

New York State Early Intervention Program
Diagnosed Conditions with a High Probability of Resulting in Developmental Delay
(Revised June 2025)

| ICD-CODES | ICD-DESCRIPTION |
|-----------|---|
| E70.21 | E70.21 – Tyrosinemia |
| E70.29 | E70.29 - Other disorders of tyrosine metabolism |
| E70.30 | E70.30 - Albinism, unspecified |
| E70.5 | E70.5 - Disorders of tryptophan metabolism |
| E70.8 | E70.8 - Other disorders of aromatic amino-acid metabolism |
| E78.71 | E78.71 - Barth syndrome |
| E78.72 | E78.72 - Smith-Lemli-Opitz syndrome |
| F82 | F82 – Dyspraxia (Specific developmental disorder of motor function) |
| F84.0 | F84.0 – Autistic disorder |
| F84.2 | F84.2 – Rett’s syndrome |
| F84.5 | F84.5 – Asperger’s syndrome |
| F84.8 | F84.8 – Other pervasive developmental disorders |
| F84.9 | F84.9 – Pervasive developmental disorder, unspecified |
| F90.2 | F90.2 – Attention-deficit hyperactivity disorder, combined type |
| F93.0 | F93.0 – Separation anxiety disorder of childhood |
| F93.8 | F93.8 – Other childhood emotional disorders |

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|-----------|---|
| F93.9 | F93.9 – Childhood emotional disorder, unspecified |
| F94.1 | F94.1 – Reactive attachment disorder of childhood |
| F94.8 | F94.8 – Other childhood disorders of social functioning |
| F98.9 | F98.9 – Unspecified behavioral and emotional disorders with onset usually occurring in childhood and adolescence |
| G12.0 | G12.0 – Infantile spinal muscular atrophy, type I [Werdnig-Hoffman] |
| G40.009 | G40.009 – Localization-related (local) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, not intractable, without status epilepticus |
| G40.201 | G40.201 – Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, not intractable, with status epilepticus |
| G40.411 | G40.411 – Other generalized epilepsy and epileptic syndromes, intractable, with status epilepticus |
| G40.821 | G40.821 – Epileptic spasms, not intractable, with status epilepticus |
| G40.822 | G40.822 – Epileptic spasms, not intractable, without status epilepticus |
| G40.823 | G40.823 – Epileptic spasms, intractable, with status epilepticus |
| G40.824 | G40.824 – Epileptic spasms, intractable, without status epilepticus |
| G40.834 | G40.834 - Dravet syndrome, intractable, without status epilepticus |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|---|
| G71.0 | G71.0 - Muscular dystrophy |
| G71.00 | G71.00 - Muscular dystrophy, unspecified |
| G71.11 | G71.11 - Myotonic muscular dystrophy |
| G71.19 | G71.19 - Other specified myotonic disorders |
| G71.2 | G71.2 - Congenital myopathies |
| G72.89 | G72.89 - Other specified myopathies |
| G72.9 | G72.9 - Myopathy, unspecified |
| G80.0 | G80.0 - Spastic quadriplegic cerebral palsy |
| G80.1 | G80.1 - Spastic diplegic cerebral palsy |
| G80.2 | G80.2 - Spastic hemiplegic cerebral palsy |
| G80.3 | G80.3 - Athetoid cerebral palsy |
| G80.4 | G80.4 - Ataxic cerebral palsy |
| G80.8 | G80.8 - Other cerebral palsy |
| G80.9 | G80.9 - Cerebral palsy, unspecified |
| G82.20 | G82.20 - Paraplegia, unspecified |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|--|
| G91.0 | G91.0 - Communicating hydrocephalus |
| G91.1 | G91.1 - Obstructive hydrocephalus |
| G91.9 | G91.9 - Hydrocephalus, unspecified |
| G93.1 | G93.1 - Anoxic brain damage, not elsewhere classified |
| H35.00 | H35.00 - Unspecified background retinopathy |
| H35.151 | H35.151 - Retinopathy of prematurity, stage 4, right eye |
| H35.152 | H35.152 - Retinopathy of prematurity, stage 4, left eye |
| H35.153 | H35.153 - Retinopathy of prematurity, stage 4, bilateral |
| H35.159 | H35.159 - Retinopathy of prematurity, stage 4, unspecified eye |
| H35.161 | H35.161 - Retinopathy of prematurity, stage 5, right eye |
| H35.162 | H35.162 - Retinopathy of prematurity, stage 5, left eye |
| H35.163 | H35.163 - Retinopathy of prematurity, stage 5, bilateral |
| H35.169 | H35.169 - Retinopathy of prematurity, stage 5, unspecified eye |
| H35.179 | H35.179 - Retrolental fibroplasia, unspecified eye |

