Mandibular hyperplasia / hypoplasia
7560.41	Mandibulofacial dysostosis
2775.0-	Mannosidosis
2703.1-	Maple-syrup-urine disease
7565.41	Marble bones
7428.01	Marcus-Gunn phenomenon
7565.63	Marden Walker syndrome
7598.61	Marfan syndrome
7481.82	Marshall syndrome
7598.33	Marshall-Smith syndrome
7573.2-	Mast cell disease
7573.2-	Mastocytosis
2710.4-	McArdle's syndrome
7510.2-	Meckel bridle isolated
7510.1-	Meckel's diverticulum
7598.81	Meckel-Grüber syndrome
7531.	Medullary cystic disease of the kidney
7513.4-	Megacolon (without aganglionosis), congenital
7432.21	Megalocornea
7507.1-	Megalogastria
7532.21	Megaureter
7570.2-	Meige disease
7432.13	Melnick Needles osteodysplasty
7444.21	Melnick-Fraser syndrome
7420.	Meningocele cranial
7411.-	Meningocele spinal
	Meningomyelocele, see Myelomeningocele
7598.73	Menkès syndrome
7400.3-	Meroacrania
7400.3-	Meroanencephaly
7514.1	Mesentery, anomaly
7565.81	Mesomelic dwarfism
7564.35	Mesomelic dysplasia
7564.52	Metaphyseal chondrodysplasia Mc Kusick type
7545.22	Metatarsus adductus
7546.03	Metatarsus valgus

7545.21	Metatarsus varus
7564.42	Metatropic dwarfism / dysplasia
2897.0-	Methaemoglobinaemia congenital
7528.53	Micropenis
7421.11	Microcephalic primordial dwarfism syndrome
7421.01	Microcephaly
7448.3-	Microcheilia
7515.2-	Microcolon
7507.01	Microgastria
7501.1-	Microglossia
5240.5-	Micrognathism
7422.5-	Microgyria
7555.81	Micromelia arm
7556.81	Micromelia leg
7431.01	Microphthalmos
7422.5-	Micropolygyria
7448.1-	Microstomia
7442.1-	Microtia
7422.41	Miller-Dieker syndrome
7451.3-	Miror image atrial arrangement (with asplenia or polysplenia)
7465.1-	Mitral atresia
7466.1-	Mitral insufficiency (congenital)
7465.2-	Mitral stenosis (congenital)
7428.21	Möbius / Moebius syndrome
7598.03	Mohr syndrome
7573.82	Mongolian blue spot
7436.82	Mongoloid slant
7574.1-	Monilethrix
7528.02	Monorchism
7556.1	Monodactyly, foot
7555.1	Monodactyly, hand
7502.9	Mouth anomaly
2775.	Mucopolysaccharidosis
7531.6	Multicystic kidney
7520.4-	Multicystic ovary
7565.71	Multiple ephiphyseal dysplasia
7524.13	Murcs association
7568.	Muscle, specified anomaly
7569.01	Muscle, unspecified anomaly
7425.1	Myelodysplasia
7412.-	Myelomeningocele
7468.6-	Myocardium malformation
358-	Myoneural disorders
3590.1-	Myopathy myotubular
2710.4-	Myophosphorylase deficiency
3592.-	Myotonic disorders
7573.8	Naevus
7598.11	Nager syndrome
7575.-	Nail, anomaly
7568.31	Nail-patella syndrome
7575.1-	Nail enlarged
7564.41	Nanism

7598.27	Nanocephalic dwarfism
7540.2	Nasal shape anomalies
7481.3-	Nasal sinus wall anomaly
5206.2-	Natal tooth
744	Neck congenital anomaly
3590.2-	Nemaline body disease
5206.3-	Neonatal tooth
7533.62	Nephritis, congenital
2754.0-	Nephrocalcinosis
7531.41	Nephronophthisis, juvenile type
7598.77	Nephropathy and deafness hereditary
7533.63	Nephrosis congenital
7538.81	Neurogenic bladder dysfunction
2880.0-	Neutropenia, congenital
2880.4	Neutropenia, cyclical
7576.4-	Nipple small
7555.04	Noack syndrome
7570.1-	Nonne-Milroy's disease
7598.69	Noonan syndrome
7540.2	Nose, abnormal shape
7481.	Nose anomaly
7428.21	Nuclear agenesis
3631.1-	Nystagmus, congenital
7540.42	Occiput flat
3591.4-	Ocular dystrophy
7560.61	Oculo-auriculo-vertebral syndrome
7560.51	Oculomandibular dysostosis
3591.5-	Oculopharyngeal dystrophy
7570.-	Oedema of legs hereditary
7535.1-	OEIS complex
7503.4	Oesophageal anomalies
7565.	Osteodystrophy
7530.03	Oligohydramnios sequence
7564.1-	Ollier disease
7567.01	Omphalocele
7575.1-	Onychauxis
7575.0-	Onychoatrophy
7434.01	Opacity corneal
7598.17	Opitz (H-H) syndrome
7560.04	Opitz trigonocephaly syndrome
7598.16	Opitz-Frias syndrome
3588.2-	Oppenheim syndrome
7435.4-	Optic disc, specified anomaly
7436.7-	Orbit anomaly
7440.31	Organ of Corti, congenital anomaly
7598.02	Oro-facio-digital syndrome type I
7598.03	Oro-facio-digital syndrome type II
7568.31	Osteo-onychodysplasia hereditary
7564.1-	Osteochondromatosis
7565.9	Osteodystrophies, unspecified
7565.0	Osteogenesis imperfecta
7565.41	Osteopetrosis

