Special Needs of Children Less Than 5 Years Old in Ukraine That May Make Them Available for Adoption by Foreigners

Note that this list was current as of September 6, 2021 and does not guarantee that the child *will* be available for adoption, only that the child *may* be available as long as other other criteria have been met.

B16 Acute hepatitis B
B17 Other acute viral hepatitis
B18.0 Chronic viral hepatitis B with delta-agent
B18.1 Chronic viral hepatitis B without delta-agent
B18.2 Chronic viral hepatitis C
B18.8 Other chronic viral hepatitis
B18.9 Chronic viral hepatitis, unspecified
B20 Human immunodeficiency virus [HIV] disease resulting in infectious and parasitic diseases
B21 Human immunodeficiency virus [HIV] disease resulting in malignant neoplasms
B22 Human immunodeficiency virus [HIV] disease resulting in other specified diseases
B23.0 Acute HIV infection syndrome
B23.1 HIV disease resulting in (persistent) generalized lymphadenopathy
B23.2 HIV disease resulting in haematological and immunological abnormalities, not elsewhere classified
B23.8 HIV disease resulting in other specified conditions
C00-C97 Malignant neoplasms
D61.0 Constitutional aplastic anemia
D81.0 Severe combined immunodeficiency [SCID] with reticular dysgenesis
D81.1 Severe combined immunodeficiency [SCID] with low T- and B-cell numbers
D81.2 Severe combined immunodeficiency [SCID] with low or normal B-cell numbers
D81.3 Adenosine deaminase [ADA] deficiency
D81.4 Nezelof syndrome
D81.5 Purine nucleoside phosphorylase [PNP] deficiency
D81.6 Major histocompatibility complex class I deficiency
Bare lymphocyte syndrome
D81.7 Major histocompatibility complex class II deficiency
D81.8 Other combined immunodeficiencies
Biotin-dependent carboxylase deficiency
D81.9 Combined immunodeficiency, unspecified
Severe combined immunodeficiency disorder [SCID] NOS
D82.0 Wiskott-Aldrich syndrome
D82.1 Di George syndrome
E 34.3 Short stature, not elsewhere classified
E70.0 Classical phenylketonuria
E70.1 Other hyperphenylalaninaemias
E70.2 Disorders of tyrosine metabolism
E71.0 Maple-syrup-urine disease
E74.0 Glycogen storage disease

E74.2 Disorders of galactose metabolism
E75.0 GM2 gangliosidosis
E75.1Other gangliosidosis
E75.2 Other sphingolipidosis
E75.3 Sphingolipidosis, unspecified
E75.4 Neuronal ceroid lipofuscinosis
E75.5 Other lipid storage disorders
E75.6 Lipid storage disorder, unspecified
E76.0 Mucopolysaccharidosis, type I
E76.1Mucopolysaccharidosis, type II
E76.2 Other mucopolysaccharidoses
E76.3 Mucopolysaccharidosis, unspecified
E76.8 Other disorders of glucosaminoglycan metabolism
E76.9 Disorder of glucosaminoglycan metabolism, unspecified
E77.0 Defects in post-translational modification of lysosomal enzymes
E83.0 Disorders of copper metabolism
E83.3 Disorders of phosphorus metabolism and phosphatases
F72 Severe mental retardation
G09 Sequelae of inflammatory diseases of central nervous system
G40.2 Localization-related (focal)(partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures
G40.3 Generalized idiopathic epilepsy and epileptic syndromes
G40.4 Other generalized epilepsy and epileptic syndromes
G40.5 Special epileptic syndromes
G80.0 Spastic quadriplegic cerebral palsy
G80.1 Spastic diplegic cerebral palsy
G80.2 Spastic hemiplegic cerebral palsy
G80.3 Dyskinetic cerebral palsy
G80.4 Ataxic cerebral palsy
G80.8O ther cerebral palsy
G80.9 Cerebral palsy, unspecified
G82.0 Flaccid paraplegia
G82.1 Spastic paraplegia
G82.2 Paraplegia, unspecified
G82.3 Flaccid tetraplegia
G82.4 Spastic tetraplegia
G82.5 Tetraplegia, unspecified
H54.0 Blindness, in both eyes
H54.1 Blindness, one eye, low vision other eye
H54.3 Unqualified visual loss, both eyes
H54.4 Blindness, one eye
H54.6 Unqualified visual loss, one eye
H90.0 Conductive hearing loss, bilateral
H90.1 Conductive hearing loss, unilateral with unrestricted hearing on the contralateral side
H90.2 Conductive hearing loss, unspecified
H90.3 Sensorineural hearing loss, bilateral