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|-----------|---|
| H35.53 | H35.53 - Other dystrophies primarily involving the sensory retina |
| H35.54 | H35.54 - Dystrophies primarily involving the retinal pigment epithelium |
| H47.031 | H47.031 - Optic nerve hypoplasia, right eye |
| H47.032 | H47.032 - Optic nerve hypoplasia, left eye |
| H47.033 | H47.033 - Optic nerve hypoplasia, bilateral |
| H47.039 | H47.039 - Optic nerve hypoplasia, unspecified eye |
| H47.099 | H47.099 - Other disorders of optic nerve, not elsewhere classified, unspecified eye |
| H47.311 | H47.311 - Coloboma of optic disc, right eye |
| H47.312 | H47.312 - Coloboma of optic disc, left eye |
| H47.313 | H47.313 - Coloboma of optic disc, bilateral |
| H47.319 | H47.319 – Coloboma of optic disc, unspecified eye |
| H54.0 | H54.0 - Blindness, both eyes |
| H54.0X33 | H54.0X33 - Blindness right eye category 3, blindness left eye category 3 |
| H54.0X34 | H54.0X34 - Blindness right eye category 3, blindness left eye category 4 |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|---|
| H54.0X35 | H54.0X35 - Blindness right eye category 3, blindness left eye category 5 |
| H54.0X43 | H54.0X43 - Blindness right eye category 4, blindness left eye category 3 |
| H54.0X44 | H54.0X44 - Blindness right eye category 4, blindness left eye category 4 |
| H54.0X45 | H54.0X45 - Blindness right eye category 4, blindness left eye category 5 |
| H54.0X53 | H54.0X53 - Blindness right eye category 5, blindness left eye category 3 |
| H54.0X54 | H54.0X54 - Blindness right eye category 5, blindness left eye category 4 |
| H54.0X55 | H54.0X55 - Blindness right eye category 5, blindness left eye category 5 |
| H54.1 | H54.1 - Blindness, one eye, low vision other eye |
| H54.10 | H54.10 - Blindness, one eye, low vision other eye, unspecified eyes |
| H54.1131 | H54.1131 - Blindness right eye category 3, low vision left eye category 1 |
| H54.1132 | H54.1132 - Blindness right eye category 3, low vision left eye category 2 |
| H54.1141 | H54.1141 - Blindness right eye category 4, low vision left eye category 1 |
| H54.1142 | H54.1142 - Blindness right eye category 4, low vision left eye category 2 |
| H54.1151 | H54.1151 - Blindness right eye category 5, low vision left eye category 1 |
| H54.1152 | H54.1152 - Blindness right eye category 5, low vision left eye category 2 |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|--|
| H54.12 | H54.12 - Blindness, left eye, low vision right eye |
| H54.1213 | H54.1213 - Low vision right eye category 1, blindness left eye category 3 |
| H54.1214 | H54.1214 - Low vision right eye category 1, blindness left eye category 4 |
| H54.1223 | H54.1223 - Low vision right eye category 2, blindness left eye category 3 |
| H54.1224 | H54.1224 - Low vision right eye category 2, blindness left eye category 4 |
| H54.1225 | H54.1225 - Low vision right eye category 2, blindness left eye category 5 |
| H54.2 | H54.2 - Low vision, both eyes |
| H54.2X11 | H54.2X11 - Low vision right eye category 1, low vision left eye category 1 |
| H54.2X12 | H54.2X12 - Low vision right eye category 1, low vision left eye category 2 |
| H54.2X21 | H54.2X21 - Low vision right eye category 2, low vision left eye category 1 |
| H54.2X22 | H54.2X22 - Low vision right eye category 2, low vision left eye category 2 |
| H54.3 | H54.3 - Unqualified visual loss, both eyes |
| H54.4 | H54.4 - Blindness, one eye |
| H54.40 | H54.40 - Blindness, one eye, unspecified eye |
| H54.6 | H54.6 - Unqualified visual loss, one eye |