1940 Neuroblastoma

7565.61	Osteopoikilosis
7456.0-	Ostium primum atrial septal defect
7455.1-	Ostium secundum atrial septal defect
7598.45	Oto-palato-digital syndrome
7520.-	Ovary, anomaly
7548.6-	Overlapping fingers
7472.6-	Overriding aorta
7521.	Oviduct, anomaly
7527.02	Ovotestis
2718.-	Oxalosis / oxaluria
7560.01	Oxycephaly
7575.1-	Pachyonychia
749	Palate cleft
7572.4-	Palmar furrows
7572.-	Palmar simian crease
7436.32	Palpebral fissure, narrow
7517.2	Pancreas annular
7517.8-	Pancreas, anomaly
2532.0-	Panhypopituitarism idiopathic
7483.03	Papilloma of larynx, congenital
7598.02	Papillon-Leage syndrome
3592.1-	Paramyotonia congenita
272.--	Paraproteinaemia
7526.09	Paraspadia
7592.3	Parathyroid gland anomaly
7581.	Patau's syndrome
7470.--	Patent ductus arteriosus
7537.0-	Patent urachus
7548.0-	Pectus carinatum
7548.11	Pectus excavatum
7598.82	Pena Shokeir type I disease
7598.64	Pena Shokeir type II disease
2439.0-	Pendred's syndrome
7526	Penis anomalies
7481.4-	Perforated nasal septum
7468.5-	Pericardium malformation
2827.3-	Persistent fetal haemoglobin, hereditary
7524.61	Persistent canal of Nuck
7515.5-	Persistent cloaca
7474.83	Persistent left posterior cardinal vein
7474.1-	Persistent left superior vena cava
7510.0-	Persistent omphalomesenteric duct
7472.3-	Persistent right aortic arch
7510.0-	Persistent vitelline duct
7547.02	Pes cavus
7547.01	Pes equinus
7546.1-	Pes planus, congenital
7596.01	Peutz-Jegher's syndrome
7555.04	Pfeiffer syndrome
7502.-	Pharyngeal anomaly
2701.--	Phenylketonuria
7608.02	Phenylketonuria, effect of maternal

7553.1	Phocomelia of lower limb
7552.1	Phocomelia of upper limb
7560.31	Pierre Robin sequence
7548.0-	Pigeon chest
7574.2-	Pili torti
2533.-	Pituitary dwarfism
7592.0	Pituitary gland, anomaly
7560.06	Plagiocephaly due to synostosis
7540.51	Plagiocephaly, oblique head deformation
7572.4-	Plantar furrows
7488.0-	Pleura anomaly
7484.23	Pneumatocele, congenital
7442.33	Pointed ear
7568.01	Poland syndrome
2730.--	Polyclonal hypergamma-globulinaemia
7434.41	Polycoria
7516.12	Polycystic disease of the liver
7531.11	Polycystic kidney of the newborn
7531.	Polycystic kidney disease
7484.1-	Polycystic lung
2896.--	Polycythaemia familial
7550.	Polydactyly
7422.5-	Polygyria
7528.21	Polyorchism
7565.11	Polyostotic fibrous dysplasia
7441.01	Polyotia
7590.42	Polysplenia syndrome
7550.4	Polysyndactyly fingers
2710.1-	Pompe's disease
7568.82	Popliteal pterygium
7568.82	Popliteal web syndrome
7424.1-	Porencephaly (not due to neonatal event)
7474.42	Portal vein - hepatic artery fistula
7435.-	Posterior segment of the eye, anomaly
7530.03	Potter sequence
7531.11	Potter type I, polycystic kidneys
7531.61	Potter type IIa renal dysplasia
7532.62	Potter type IIb renal dysplasia
7531.21	Potter type III, adult polycystic kidney disease
7531.7-	Potter type IV
7540.1-	Potter's facies
7598.72	Prader-Willi syndrome
2552.7-	Precocious puberty with adrenocortical hyperfunction
7543.1	Predislocation / preluxation of hip
3342.--	Primary cerebellar degeneration
2598.0-	Progeria
7538.3-	Prolapsed bladder (mucosa)
7523.13	Prolapsed uterus
7528.4	Prostate, anomaly
7567.21	Prune belly syndrome
7564.52	Pseudo-achondroplasia
7527.	Pseudo-hermaphroditism

