**Massachusetts Department of Public Health**

**Early Intervention**

**DIAGNOSED CONDITIONS LIST**

*(Effective: October 1, 2015)*

All children diagnosed with one of the following non-asterisked established conditions that have a high probability of resulting in developmental delay, are eligible for early intervention services until the child’s third birthday, unless a change in the status of a diagnosis or condition resolves with medical/surgical treatment. Those diagnoses or conditions that may improve are denoted on the list with an asterisk \* and are underlined. These conditions can be entered on only ONE evaluation and will have an eligibility timeframe of one year. The evaluation having this condition does NOT have to be the initial evaluation. No other evaluations for this child can have this same asterisked established condition.

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| **SENSORY CONDITIONS** |
| *BLINDNESS* |  |
| H54.0 |  | Blindness, both eyes (Note: 10/1/17 not valid for insurers) |
| H54.40 |  | Blindness, one eye |
| H47.619 |  | Cortical Blindness/Cortical Vision Impairment |
| H47.20 |  | Optic Nerve Atrophy |
| H35.179 | \* | Retinopathy of Prematurity |
| *LOW VISION* |
| H54.2 |  | Low vision, both eyes (Note: 10/1/17 not valid for insurers) *(20/70 best acuity with correction)* |
| H54.52 | \* | Low vision, left eye, normal vision right eye (Note: 10/1/17 not valid for insurers) |
| H54.51 | \* | Low vision, right eye, normal vision left eye (Note: 10/1/17 not valid for insurers) |
| *HEARING LOSS (permanent)* |
| H91.93 |  | Hearing impairment, bilateral |
| H91.92 |  | Hearing impairment, left ear |
| H91.91 |  | Hearing impairment, right ear |
| H90.3 |  | Neural hearing loss/auditory neuropathy |
| **CANCERS** |
| C80.1 |  | Cancer, Other *(not included on this list)* |
| C95.90 |  | Leukemia |
| C82.50 |  | Lymphoma |
| C71.9 |  | Malignant neoplasm of brain |
| C64.9 |  | Malignant neoplasm of kidney |
| C41.9 |  | Osteosarcoma |
| C49.3 |  | Rhabdomyosarcoma |
| **CARDIAC CONDITIONS** |
| Q21.2 |  | Atrioventricular canal defect |
| I42.9 |  | Cardiomyopathy |
| Q25.1 |  | Coarctation of the aorta |
| I27.0 |  | Hypertension, pulmonary |
| Q23.4 |  | Hypoplastic left heart syndrome |
| Q24.9 |  | Major cardiac anamoly, other *(not included on this list)* |
| Q21.3 |  | Tetralogy of Fallot |
| Q20.3 |  | Transposition of great vessels |
| **CENTRAL NERVOUS SYSTEM DISORDERS** |
| Q04.0 |  | Aicardi syndrome |
| Q00.0 |  | Anencephaly |
| G11.3 |  | Ataxia-Telangiectasia |
| G37.9 |  | Brain Sclerosis |
| G12.22 |  | Bulbar palsy |
| Q04.9 |  | Cerebral atrophy, congenital |
| G80.9 |  | Cerebral palsy |
| I61.9 |  | Cerebrovascular accident *(CVA)* |
| Q04.6 |  | Congenital Schizencephaly |
| Q03.1 |  | Dandy-Walker malformation |
| G80.1 |  | Diplegia |
| G04.90 |  | Encephalitis |
| Q01.9 |  | Encephalocele |
| G93.40 |  | Encephalopathy |
| G40.901 |  | Epilepsy |
| **CENTRAL NERVOUS SYSTEM DISORDERS** *(Continued)* |
| G81.90 |  | Hemiparesis/Hemiplegia |
| Q04.2 |  | Holoprosencephaly |
| Q04.3 |  | Hydranencephaly  |
| Q04.3 |  | Hypoplasia of the brain |
| G40.401 |  | Hypsarrhythmia |
| P52.21 | \* | Intraventricular hemorrhage *(grade 3)* |
| P52.22 | \* | Intraventricular hemorrhage *(grade 4)* |
| E75.25 |  | Leukodystrophy/Canavan disease |
| Q04.3 |  | Lissencephaly |
| G03.9 |  | Meningitis *with negative long-term effects* |
| Q02 |  | Microcephaly |
| P96.1 | \* | Neonatal Abstinence Syndrome |
| Q04.3 |  | Polymicrogyria |
| G40.401 |  | Spasms, infantile |
| Q05.9 |  | Spina bifida/Myelomenigocele |
| P11.5 |  | Spinal Cord Injury at birth |
| S14.109A |  | Spinal Cord Injury not at birth, *cervical* spinal cord |
| S34.109A |  | Spinal Cord Injury not at birth, *lumbar* spinal cord |
| S34.139A |  | Spinal Cord Injury not at birth, *sacral* spinal cord |
| S24.109A |  | Spinal Cord Injury not at birth, *thoracic* spinal cord |
| Q05.9 |  | Spinal Lipomeningocele |
| Q85.8 |  | Sturge-Weber disease |
| **CHROMOSOMAL DISORDERS** |
| Q93.4 |  | Cat Cry Syndrome *(Cri-du-Chat)* |
| Q99.9 |  | Chromosomal Anomaly  *(including trisomies, deletions,* d*uplications,*  *translocations,inversions, rings & isochromosome)* *Exceptions: Klinfelter’s Syndrome, Turner Syndrome)* |
| Q87.1 |  | Cornelia De Lange Syndrome |
| Q90.9 |  | Down Syndrome |
| Q99.2 |  | Fragile X syndrome |
| Q87.1 |  | Prader-Willi Syndrome |
| Q93.88 |  | Smith-Magenis Syndrome |
| **METABOLIC DISORDERS** |
| E72.9 |  | Amino acid metabolism disorder |
| E88.1 |  | Congenital lipodystrophy |
| E74.21 | \* | Galactosemia |
| E75.10 |  | Gangliosidosis |
| E74.00 |  | Glycogen storage disease |
| E76.1 |  | Hunter syndrome |
| E83.39 |  | Hypophosphatasia |
| E71.310 |  | LCHAD *(Long Chain Acyl CoA Dehydrogenase*  *Deficiency)* |
| E71.0 | \* | Maple syrup urine disease |
| E83.09 |  | Menkes Syndrome |
| E71.120 |  | Methylmalonic acidemia |
| E76.3 |  | Mucopolysaccharidosis |
| E70.0 | \* | Phenylketonuria *(PKU)* |
| E71.121 |  | Propionic acidemia |
| Q87.1 |  | Sjogren-Larsson Syndrome |
| E75.02 |  | Tay-Sachs disease  |