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|-----------|---|
| H54.60 | H54.60 - Unqualified visual loss, one eye, unspecified |
| H54.7 | H54.7 - Unspecified visual loss |
| H54.8 | H54.8 - Legal blindness, as defined in USA |
| H55.03 | H55.03 – Visual deprivation nystagmus |
| H90.0 | H90.0 - Conductive hearing loss, bilateral |
| H90.11 | H90.11 - Conductive hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side |
| H90.12 | H90.12 - Conductive hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side |
| H90.2 | H90.2 - Conductive hearing loss, unspecified |
| H90.3 | H90.3 - Sensorineural hearing loss, bilateral |
| H90.41 | H90.41 - Sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side |
| H90.42 | H90.42 - Sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side |
| H90.5 | H90.5 - Unspecified sensorineural hearing loss |
| H90.6 | H90.6 - Mixed conductive and sensorineural hearing loss, bilateral |

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|-----------|--|
| H90.71 | H90.71 - Mixed conductive and sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side |
| H90.72 | H90.72 - Mixed conductive and sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side |
| H90.8 | H90.8 - Mixed conductive and sensorineural hearing loss, unspecified |
| H90.A11 | H90.A11 – Conductive hearing loss, unilateral, right ear with restricted hearing on the contralateral side |
| H90.A12 | H90.A12 – Conductive hearing loss, unilateral, left ear with restricted hearing on the contralateral side |
| H90.A21 | H90.A21 - Sensorineural hearing loss, unilateral, right ear, with restricted hearing on the contralateral side |
| H90.A22 | H90.A22 - Sensorineural hearing loss, unilateral, left ear, with restricted hearing on the contralateral side |
| H90.A31 | H90.A31 - Mixed conductive and sensorineural hearing loss, unilateral, right ear with restricted hearing on the contralateral side |
| H90.A32 | H90.A32 - Mixed conductive and sensorineural hearing loss, unilateral, left ear with restricted hearing on the contralateral side |
| H91.3 | H91.3 - Deaf nonspeaking, not elsewhere classified |
| H91.8X3 | H91.8X3 - Other specified hearing loss, bilateral |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|--|
| P07.01 | P07.01 - Extremely low birth weight newborn, less than 500 grams |
| P07.02 | P07.02 - Extremely low birth weight newborn, 500-749 grams |
| P07.03 | P07.03 - Extremely low birth weight newborn, 750-999 grams |
| P27.0 | P27.0 - Wilson-Mikity syndrome |
| P52.22 | P52.22 - Intraventricular (nontraumatic) hemorrhage, grade 4 of newborn |
| P57.8 | P57.8 - Other specified kernicterus |
| P57.9 | P57.9 - Kernicterus, unspecified |
| P91.2 | P91.2 - Neonatal cerebral leukomalacia |
| P91.821 | P91.821 - Neonatal cerebral infarction, right side of brain |
| P91.822 | P91.822 - Neonatal cerebral infarction, left side of brain |
| P91.823 | P91.823 - Neonatal cerebral infarction, bilateral |
| P96.1 | P96.1 - Neonatal withdrawal symptoms from maternal use of drugs of addiction |
| Q01.0 | Q01.0 - Frontal encephalocele |
| Q01.1 | Q01.1 - Nasofrontal encephalocele |
| Q01.2 | Q01.2 - Occipital encephalocele |

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|-----------|---|
| Q01.8 | Q01.8 - Encephalocele of other sites |
| Q01.9 | Q01.9 - Encephalocele, unspecified |
| Q02 | Q02 – Microcephaly |
| Q03.0 | Q03.0 - Malformations of aqueduct of Sylvius |
| Q03.1 | Q03.1 - Atresia of foramina of Magendie and Luschka |
| Q03.8 | Q03.8 - Other congenital hydrocephalus |
| Q03.9 | Q03.9 - Congenital hydrocephalus, unspecified |
| Q04.0 | Q04.0 - Agenesis of corpus callosum |
| Q04.1 | Q04.1 – Arhinencephaly |
| Q04.2 | Q04.2 – Holoprosencephaly |
| Q04.3 | Q04.3 - Other reduction deformities of brain |
| Q04.4 | Q04.4 - Septo-optic dysplasia of brain |
| Q04.5 | Q04.5 – Megalencephaly |
| Q04.6 | Q04.6 - Congenital cerebral cysts |
| Q04.8 | Q04.8 - Other specified congenital malformations of brain |

