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.

SENSORY CONDITIONS

BLINDNESS

H54.0		Blindness, both eyes
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 *(20/70 best acuity with correction)*
H54.52	*	Low vision, left eye, normal vision right eye
H54.51	*	Low vision, right eye, normal vision left eye

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, duplications, 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