DELAWARE HEALTH AND SOCIAL SERVICES
Division of Public Health
Birth to Three Early Intervention Program

birth to three
Early Intervention Program

# Established Conditions List

| | |
|---|---|
| Date: | April 27, 2021 |
| From: | Birth to Three Early Intervention Program – Administration |
| Approval Date: | June 29, 2023 |
| Signature : | Nicole Topper |
| Effective Date: | May 27, 2021 |
| Last Review Date: | June 22, 2023 |

## Purpose

The Birth to Three Early Intervention Program (the Program) maintains an Established Condition List that identifies prevalent physical or mental conditions that would qualify infants and toddlers for Part C services, in accordance with §303.21(a)(2) of the Individuals with Disabilities Education Act. The list is not exhaustive; if an infant or toddler is referred to the Program with a diagnosed physical or mental condition that is not on the Established Condition list, but meets the criteria set forth in §303.21 (2), the infant or toddler may be determined eligible for Part C services.

To appropriately reflect the evolving needs of children and families in Delaware, the established conditions list has been updated and will be implemented effective immediately upon administrative signature and approval.

**"*" = follow until 12 months post referral**

**"++" =applies until second birthday**

Established Conditions List

***BIRTH MANDATE/Delaware***

| | ***IDEA designations: sensory impairments; severe attachment disorders*** |
|---|---|
| F84.0 | Autism Spectrum disorder |
| H93.3X9 | Auditory Neuropathy Spectrum Disorder (ANSD) (Disorder of Acoustic Nerve) |
| H90.2 | Conductive hearing loss, unspecified |
| H90.8 | Mixed conductive and sensorineural hearing loss, unspecified |
| H90.5 | Sensorineural Hearing Loss (Unspec) |
| H90.4 | Sensorineural hearing loss, unilateral |
| H91.93 | Unspecified hearing loss - Bilateral |
| H90.2 | Cond. hearing loss, unspecified |
| H90.8 | Mixed cond & sensorineural hearing loss, unspec |
| H54 | Blindness NOS |
| H47.619 | Cortical Blindness, unspecified |
| H47.619 | Cortical Blindness, unspecified |

***MULTIPLE SYSTEM/ GENETIC/ CHROMOSOMAL***

| | ***IDEA designation: chromosomal abnormalities; genetic or congenital disorders; inborn errors of metabolism*** |
|---|---|
| Q93.5 | Angelman Syndrome |
| Q97.0 | Chromosomal Anomalies – Karyotype 47, XXX |
| E70.0 | Classical phenylketonuria |
| Q90.9 | Down Syndrome |
| Q99.2 | Fragile X Syndrome |
| Q75.4 | Franceschetti–Klein (Wildervanck) Syndrome |
| E75.22 | Gaucher Disease (Lipidoses) |
| E71.0 | Maple-Syrup-Urine Disease |
| G71.0 | Muscular Dystrophy |
| Q85.02 | Neurofibromatosis type- 2 |
| Q78.0 | Osteogenesis imperfecta |
| Q87.0 | Saethre–Chotzen syndrome (aka Acrocephalosyndactyly type III) |
| Q87.2 | TAR Syndrome: (Thrombocytopenia with Absent Radius Syndrome) |
| Q75.4 | Treacher Collins Syndrome |
| Q97.0 | Triple X Syndrome (Chromosomal Anom) |
| Q91.7 | Trisomy 13 (Patau's Syndrome) Unspec |
| Q91.3 | Trisomy 18 (Edwards Syndrome) Unspec |
| Q85.1 | Tuberous sclerosis |
| Q96.9 | Turner-Ullrich Syndrome |

