CNMI Early Intervention Program

Established Condition List (but not limited to) that indicate automatic eligibility for Early Intervention Services

Infant or Toddler with a Disability

§303.21 (2) Has a diagnosed physical or mental condition that-

(i) Has a high probability of resulting in developmental delay; and

(ii) Includes conditions such as chromosomal abnormalities; genetic or congenital disorders; sensory impairments; inborn errors of metabolism; disorders reflecting disturbances of the development of the nervous system; congenital infections; severe attachment disorders; and disorders secondary to exposure to toxic substances, including fetal alcohol syndrome.

Chromosomal Abnormalities/Genetic or Congenital Disorders
Albinism
Albright Hereditary
Angelman Syndrome (Happy Puppet Syndrome)
Achondroplasia (dwarfism)
Acrocallosal Syndrome, Schinzel Type (Absence of Corpus Callosum, Schinzel Type, ACS Hallux Duplication, Postaxial Polydactyly, Absence of Corpus Callosum, Schinzel Acrocallosal Syndrome, ACLS)
Adrenoleukodystrophy
Amelia
Antley-Bixler Syndrome (Multisynostotic Osteodysgenesis, Craniosynostosis, Choanal Atresia, Radial Humeral Synostosis, Trapezoidocephaly-Multiple Synostosis Syndrome, ABS, Multisynostotic Osteodysgenesis with Long Bone Fractures)
Apert Syndrome (Acrocephalosyndactyly)
Arthrogryposis Multiplex Congenita
Ataxia
Ataxia-Telangiectasia Syndrome (Louis-Bar Syndrome)
Beals Syndrome (Congential Contractural Arachnodactyly, Hecht-Beals Syndrome)
Beckwith-Wiedemann Syndrome
Canavan Disease
Cardio-Facio-Cutaneo Syndrome
Cerebral Lipdosis
Cerebro-Oculo-Factio-Skeletal (COFS) Syndrome
CHARGE Syndrome/Association
Chromosome 10p+, 11p-, 12p-, 13q-, 13q+, 18q-, 21q-, 22q-, 3q+, 4q-, 4Q+, 5p- Syndromes
Coffin-Lowry Syndrome
Coffin-Siris Syndrome
Cornelia de Lange Syndrome (Brachmann de Lange
Cri-du-chat Syndrome (Deletion 5p Syndrome)
Cystic Fibrosis

Dandy Walker Syndrome
Down Syndrome (Trisomy 21)
Duchenne Muscular Dystrophy
Dyggve-Melchio-Clausen Syndrome (DMC Disease, DMC Syndrome, Smith-McCort Dysplasia)
Fanconi Syndrome
Fragile X Syndrome
Fraser Syndrome (Cryptophthalmos Syndrome, Meyer-Schwickerath's syndrome, Fraser-Francois syndrome, Ullrich-Feichtiger syndrome)
Gaucher Syndrome (Glucosylceramide storage disease; GSDI)
Glutaric Aciduria Type I & Type II
Hypothyroidism (congenital)
Jeune Syndrome
Joubert Syndrome
Klinefelter Syndrome
Krabbe's disease
Lissencephaly Syndrome (Miller-Dieker Syndrome, Agyria)
Mucolipidosis II, III
Noonan Syndrome
Organic Acidemias
Pelizaeus-Merzbacher disease
Peroxisomal Disorders
Phenylketonuria (PKU)
Phelan-McDermid syndrome
Prader-Willi Syndrome
Rubenstein-Taybi Syndrome
Schwartz-Jampel Syndrome
Steinert Myotonic Dystrophy Syndrome (Curschmann-Batten-Steinert syndrome)
Treacher-Collins Syndrome
Trisomy 8
Trisomy 9
Tetrasomy 12p
Trisomy 13 (Patau Syndrome)
Trisomy 18 (Edward's Syndrome)
Tuberous Sclerosis Complex
Urea Cycle Defect
Very long chain fatty acid storage diseases
Waardenburg Syndrome, Types I and II
Walker-Warburg Syndrome (XO)
Williams Syndrome
Zellweger Syndrome (Cerebro-Hepato-Renal Syndrome)

Sensory Impairments
Auditory Neuropathy
Blindness/Visual Impairments
Hearing Loss
Optic Nerve Hypoplasia (DeMorsier's Syndrome, Septo Optic Dysplasia)
Progressive hearing loss as related to syndromes such as neurofibromatosis, osteopetrosis, and Usher's
Retinopothy of Prematurity (ROP)

Inborn Errors of Metabolism
Galactosemia
Glycogen Storage Disease
Hunter Syndrome
Hurlur-Sheie Syndrome
Hyperadrenolcortical/ Hypoadrenalcortical Function
Hyperparathyroidism
Hyperpituitary/Hypopituitary
Infant PKU
Infantile Gaucher Disease
Lesch-Nyhan Syndrome
Maple Syrup Disease
Murquio Syndrome
Maroteaux Syndrome
Neimann Pick Disease
Sanflippo Syndrome
Shceie Syndrome
Sly Syndrome
Tay Sachs Disease

Disorders Reflecting Disturbances of the Development of the Nervous System
Agyria (Miller-Dieker lissencephaly syndrome (MDLS), agyria syndrome, agyriapachygyria syndrome, classical lissencephaly)
Aicardi Syndrome
Alpers Syndrome/Disease
Apert Syndrome (Acrocephalosyndactyly)
Aphasia
Arachnoid cyst with neuro-developmental delay
Arhinencephaly (Holprosencephaly)
Arnold-Chiari syndrome, type II (Malformation d'Arnold-Chiari)
Ataxia
Cerebral Palsy
CNS Aneurysm with Neuro-Developmental Delay
CNS Tumor with Neuro- Developmental Delay
Encephalopathy, congenital only
Encephalopathy, Static

Erb's Palsy (Brachial Plexus Injury, Perinantal Origin)
Holoprosencephaly
Hypertonia (persistent only)
Hypophosphotasia-Infantile
Lennox-Gastaut Syndrome
Intraventricular hemorrhage (III or IV)
Meningocele (cervical)
Miller-Dieker Syndrome
Mitochondrial Disorder
Multiple anomalies of the brain
Myopathy
Neural Tube Defect
Spinocerebellar Disorders
TAR (Thrombocytopenia-Absent Radii syndrome)
Traumatic Brain Injury (Head Trauma**)**

Congenital Infections
Cytomegalovirus (CMV)
Herpes
HIV+
Rubella
Syphillis
Taxoplasmosis

Severe Attachment Disorders
Anxiety Disorders of Infancy and Early Childhood
Depression of Infancy and Early Childhood
Infantile Anorexia

Disorders Secondary to Exposure to Toxic Substances
Fetal Alcohol Syndrome

Others
Asperger's Disorder
Autism Spectrum Disorder (ASD)
Substantiated Child Abuse/Child Neglect
Childhood Disintegrative Disorder
Failure to Thrive (FTT)
Pervasive Developmental Disorder (PDD)
Rett's Syndrome
Hydrocephalus (congenital or acquired)
Low Birth Weight (<1,500 grams at birth)
Any loss of skills
Lack of (no) verbal skills by 16 months of age