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| ICD-CODES | ICD-DESCRIPTION |
|-----------|---|
| Q05.0 | Q05.0 - Cervical spina bifida with hydrocephalus |
| Q05.1 | Q05.1 - Thoracic spina bifida with hydrocephalus |
| Q05.2 | Q05.2 - Lumbar spina bifida with hydrocephalus |
| Q05.3 | Q05.3 - Sacral spina bifida with hydrocephalus |
| Q05.4 | Q05.4 - Unspecified spina bifida with hydrocephalus |
| Q05.5 | Q05.5 - Cervical spina bifida without hydrocephalus |
| Q05.6 | Q05.6 - Thoracic spina bifida without Hydrocephalus |
| Q05.7 | Q05.7 - Lumbar spina bifida without hydrocephalus |
| Q05.8 | Q05.8 - Sacral spina bifida without hydrocephalus |
| Q05.9 | Q05.9 - Spina bifida, unspecified |
| Q07.01 | Q07.01 - Arnold-Chiari syndrome with spina bifida |
| Q07.02 | Q07.02 - Arnold-Chiari syndrome with hydrocephalus |
| Q07.03 | Q07.03 - Arnold-Chiari syndrome with spina bifida and hydrocephalus |
| Q11.2 | Q11.2 – Microphthalmos |
| Q13.1 | Q13.1 – Absence of iris |

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|-----------|---|
| Q13.4 | Q13.4 - Other congenital corneal malformations |
| Q14.2 | Q14.2 - Congenital malformation of optic disc |
| Q16.0 | Q16.0 - Congenital absence of (ear) auricle |
| Q16.1 | Q16.1 - Congenital absence, atresia and stricture of auditory canal (external) |
| Q16.3 | Q16.3 - Congenital malformation of ear ossicles |
| Q16.4 | Q16.4 - Other congenital malformations of middle ear |
| Q16.5 | Q16.5 - Congenital malformation of inner ear |
| Q16.9 | Q16.9 - Congenital malformation of ear causing impairment of hearing, unspecified |
| Q35.1 | Q35.1 - Cleft hard palate |
| Q35.3 | Q35.3 - Cleft soft palate |
| Q35.5 | Q35.5 - Cleft hard palate with cleft soft palate |
| Q35.7 | Q35.7 - Cleft uvula |
| Q35.9 | Q35.9 - Cleft palate, unspecified |
| Q36.0 | Q36.0 - Cleft lip, bilateral |
| Q36.1 | Q36.1 - Cleft lip, median |

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|-----------|---|
| Q36.9 | Q36.9 - Cleft lip, unilateral |
| Q37.0 | Q37.0 - Cleft hard palate with bilateral cleft lip |
| Q37.1 | Q37.1 - Cleft hard palate with unilateral cleft lip |
| Q37.2 | Q37.2 – Cleft soft palate with bilateral cleft lip |
| Q37.3 | Q37.3 - Cleft soft palate with unilateral cleft lip |
| Q37.4 | Q37.4 - Cleft hard and soft palate with bilateral cleft lip |
| Q37.5 | Q37.5 - Cleft hard and soft palate with unilateral cleft lip |
| Q37.8 | Q37.8 - Unspecified cleft palate with bilateral cleft lip |
| Q37.9 | Q37.9 - Unspecified cleft palate with unilateral cleft lip |
| Q71.61 | Q71.61 - Lobster-claw right hand |
| Q71.62 | Q71.62 - Lobster-claw left hand |
| Q71.63 | Q71.63 - Lobster-claw hand, bilateral |
| Q71.899 | Q71.899 - Other reduction defects of unspecified upper limb |
| Q71.90 | Q71.90 - Unspecified reduction defect of unspecified upper limb |