2754.2-	Pseudohypoparathyroidism
3588.3-	Pseudoparalysis atonica, congenital
7445.--	Pterygium colli
7598.48	Pterygium multiple
7436.01	Ptosis, congenital
7473.34	Pulmonary arteriovenous aneurysm
7473.3	Pulmonary artery, anomaly
7460.-	Pulmonary valve, anomaly
7474.	Pulmonary veins, anomaly
2775.5-	Pyknodysostosis
7505.1	Pyloric stenosis
7505.8-	Pyloric obstruction, other congenital
7552.68	Radial aplasia-thrombocytopenia (TAR) syndrome
7548.43	Radial deviation of hand
7552.	Radius anomalies
7502.2-	Ranula
7552.	Reduction deformity of arm
7422.-	Reduction deformity of brain
7553.	Reduction deformity of leg
7476.-	Renal artery, anomaly
7533.5-	Renal calculi, congenital
7533.3-	Renal ectopia
748	Respiratory system anomaly
7435.18	Retina, specified anomaly
7435.1-	Retinal aneurysm, congenital
3627.-	Retinal dystrophy hereditary
3627.0-	Retinitis pigmentosa
7532.22	Retrocaval ureter
7528.52	Retroscrotal penis
7563.-	Ribs, anomaly
2753.0-	Rickets vitamin-D-resistant
7540.62	Ridge metopic suture
7434.6-	Rieger's syndrome
7472.53	Right aortic arch with left ligamentum arteriosum
7428.1-	Riley-Day syndrome
	Ring chromosome-see Chromosomes
7598.14	Roberts syndrome
7598.06	Robinow dwarfism
7598.06	Robinow syndrome
7556.11	Rocker bottom foot
7454.0-	Roger's disease
7514.02	Rotation of intestine reversed
2774.3-	Rotor's syndrome
7710.11	Rubella embryopathy
7598.41	Rubinstein-Taybi syndrome
7598.23	Russell-Silver syndrome
7561.7	Sacrococcygeal anomaly
7555.03	Saethre-Chotzen syndrome
7502. *	Salivary glands or ducts, anomaly
7560.05	Scaphocephaly
7568.81	Schwartz-Jampel syndrome
7474.84	Scimitar syndrome

52405 Retrognathism

* 755603 Sandalen lücke

7434.-	Sclera or anterior part of eye, anomaly
7542.	Scoliosis postural
7528.	Scrotum, anomaly
7598.27	Seckel syndrome
7523.82	Septate uterus
7428.4-	Septo-optic dysplasia
7485.2-	Sequestration of lung tissue
7548.14	Shield chest
7564.05	Short rib syndromes (with or without polydactyly),
2826.--	Sickle-cell anaemia
2824.6-	Sickle-cell thalassaemia
2825.--	Sickle-cell trait
7598.23	Silver syndrome
7456.1- *	Single common atrium
7475.--	Single umbilical artery
7453.3	Single ventricle
4266.--	Sino atrial block
7444.02	Sinus branchial associated with first branchial arch
7444.1-	Sinus preauricular
7598.44	Sirenomelia
7593.	Situs inversus
7507.31	Situs inversus of the stomach isolated
7471.2-	Sjogren-Larsson syndrome
7573.-	Skin anomalies
7560.-	Skull, anomaly
7540	Skull, anomaly
7598.28	Smith-Lemli-Opitz type I syndrome
7598.49	Smith-Lemli-Opitz type II syndrome
7598.32	Sotos syndrome
7548.5-	Spade-like hand
7433.1-	Spherical lens
2820.--	Spherocytosis, hereditary
7433.1-	Spherophakia
7419.-	Spina bifida aperta
7561.01	Spina bifida occulta uncomplicated
7425.8-	Spinal cord anomaly, other specified
7429.1-	Spinal cord, unspecified anomaly
3351.--	Spinal muscular atrophy
7561.-	Spine anomaly
334.--	Spinocerebellar disease
7590.-	Spleen, anomaly
7556.1	Split foot
7555.1	Split hand
7564.61	Spondylo-epiphyseal dysplasia congenita, autosomal dominant form
7564.62	Spondylo-thoracic dysplasia
7561.3-	Spondylolisthesis, congenital
7565.62	Spondylothoracic dysplasia
7531.11	Sponge kidney of the newborn
7575.2-	Spoon nails
7555.52	Sprengel's deformity
2728.1-	Steatosis
7541.-	Sternocleidomastoid anomaly