H90.4 Sensorineural hearing loss, unilateral with unrestricted hearing on the contralateral side
H90.5 Sensorineural hearing loss, unspecified
H90.6 Mixed conductive and sensorineural hearing loss, bilateral
H90.7 Mixed conductive and sensorineural hearing loss, unilateral with unrestricted hearing on the contralateral side
H90.8 Mixed conductive and sensorineural hearing loss, unspecified
transient ischaemic deafness (H93.0)
H91.0 Ototoxic hearing loss
H91.1 Presbycusis
H91.2 Sudden idiopathic hearing loss
H91.3 Deaf mutism, not elsewhere classified
H91.8 Other specified hearing loss
H91.9 Hearing loss, unspecified
I27.0 Primary pulmonary hypertension
I42.0 Dilated cardiomyopathy
I42.1Obstructive hypertrophic cardiomyopathy
I42.2Other hypertrophic cardiomyopathy
I42.5 Other restrictive cardiomyopathy
L40.0 Psoriasis vulgaris
L40.1 Generalized pustular psoriasis
L40.2 Acrodermatitis continua
L40.3 Pustulosis palmaris et plantaris
L40.4 Guttate psoriasis
L40.5 Arthropathic psoriasis
L40.8 Other psoriasis
L40.9 Psoriasis, unspecified
N18.0 Chronic kidney disease
N18.1Chronic kidney disease, stage 1
N18.2Chronic kidney disease, stage 2
N18.3Chronic kidney disease, stage 3
N18.4Chronic kidney disease, stage 4
N18.5Chronic kidney disease, stage 5
N18.9Chronic kidney disease, unspecified
N25.0 Renal osteodystrophy
N25.1 Nephrogenic diabetes insipidus
N26 Unspecified contracted kidney
N27.1 Small kidney, bilateral
Q01.0 Frontal encephalocele
Q01.1 Nasofrontal encephalocele
Q01.2 Occipital encephalocele
Q01.8 Encephalocele of other sites
Q01.9 Encephalocele, unspecified
Q02 Microcephaly
Q03.0 Malformations of aqueduct of Sylvius
Q03.1 Atresia of foramina of Magendie and Luschka
Q03.8 Other congenital hydrocephalus

Q03.9 Congenital hydrocephalus, unspecified
Q04.0 Congenital malformations of corpus callosum
Q04.1 Arhinencephaly
Q04.2 Holoprosencephaly
Q04.3 Other reduction deformities of brain
Q04.4 Septo-optic dysplasia
Q04.5 Megalencephaly
Q04.6 Congenital cerebral cysts
Q04.8 Other specified congenital malformations of brain
Q04.9 Congenital malformation of brain, unspecified
Q05.0 Cervical spina bifida with hydrocephalus
Q05.1 Thoracic spina bifida with hydrocephalus
Q05.2 Lumbar spina bifida with hydrocephalus
Q05.3 Sacral spina bifida with hydrocephalus
Q05.4 Unspecified spina bifida with hydrocephalus
Q05.5 Cervical spina bifida without hydrocephalus
Q05.6 Thoracic spina bifida without hydrocephalus
Q05.7 Lumbar spina bifida without hydrocephalus
Q05.8 Sacral spina bifida without hydrocephalus
Q05.9 Spina bifida, unspecified
Q06.2 Diastematomyelia
Q07.0 Arnold-Chiari syndrome
Q11.0 Cystic eyeball
Q11.1 Other anophthalmos
Q11.2 Microphthalmos
Q11.3 Macrophthalmos
Q16.0 Congenital absence of (ear) auricle
Q16.1 Congenital absence, atresia and stricture of auditory canal (external)
Q16.2 Absence of eustachian tube
Q16.3 Congenital malformation of ear ossicles
Q16.4 Other congenital malformations of middle ear
Q16.5 Congenital malformation of inner ear
Q16.9 Congenital malformation of ear causing impairment of hearing, unspecified
Q20.0 Common arterial trunk
Q20.1Double outlet right ventricle
Q20.2 Double outlet left ventricle
Q20.3 Discordant ventriculoarterial connection
Q20.4 Double inlet ventricle
Q20.5 Discordant atrioventricular connection
Q20.6 Isomerism of atrial appendages
Q20.8 Other congenital malformations of cardiac chambers and connections
Q20.9 Congenital malformation of cardiac chambers and connections, unspecified
Q21.0 Ventricular septal defect
Q21.1 Atrial septal defect
Q21.2 Atrioventricular septal defect
Q21.3 Tetralogy of Fallot

