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:	April 16, 2024
Signature :	Nicole Topper
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Last Review Date:	April 16, 2024

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
H54.7	Vision Loss 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
Q87.89	Bardet-Biedl Syndrome
Q87.3	Beckwith-Wiedemann Syndrome
Q99.9	Chromosomal Abnormality NOS (Deletion/Dup)
Q99.9	HNRNPU-NDD
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)
Q98.4	Klinefelter's Syndrome
E72.03	Lowe's Syndrome
E71.0	Maple-Syrup-Urine Disease
G71.0	Muscular Dystrophy
Q85.02	Neurofibromatosis type- 2
Q87.19	Noonan Syndrome
Q78.0	Osteogenesis imperfecta
Q87.0	Saethre–Chotzen syndrome (aka Acrocephalosyndactyly type III)

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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
Q90.9	Trisomy 21 (Down Syndrome) Unspec
Q85.1	Tuberous sclerosis
Q96.9	Turner-Ullrich Syndrome

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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*
E75.4	NCL: Ceroid-Lipofuscinosis, Neuronal

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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)*

PRENATAL SUBSTANCE EXPOSURE

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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

Q39.1	Atresia of Esophagus with trachea-esophageal fistula
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)
Q20.3	Discordant Ventriculoarterial Connection
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	Heart Failure *, unspecified

ENDOCRINE

E03.1	Hypothyroidism (congenital)*
E23.2	Diabetes Insipidus
E11.9	Diabetes, Diabetic - unspecified
E20.1	Pseudohypoparathyroidism

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STRUCTURAL

Z89.431	Acquired Absence of Right Foot
Z89.432	Acquired Absence of Left Foot
Z89.439	Acquired Absence of Unspecified Foot
Q72.03	Amelia of the Lower Extremities (bilateral)
S68.412S	Amputation of Left Hand @ Wrist
S68.411S	Amputation of Right Hand @ Wrist
Q76.49	Caudel Regression Syndrome
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
Q31.5	Laryngomalacia
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)
E74.02	Pompe Syndrome
G46.5	PURA syndrome
Q477	Alagille Syndrome *

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