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| **SKELETAL DISORDERS** |
| M08.00 |  | Arthritis, juvenile rheumatoid |
| Q74.3 |  | Arthrogryposis |
| Q87.40 |  | Beals Syndrome |
| Q74.0 |  | Cleidocranial Dysostosis |
| Q75.0 |  | Craniostenosis |
| Q75.1 |  | Crouzon's Syndrome |
| Q87.0 |  | Mobius sequence |
| Q87.0 |  | Nager-de Reynier Syndrome |
| Q78.0 |  | Osteogenesis imperfecta |
| Q72.90 |  | Proximal Focal Fibula Deficiency *(PFFD w/fibula hemimelia)* |
| Q68.1 | \* | Radial Club Hand |
| **OTHER DISORDERS** |
| Q73.0 |  | Absence of limb |
| Q77.4 |  | Achondroplasia |
| P02.9 |  | ADAM Complex |
| B20 |  | AIDS |
| Q44.7 |  | Alagille Syndrome |
| Q93.5 |  | Angelman Syndrome |
| Q87.0 |  | Apert Syndrome |
| D61.9 |  | Aplastic anemia |
| F84.0 |  | Autism spectrum disorder |
| Q87.3 |  | Beckwith-Wiedemann Syndrome |
| D61.01 |  | Blackfan-Diamond Syndrome |
| Q82.3 |  | Bloch-Sulzberger Syndrome |
| Q92.8 |  | Cat Eye Syndrome |
| Q89.8 |  | CHARGE Syndrome |
| Q37.9 | \* | Cleft lip and palate |
| Q35.9 | \* | Cleft palate |
| Q87.1 |  | Cockayne Syndrome |
| P35.1 |  | Congenital CMV - Cytomegalovirus |
| Q89.4 |  | Conjoined twin |
| E84.9 |  | Cystic fibrosis |
| D82.1 |  | DiGeorge Syndrome |
| Q79.6 |  | Ehlers-Danlos Syndrome |
| D22.9 |  | Epidermal Nevus Syndrome |
| Q81.9 |  | Epidermolysis Bullosa |
| N04.9 |  | Epstein's Syndrome |
| P14.0 |  | Erb's palsy *(Brachial Plexis injury)* |
| R62.51 | \* | Failure to Thrive |
| D61.09 |  | Fanconi Anemia |
| P04.3 |  | Fetal Alcohol effects |
| Q86.0 |  | Fetal Alcohol Syndrome |
| Q79.3 |  | Gastroschisis |
| Q87.0 |  | Goldenhar Syndrome |
| Q67.4 |  | Hemifacial Microsomia |
| P35.2 |  | Herpes, congenital  |
| P83.2 |  | Hydrops-Fetalis |
| E03.1 |  | Hypothyroidism, congenital |
| P57.9 |  | Kernicterus |
| Q76.1 |  | Klippel-Feil Syndrome |
| **OTHER DISORDERS** *(Continued)* |
| Q74.8 |  | Larsen Syndrome |
| G31.82 |  | Leigh's Disease |
| K76.9 |  | Liver disease and/or dysfunction |
| Q78.5 |  | Metaphyseal dysplasia |
| G71.3 |  | Mitochondrial myopathy |
| I67.5 |  | Moyamoya disease |
| G71.0 |  | Muscular dystrophy/Duchenne's  |
|  |  |   *(includes progressive muscular atrophy)* |
| D46.9 |  | Myelodysplasia |
| G71.11 |  | Myotonic dystrophy |
| Q85.00 |  | Neurofibromatosis |
| Q87.1 |  | Noonan Syndrome |
| J38.00 |  | Paralysis, vocal cords |
| Q87.0 | \* | Pierre Robin Syndrome |
| F84.2 |  | Rett Syndrome |
| P35.0 |  | Rubella, congenital |
| Q87.2 |  | Rubinstein-Taybi Syndrome |
| Q87.1 |  | Russell-Silver Syndrome |
| T74.4XXA |  | Shaken Baby Syndrome |
| E78.72 |  | Smith-Lemli-Optiz Syndrome |
| Q87.3 |  | Sotos Syndrome |
| Q89.8 |  | Stickler Syndrome |
| Q68.0 | \* | Torticollis, congenital |
| P37.1 |  | Toxoplasmosis, congenital |
| S06.1X0A |  | Traumatic Head Injury |
| Q75.4 |  | Treacher Collins Syndrome |
| Q85.1 |  | Tuberous Sclerosis |
| Q87.2 |  | VACTER Syndrome |
| Q79.8 |  | Waardenburg-Klein Syndrome |
| Q87.89 |  | Williams Syndrome |
| D82.0 |  | Wiskott-Aldrich Syndrome |