Q21.4 Aortopulmonary septal defect
Q21.8 Other congenital malformations of cardiac septa
Q21.9 Congenital malformation of cardiac septum, unspecified
Q22.5 Ebstein's anomaly
Q22.6 Hypoplastic right heart syndrome
Q23.4 Hypoplastic left heart syndrome
Q25.0 Patent ductus arteriosus
Q25.1 Coarctation of aorta
Q25.2 Atresia of aorta
Q25.3 Stenosis of aorta
Q25.4 Other congenital malformations of aorta
Q25.5 Atresia of pulmonary artery
Q25.6 Stenosis of pulmonary artery
Q25.7 Other congenital malformations of pulmonary artery
Q25.8 Other congenital malformations of great arteries
Q25.9 Congenital malformation of great arteries, unspecified
Q26.2 Total anomalous pulmonary venous connection
Q27.3 Peripheral arteriovenous malformation
Q28.2 Arteriovenous malformation of cerebral vessels
Q28.3 Other malformations of cerebral vessels
Q33.0 Congenital cystic lung
Q33.1 Accessory lobe of lung
Q33.2 Sequestration of lung
Q33.3 Agenesis of lung
Q33.4 Congenital bronchiectasis
Q33.5 Ectopic tissue in lung
Q33.6 Hypoplasia and dysplasia of lung
Q33.8 Other congenital malformations of lung
Q33.9 Congenital malformation of lung, unspecified
Q39.0 Atresia of oesophagus without fistula
Q39.1 Atresia of oesophagus with tracheo-oesophageal fistula
Q41.0 Congenital absence, atresia and stenosis of duodenum
Q41.1 Congenital absence, atresia and stenosis of jejunum
Q41.2 Congenital absence, atresia and stenosis of ileum
Q41.8 Congenital absence, atresia and stenosis of other specified parts of small intestine
Q41.9 Congenital absence, atresia and stenosis of small intestine, part unspecified
Q45.3 Other congenital malformations of pancreas and pancreatic duct
Q60.0 Renal agenesis, unilateral
Q60.1 Renal agenesis, bilateral
Q60.2 Renal agenesis, unspecified
Q60.3 Renal hypoplasia, unilateral
Q60.4 Renal hypoplasia, bilateral
Q60.5 Renal hypoplasia, unspecified
Q60.6 Potter syndrome
Q61.1 Polycystic kidney, autosomal recessive
Q61.2 Polycystic kidney, autosomal dominant