7572.--* Simian crease

7563.-	Sternum, anomaly
7560.32	Stickler syndrome
7507.-	Stomach, anomaly
7461.14	Straddling of tricuspid valve
7520.1-	Streak ovary
7440.0-	Stricture auditory canal
7436.5-	Stricture lacrimal duct
7524.02	Stricture of cervix
7532.1-	Stricture ureter or ureteropelvic junction
7596.12	Sturge-Weber syndrome
7468.3	Subaortic stenosis
7502.34	Sublingual glands, anomaly
7543.12	Subluxation of hip
7502.33	Submaxillary glands, anomaly
2713.4-	Sucrose intolerance
2713.5-	Sucrose-isomaltose intolerance
2713.7-	Sucrosuria
7483.65	Sulcus glottitis
2775.8-	Sulphatases deficiency multiple
2704.-	Sulphur-bearing amino-acid metabolism disturbances
7486.2	Supernumerary lung tissue
7483.43	Supernumerary bronchus
7533.0-	Supernumerary kidney
7562.--	Supernumerary cervical rib
7561.74	Supernumerary sacrococcygeal vertebrae
5201.--	Supernumerary teeth
7561.54	Supernumerary thoracic vertebrae
7561.64	Supernumerary lumbar vertebrae
7472.2-	Supraaortic stenosis
7472.2-	Supravalvular aortic stenosis
7551.04	Symphalangism of fingers
7551.	Syndactyly fingers / toes
7551.04	Synostosis of fingers
7551.25	Synostosis of toes
7555.32	Synostosis radio-ulnar
7712.5-	Syphilis, congenital
7425.5-	Syringomyelia
427-	Tachycardia , paroxysmal
7574.3-	Taenzer's hair
7441.2-	Tag auricle
7441.1-	Tag preauricular
7546.02	Talipes calcaneovalgus
7545.1-	Talipes calcaneovarus
7546.01	Talipes equinovalgus
7545.0-	Talipes equinovarus
3627.5-	Tapetoretinal dystrophy
7552.68	TAR syndrome
7564.62	Tarcho Levin syndrome
7451.12	Taussig-Bing anomaly
7598.45	Taybi syndrome
7568.	Tendon, anomaly
7527.12	Testicular feminization syndrome

7528.	Testis, anomaly
	Tétraploidy - see Chromosomes
	Tetrasomy - see Chromosomes
2824.-	Thalassaemia
7564.45	Thanatophoric dysplasia
7506.01	Thoracic stomach, partial
7548.12	Thorax, anomalies
2873.1-	Thrombocytopenic purpura, congenital
2873.2-	Thrombocytopenia with absent radius (TAR) syndrome
7550.1	Thumb anomalies
2792.1-	Thymic dysplasia with immune deficiency
7592.4	Thymus, anomaly
7592.21	Thyroglossal duct anomaly
7592.1	Thyroid gland anomaly
7501.-	Tongue, anomaly
5206.1-	Tooth impacted
5203.--	Tooth, mottled
7483.-	Trachea, anomaly
	Translocation - see Chromosomes
7515.1-	Transposition of appendix, colon and intestine
7451.	Transposition of great vessels
7507.32	Transposition of stomach
7560.41	Treacher-Collins syndrome
7598.13	Trichorhinophalangeal, type I syndrome
7481.83	Trichorhinophalangeal, type II syndrome
7574.6-	Trichorrhexis nodosa
7462.-	Tricuspid valve, anomaly
7560.03	Trigonocephaly
7550.15	Triphalangeal thumb
	Triploidy - see Chromosomes
	Trisomy - see Chromosomes
2778.1-	Trypsinogen deficiency
7712.4-	Tuberculosis, congenital
7595.01	Tuberous sclerosis
7586.-	Turner's syndrome
7560.01	Turricephaly
7594.	Twins, conjoined
7574.2-	Twisted hair
2702.2-	Tyrosinosis
2702.3-	Tyrosinuria
7468.94	Uhl's disease (parchement right ventricle)
7525.	Undescended testicle
7552.	Upper limb, reduction deformities
7555.	Upper limb, other anomalies
7537.-	Urachus, anomaly
7532.21	Ureter dilated
7534.8-	Ureter, other specified anomaly
7534.3-	Ureterocele
7538.	Urethra specified anomaly
7536.01	Urethral valves posterior
7573.2-	Urticaria pigmentosa
7523.8	Uterus, anomaly

