Nevada Part C: Medical/Auto Eligible list for Nevada Early Intervention Services

The below categories represent a comprehensive list of diagnostic categories which are considered to ensure that certain infants/toddlers are automatically eligible for Part C Early Intervention Services. However, it is the physicians within the Early Intervention System (EIS) who make final determinations. Any referral coming into the EIS must be accompanied by physician documentation to support eligibility.

Auto eligibility categories include:

Arthrogryposis/Major Limb Malformations – Rare condition characterized by stiff joints and abnormally developed muscles and not a specific diagnosis but a clinical finding:

 Amniotic Band Syndrome Amputations

Central Nervous System Disorders/Diseases – Affect brain, spinal cord and other nervous tissues throughout the body:

- Encephalopathy
- Bacterial or Herpes Meningitis
- Infantile Spasms
- Seizure Disorder
- Sensory feeding or other
- Abnormal neurologic exam

- degenerative Disorder

Chromosomal Abnormalities - Existing in the genetic structure of the infant's chromosomes:

- Angelman Syndrome
- Velo-Cardio-Facial

• Williams Syndrome

- Craniofacial Malformations Diverse group of deformities in the growth of the head and facial bones:
 - Williams Syndrome
- Goldenhar Syndrome

- DiGeorge Sequence
- Treacher-Collins Syndrome
- Cleft Palate: soft or hard (not cleft lip) Congenital Aural Atresial/Microtia

Neurocutaneous Syndromes – genetic disorders leading to tumor growth in the body which affects development:

- Neurofibromatosis
- Tuberous Sclerosis

• Sturge-Weber Syndrome

Muscular Dystrophy

Hemplegia

Tracheoesophageal Fistula

Neuromuscular Disorders -- Affects nerves and impairs functioning of muscles:

- Erb's Palsy
- Cerebral Palsy
- Mitochondrial Disease • Myopathies
- Spinal Muscular Atrophy
- Diaphragmatic Hernia
- Extracorporeal Membrane Oxygenation (ECMO)
- Approved by IDEA Part C Office: May 2014

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- Cri-du-chat Syndrome
- Fragile X Syndrome
- Unbalanced Translocations
- Grade II with atypical tone

Hydrocephalus

• Traumatic Brain Injury

Shaken baby syndrome

IVH Grade III or IV and

- Major brain malformations Periventricular Leukomalacia
- Kernicterus Neuro-

- CHARGE Syndrome/Association
- Down Syndrome Klinefelter Syndrome
 - - - Pierre-Robin Sequence

 - Prader-Willi Syndrome

- Trisomy Syndromes/Deletions (all)

Other Disorders or Conditions associated with, or has a significant potential of, leading to a developmental delay:

- Autism Spectrum Disorder
- Child Maltreatment Syndrome • Exposure to Toxic Substances
- Cyanotic Heart Disease
- Fetal Alcohol Syndrome

• Central Visual impairment (Cortical –Cerebral)

Unilateral or Bilateral hearing loss (permanent)

- Cystic Fibrosis Tracheostomy
- Hypoxic lschemic Encephalopathy Inborn Errors of Metabolism Organic and/or Psychosocial Failure to Thrive

• Cortical visual processing disorder

equal to or greater than 40db

- Perinatal Drug Affect with Microcephaly
- Extreme Prematurity (</= 1,000 grams/2.20 pounds or <=27 weeks gestation up to 18 months corrected)

Sensory Impairment – Vision and hearing loss which may lead to learning and other delays:

- Cerebral visual impairment
- Optic Nerve Hypoplasia (ONH)
- Septo-Optic Dysplasia (SOD)
- Retinopathy of prematurity (with laser treatment)

NOTE: Failing the newborn hearing screening is not a part c eligible until hearing loss is confirmed.

Spinal Cord Injuries or Defects – Usually in utero and leads to structure defects involving the spine and spinal cord:

- Myelomeningocele
- Tethered Spinal Cord Syndrome
- Spina Bifida with Tethered Cord (unless treated)

TORCH Infections – Perinatal infections which can lead to fetal anomalies:

• CMV

• Herpes

Other Rubella

Toxoplasmosis

FURTHER DISTINCTIONS:

The above categories are not a restrictive/inclusive list and rely on doctors, who have the expertise, to make informed clinical decisions to include or exclude similar disorders which have not been included above. This includes symptomatology that leads to, or has significant risk of, a developmental delay, including: Acute Lymphoid Leukemia, Chronic Myelogenous Leukemia, and Cancers.

Some conditions can be ameliorated with medical/surgical attention, at which point the child can exit the program. The above list includes specific disorders or diseases; however, there are treatments which or symptoms may necessitate the infant/toddler be determined auto-eligible if supporting documentation includes appropriate references, including: NG-tube, G-tube dependent upon discharge, cool cap treatment requirements and others. Any time variance from categories is needed, appropriate documentation must be included in the file with the decision.

NOTE: At any point a child is no longer eligible for early intervention, due to lack of supporting documentation to support eligibility (and informed clinical opinion is that the child is not eligible), received appropriate medical/surgical intervention, has met developmental milestones, etc – the infant/toddler is to be discharged from early intervention services.