Q61.3 Polycystic kidney, unspecified
Q61.4 Renal dysplasia
Q61.5 Medullary cystic kidney
Q61.9 Cystic kidney disease, unspecified
Q67.5 Congenital deformity of spine
Q71.0 Congenital complete absence of upper limb(s)
Q71.1 Congenital absence of upper arm and forearm with hand present
Q71.2 Congenital absence of both forearm and hand
Q71.3 Congenital absence of hand and finger(s)
Q71.4 Longitudinal reduction defect of radius
Q71.5 Longitudinal reduction defect of ulna
Q71.6 Lobster-claw hand
Q71.8 Other reduction defects of upper limb(s)
Q71.9 Reduction defect of upper limb, unspecified
Q72.0 Congenital complete absence of lower limb(s)
Q72.1 Congenital absence of thigh and lower leg with foot present
Q72.2 Congenital absence of both lower leg and foot
Q72.3 Congenital absence of foot and toe(s)
Q72.4 Longitudinal reduction defect of femur
Q72.5 Longitudinal reduction defect of tibia
Q72.6 Longitudinal reduction defect of fibula
Q72.7 Split foot
Q72.8 Other reduction defects of lower limb(s)
Q72.9 Reduction defect of lower limb, unspecified
Q73.0 Congenital absence of unspecified limb(s)
Q74.3 Arthrogryposis multiplex congenital
Q75.0 Craniosynostosis
Q75.1 Craniofacial dysostosis
Q76.1 Klippel-Feil syndrome
Q77.0 Achondrogenesis
Q77.2 Short rib syndrome
Q77.4 Achondroplasia
Q77.6 Chondroectodermal dysplasia
Q77.7 Spondyloepiphyseal dysplasia
Q77.8 Other osteochondrodysplasia with defects of growth of tubular bones and spine
Q78.0 Osteogenesis imperfecta
Q78.2 Osteopetrosis
Q79.0 Congenital diaphragmatic hernia
Q79.1 Other congenital malformations of diaphragm
Q79.2 Exomphalos
Q79.3 Gastroschisis
Q79.4 Prune belly syndrome
Q79.5 Other congenital malformations of abdominal wall
Q79.6 Ehlers-Danlos syndrome
Q79.8 Other congenital malformations of musculoskeletal system
Q79.9 Congenital malformation of musculoskeletal system, unspecified

Q80.0 Ichthyosis vulgaris
Q80.1 X-linked ichthyosis
Q80.2 Lamellar ichthyosis
Q80.3 Congenital bullous ichthyosiform erythroderma
Q80.4 Harlequin fetus
Q80.8 Other congenital ichthyosis
Q80.9 Congenital ichthyosis, unspecified
Q81.0 Epidermolysis bullosa simplex
Q81.1 Epidermolysis bullosa letalis
Q81.2 Epidermolysis bullosa dystrophica
Q81.8 Other epidermolysis bullosa
Q81.9 Epidermolysis bullosa, unspecified
Q85.0 Neurofibromatosis (nonmalignant)
Q85.1 Tuberous sclerosis
Q85.8 Other phakomatoses, not elsewhere classified
Q86.0 Fetal alcohol syndrome (dysmorphic)
Q86.1 Fetal hydantoin syndrome
Q86.2 Dysmorphism due to warfarin
Q86.8 Other congenital malformation syndromes due to known exogenous causes
Q87.0 Congenital malformation syndromes predominantly affecting facial appearance
Q87.1 Congenital malformation syndromes predominantly associated with short stature
Q87.2 Congenital malformation syndromes predominantly involving limbs
Q87.3 Congenital malformation syndromes involving early overgrowth
Q87.4 Marfan syndrome
Q87.5 Other congenital malformation syndromes with other skeletal changes
Q87.8 Other specified congenital malformation syndromes, not elsewhere classified
Q89.0 Congenital malformations of spleen
Q89.1 Congenital malformations of adrenal gland
Q89.2Congenital malformations of other endocrine glands
Q89.3 Situs inversus
Q89.4 Conjoined twins
Q89.7 Multiple congenital malformations, not elsewhere classified
Q89.8 Other specified congenital malformations
Q89.9 Congenital malformation, unspecified
Q90.0 Trisomy 21, meiotic nondisjunction
Q90.1 Trisomy 21, mosaicism (mitotic nondisjunction)
Q90.2 Trisomy 21, translocation
Q90.9 Down syndrome, unspecified
Q91 Edwards syndrome and Patau syndrome
Q92 Other trisomies and partial trisomies of the autosomes, not elsewhere classified
Q93 Monosomies and deletions from the autosomes, not elsewhere classified
Q95 Balanced rearrangements
Q96 Turner syndrome
Q97 Other sex chromosome abnormalities, female phenotype, not elsewhere classified
Q98 Other sex chromosome abnormalities, male phenotype, not elsewhere classified
Q99.0 Chimera 46,XX/46,XY

Q99.1 46,XX true hermaphrodite
Q99.2 Fragile X chromosome
Q99.8 Other specified chromosome abnormalities
Q99.9 Chromosomal abnormality, unspecified