7598.95	VACTERL association
7524.8	Vagina, anomaly
7546.-	Valgus deformities of feet
7536.01	Valves, urethral, posterior
7492.91	Van der Woude syndrome
7545.-	Varus deformities of feet
7528.4	Vas deferens, anomaly
7472.	Vascular ring
7598.95	VATER association
7451.2	Ventricular inversion
7454.-	Ventricular septal defect
7561.	Vertebrae anomaly
7532.-	Vesicoureteral reflux
2552.8-	Virilism
7435.01	Vitreous humour, specified anomaly
7483.64	Vocal cord palsy
7514.03	Volvulus due to abnormal rotation or fixation
2710.0-	Von Gierke's syndrome
7596.21	Von Hippel-Lindau syndrome
2750.2-	Von Recklinghausen-Appelbaum disease
2864.--	Von Willebrand's disease
7565.02	Vrolic syndrome
7442.34	Vulcan ear
7524.4	Vulva, anomaly
7590.81	Wandering spleen
7598.34	Weaver syndrome
7482.-	Web of larynx
7551.1-	Webbed fingers
7445.--	Webbed neck
7555.54	Webbed shoulder
7551.3-	Webbed toes
3350.--	Werdnig-Hoffmann disease
2598.3-	Werner's syndrome
7598.07	Whistling face syndrome
288-	White blood cells, anomaly
7574.72	Widow's peak
7598.74	Wiedemann-Beckwith syndrome
7561.12	Wildervanck syndrome
7598.08	Williams syndrome
2751.	Wilson's disease
7598.07	Windmill vane hand syndrome
2791.1-	Wiskott-Aldrich syndrome
7583.2	Wolf-Hirschhorn syndrome
4267.1-	Wolff-Parkinson-White syndrome
7573.60	Xeroderma pigmentosum
7598.75	Zellweger syndrome

ANNEX 2 : List of minor anomalies

Report of cases with the following anomalies are **not to be transmitted** to the EUROCAT Central Registry **unless** occurring in combination with other specified anomalies:

Anomalies of eye
- Stenosis or stricture of lacrimal duct (74365)

Anomalies of ear
- Minor or unspecified anomaly of ear (7443)
- Preauricular appendage, tag or lobule (74411)
- Other appendage, tag or lobule (74412)

Cardiovascular system
- Functional or unspecified cardiac murmur (7852)
- Absence or hypoplasia of umbilical artery, single umbilical artery (7475)
- Patent ductus arteriosous (7470) (in babies <37 weeks or <2.500 gr)

Digestive system
- Tongue tie (7500)

External genitalis
- Undescended testicle (7525) and unspecified ectopic testis (75253)
- Congenital hydrocele or hydrocele of testis (7786)
- Phymosis (605)
- Hypospadias when the meatus lies before the coronary sulcus, glandular or 1st degree hypospadias (75260)

Limbs
- Clicking hip (75432)
- Clubfoot of postural origin (75473)
- Postural or unspecified metatarsus varus or metatarsus adductus (75452)
- Postural or unspecified talipes calcaneovalgus or pes calcaneovalgus (75460)
- Minor or unspecified anomalies of toe such as hallux valgus, hallux varus, or "orteil en marteau" (75560)

Other musculoskeletal anomalies and anomalies of the integument
- Spina bifida occulta uncomplicated (75610)
- Pectus excavatum (75636 or 75481)
- Minor or unsepecified anomaly of nose (74819)
- Minor or unsepecified deformity of face (74491)
- Minor anomaly of nipple (75768)
- Accessory or ectopic nipple (75765)
- Congenital umbilical hernia (5531), inguinal hernia (550), para umbilical (5531), ventral or incisional (5532), hiatus hernia (7506).
- Abnormal palmar crease (7572).
- Skin tag with surface less than 4 cm^2 : skin tag (75731), naevus (75738), angioma (2280), haemangioma (2280), glomus tumor (2280), lymphangioma (2281), birthmark (75738)
- Sacral dimple (7578 or 6851)