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Established Conditions List

***NEUROLOGIC: congenitally or postnatally acquired***

| ***IDEA designation: disorders reflecting disturbance of the development of the nervous system*** | |
|---|---|
| Q04.0 | Agenesis of Corpus Callosum |
| G93.1 | Anoxic brain damage |
| Q07.02 | Arnold-Chiari syndrome with hydrocephalus |
| G60.0 | Atrophy, Charcot-Marie-tooth Syndrome |
| I63.50 | Cerebral Artery Occlusion NOS |
| G80.8 | Cerebral Diplegia / Hemiplegia (congenital) |
| P10.1 | Cerebral Hem. due to birth injury (grade III & IV bleed) |
| G80.9 | Cerebral Palsy, unspecified |
| G80.0 | Cerebral Spastic Quadriplegia |
| G04.90 | Encephalitis (postnatally acquired) |
| Q07.9 | Encephalopathy (congenital) |
| Q01 | Encephalocele |
| G93.40 | Encephalopathies (Degenerative) NOS |
| D18.02 | Hemangioma of intracranial struct. |
| P91.660 | HIE/Body Cooling (Hypoxic Ischemic Enceph.) * |
| Q04.2 | Holoprosencephaly |
| Q03.8 | Hydrocephalus (congenital); Other |
| Q07.9 | Hypomyelination affecting Meningeal bands or folds (congenital) |
| I61.9 | Intracerebral (nontraumatic) Hemorhhage * |
| G06.0 | Intracranial abscess and granuloma |
| P52.0 | Intraventricular Hem./Newborn, only Grade III & IV |
| E75.4 | Jansky-Bielschowsky Amaurotic Idiocy |
| E75.23 | Krabbe disease |
| G04.81 | Leukoencephalitis |
| P91.2 | Leukomalacia, Neonatal Cerebral |
| Q05.9 | Lipomyelomeningocele |
| Q04.9 | Malformation of Brain (Cong/Unspec) * |
| C71.8 | Malig neoplasm of overlapping sites of brain |
| Q04.5 | Megalencephaly |
| G00.2 | Meningitis – Streptococcal* |
| A02.21 | Meningitis due /Salmonella infect.* |
| G03.9 | Meningitis NOS* |
| Q02 | Microcephalus, microcephaly (under 3%) |
| Q04.3 | Microgyria |
| G70.9 | Myoneural disorder |
| G71.11 | Myotonic Dystrophy |
| Q01.1 | Nasofrontal Encephalocele* |

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| | |
|---|---|
| E75.4 | NCL: Ceroid-Lipofuscinosis, Neuronal |
| E75.4 | NCL: Batten-Mayou Disease |
| E75.4 | NCL: Kuf's Disease (NCL) |
| H55.01 | Nystagmus (congenital) |
| G91.1 | Obstructive Hydrocephalus |
| H47.039 | Optic Nerve Hypoplasia Unspec |
| Q04.8 | Other Specified Congenital Malformation of the Brain * |
| Q04.3 | Polymicrogyria |
| Q04.6 | Porencephalic Cysts (Congenital) |
| Q04.6 | Schizencephaly |
| Q05.4 | Spina Bifida |
| G12.0 | Spinal Muscular Atrophy (Werdnig-Hoffman) |
| E75.02 | Tay Sachs (Nervous System Disorder) |

***NEUROLOGIC - SEIZURES***

| | |
|---|---|
| P90 | Convulsions of Newborn* (only if they are intractable, or with ongoing medication treatment) |
| R56.9 | Convulsion NOS* (only if they are intractable, or with ongoing medication treatment) |
| G40.823 | Epil spasms, **intractable**, w/ status epilepticus |
| G40.824 | Epil spasms, **intractable**, w/o status epilepticus |
| G40.919 | Epilepsy, unspec, **intractable**, w/o status epilepticus |
| G40.911 | Epilepsy, unspec, **intractable**, w/status epilepticus |

***INFECTIONS***

| ***IDEA designation: congenital infections*** | |
|---|---|
| P35.1 | Cytomegalovirus Infection, Congenital (only if symptomatic) |
| P35.0 | Rubella (Prenatally acquired) |
| A53.9 | Syphilis (Prenatally acquired) |
| B58.9 | Toxoplasmosis (Prenatally acquired) |
| P39.9 | Varicella (Perinatally acquired) |
| A92.5 | Zika Disease only with Microcephaly (not just exposure) |