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| Q72.899 | Q72.899 - Other reduction defects of unspecified lower limb |
| Q73.0 | Q73.0 - Congenital absence of unspecified limb(s) |
| Q73.1 | Q73.1 - Phocomelia, unspecified limb(s) |
| Q73.8 | Q73.8 - Other reduction defects of unspecified limb(s) |
| Q74.3 | Q74.3 - Arthrogryposis multiplex congenital |
| Q74.8 | Q74.8 - Other specified congenital malformations of limb(s) |
| Q75.3 | Q75.3 – Macrocephaly |
| Q86.0 | Q86.0 - Fetal alcohol syndrome (dysmorphic) |
| Q87.1 | Q87.1 - Congenital malformation syndromes predominantly associated with short stature |
| Q87.11 | Q87.11 - Prader-Willi Syndrome |
| Q87.2 | Q87.2 - Congenital malformation syndromes predominantly involving limbs |
| Q87.3 | Q87.3 - Congenital malformation syndromes involving early overgrowth |
| Q87.5 | Q87.5 - Other congenital malformation syndromes with other skeletal changes |
| Q87.81 | Q87.81 - Alport syndrome |
| Q87.89 | Q87.89 - Other specified congenital malformation syndromes, not elsewhere classified |

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| Q89.4 | Q89.4 - Conjoined twins |
| Q89.7 | Q89.7 - Multiple congenital malformations, not elsewhere classified |
| Q89.8 | Q89.8 - Other specified congenital malformations |
| Q90.0 | Q90.0 - Trisomy 21, nonmosaicism (meiotic nondisjunction) |
| Q90.1 | Q90.1 - Trisomy 21, mosaicism (mitotic nondisjunction) |
| Q90.2 | Q90.2 - Trisomy 21, translocation |
| Q90.9 | Q90.9 - Down syndrome, unspecified |
| Q91.0 | Q91.0 - Trisomy 18, nonmosaicism (meiotic nondisjunction) |
| Q91.1 | Q91.1 - Trisomy 18, mosaicism (mitotic nondisjunction) |
| Q91.2 | Q91.2 - Trisomy 18, translocation |
| Q91.3 | Q91.3 - Trisomy 18, unspecified |
| Q91.4 | Q91.4 - Trisomy 13, nonmosaicism (meiotic nondisjunction) |
| Q91.5 | Q91.5 - Trisomy 13, mosaicism (mitotic nondisjunction) |
| Q91.6 | Q91.6 - Trisomy 13, translocation |
| Q91.7 | Q91.7 - Trisomy 13, unspecified |

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| Q92.1 | Q92.1 - Whole Chromosome Trisomy Mosaicism |
| Q92.8 | Q92.8 - Other specified trisomies and partial trisomies of autosomes |
| Q93.59 | Q93.59 - Other deletions of part of a chromosome |
| Q93.51 | Q93.51 - Angelman syndrome |
| Q99.2 | Q99.2 - Fragile X chromosome |
| S14.109A | S14.109A - Unspecified injury at unspecified level of cervical spinal cord, initial encounter |
| S24.109A | S24.109A - Unspecified injury at unspecified level of thoracic spinal cord, initial encounter |
| S34.109A | S34.109A - Unspecified injury to unspecified level of lumbar spinal cord, initial encounter |
| S34.139A | S34.139A - Unspecified injury to sacral spinal cord, initial encounter |
| F43.10 | F43.10 - Post-traumatic stress disorder, unspecified |
| F43.12 | F43.12 - Post-traumatic stress disorder, chronic |
| G40.901 | G40.901 - Epilepsy, unspecified, not intractable, with status epilepticus |
| G40.909 | G40.901 - Epilepsy, unspecified, not intractable, with status epilepticus |
| G47.35 | G47.35 - Congenital central alveolar hypoventilation syndrome |

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| G40.109 | G40.109 - Local-rel (focal) (partial) symptomatic epilepsy and epileptic syndromes w/ simple partial seizures, not intractable, w/o status epilepticus |
| G40.89 | G40.89 - Other seizures |
| P91.60 | P91.60 - Hypoxic ischemic encephalopathy [HIE], unspecified |
| P91.61 | P91.61 - Mild hypoxic ischemic encephalopathy [HIE] |
| P91.62 | P91.62 - Moderate hypoxic ischemic encephalopathy [HIE] |
| P91.63 | P91.62 - Moderate hypoxic ischemic encephalopathy [HIE] |
| Q93.52 | Q93.52 - Phelan-McDermid syndrome |