***INFECTIONS: other***

| | |
|---|---|
| A48.51 | Botulism (Infant)* |

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# Established Conditions List

## *PRENATAL SUBSTANCE EXPOSURE*

| *IDEA designation: disorders secondary to exposure of substances, including fetal alcohol syndrome* | |
|---|---|
| Q86.0 | Fetal Alcohol Syn. (dysmorphic, not just alcohol exposure) |
| P96.1 | NAS - Neonatal Abstinence Syndrome ++ |
| P04.1 | Phenytoin Exposed (Prenatally) |

## *CARDIOPULMONARY*

| | |
|---|---|
| P27.1 | Bronchopulmonary dysplasia (BPD)* - Only if Oxygen dependent |
| I42.8 | Cardiomyopathy (Primary)* |
| Q25.1 | Coarctation of Aorta (pre/postductal)* |
| J44.9 | Chronic/Obstructive Lung Disease |
| Q20.8 | Cor Biloculare (Cong. Malf. of Cardiac Chambers and Connections) |
| Q20.4 | Double inlet ventricle |
| Q20.2 | Double Outlet left ventricle |
| Q20.1 | Double outlet right ventricle |
| Z92.81 | ECMO - Extracorporeal Membrane Oxygenation |
| Z94.1 | Heart Transplant Status |
| I11.0 | Hypertensive Heart Disease w/ Heart Failure |
| Q23.4 | Hypoplastic Left Heart Syndrome |
| Q22.6 | Hypoplastic Right Heart Syndrome |
| Q24.9 | Malform of the Heart, Congenital/Unspec* |
| I21.3 | Myocardial Infarction Unspec (STEMI) |
| Z99.81 | Oxygen Dependent (Supplemental) |
| Q22 | Pulmonary Valve Atresia |
| Z99.11 | Respirator or Ventilator Dependent |
| Z93.0 | Tracheostomy Status |
| Q21.3 | Tetralogy of Fallot |
| I50.9 | 150.9 Heart Failure *, unspecified |

## *ENDOCRINE*

| | |
|---|---|
| E03.1 | Hypothyroidism (congenital)* |
| E23.2 | Diabetes Insipidus |
| E11.9 | Diabetes, Diabetic - unspecified |

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# Established Conditions List

***STRUCTURAL***

| | |
|---|---|
| S68.412S | Amputation of Left Hand @ Wrist |
| S68.411S | Amputation of Right Hand @ Wrist |
| Q35.9 | Cleft Palate* post repair |
| Q75.9 | Cong Malform of Skull and Face Unspec* |
| Q16.4 | Congenital Malformation of middle ear NOS |
| Q17.9 | Congenital Malformations of Ear NOS |
| Q17.2 | Microtia |
| Q60.2 | Renal agenesis, bilateral, unspecified |

| ***TRAUMA RELATED*** | |
|---|---|
| T74.4 | Shaken Infant Syndrome |
| S06.2X0D | Traumatic Brain Injury, Diffused w/o LOC |

| ***OTHER*** | |
|---|---|
| Q74.3 | Arthrogryposis |
| Z94.81 | Bone Marrow Transplant* |
| P07.01 | Ex low weight nb, less 500 gr* |
| P07.02 | Ex low weight nb, less 500-749 gr* |
| P07.03 | Ex low weight nb, less 750-999 gr* |
| P07.26 | Less than 28 weeks gestation * |
| E74.21 | Galactosemia* |
| E74.00 | Glycogen storage disorder |
| E76.02 | Hurler-Scheie Syndrome |
| R20.1 | Hypoesthesia of Skin |
| Z77.011 | Lead Exposure (Toxic) If level **5 ug/dl** or higher* |
| P77.9 | Necrotizing enterocolitis/NB, unspec* (only if post surgical) |
| H35.109 | Retinopathy of Prematurity, unspec elig., (Grade 3 and above only) |
| G46.5 | PURA syndrome |
| Q477 | Alagille Syndrome * |

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