Mississippi First Steps Early Intervention
Qualifying Diagnosed Condition List

The following diagnoses have been determined to meet the definition of *a diagnosed physical or mental condition that has a high probability of resulting in developmental delay* as included in the Mississippi definition of an infant or toddler with a disability according to 34 CFR §303.21.

Please contact the state First Steps Ealy Intervention office if there is a diagnosis not included on this list you feel should be considered for inclusion.

DIAGNOSIS CODE	DESCRIPTION
Q13.1	Absence of iris- Aniridia
Q77.4	Achondroplasia
Z89.439	Acquired absence of unspecified foot
M21.549	Acquired clubfoot, unspecified foot
E22.0	Acromegaly and pituitary gigantism
A85.1	Adenoviral Encephalitis
E71.522	Adrenomyeloneuropathy
G31.81	Alpers disease
Q87.81	Alport syndrome
Q04.1	Arhinencephaly
R27.0	Ataxia, unspecified
Q39.1	Atresia of esophagus with tracheo-esophageal fistula
Q03.1	Atresia of foramina of Magendie and Luschka
Q25.5	Atresia of pulmonary artery
Q21.2	Atrioventricular septal defect
F84.9	Atypical Autism
F84.0	Autistic Disorder
G00.9	Bacterial meningitis, unspecified
D56.1	Beta thalassemia
G54.0	Brachial Plexus Disorders; Thoracic Outlet Syndrome
R01.1	Cardiac murmur, unspecified
I67.1	Cerebral aneurysm, nonruptured
I63.9	Cerebral infarction, unspecified
G80.9	Cerebral palsy, unspecified
Q05.5	Cervical Spina Bifidia without hydrocephalus
E70.330	Chediak Higashi (Steinbrinck) Syndrome
Q30.0	Choanal atresia
Q77.3	Chondrodysplasia Punctata
Q99.9	Chromosomal abnormality, unspecified
C91.10	Chronic lymphocytic leukemia of B-cell type not having achieved remission
E70.0	Classical phenylketonuria
Q36.9	Cleft lip, unilateral
Q35.9	Cleft palate, unspecified
D81.9	Combined immunodeficiency, unspecified
H90.0	Conductive hearing loss, bilateral

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DIAGNOSIS CODE	DESCRIPTION
H90.12	Conductive hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side
H90.11	Conductive hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side
H90.2	Conductive hearing loss, unspecified
H35.53	Cone Dystrophy
Q73.0	Congenital absence of unspecified limb(s)
Q42.3	Congenital absence, atresia and stenosis of anus without fistula
Q41.1	Congenital absence, atresia and stenosis of jejunum
Q16.1	Congenital absence, atresia and stricture of auditory canal (external)
G70.2	Congenital and developmental myasthenia
Q12.0	Congenital cataract
Q04.6	Congenital cerebral cysts
P35.1	Congenital cytomegalovirus infection
Q68.1	Congenital deformity of finger(s) and hand
Q15.0	Congenital glaucoma
P35.2	Congenital herpes viral [herpes simplex] infection
Q03.9	Congenital hydrocephalus, unspecified
E03.1	Congenital hypothyroidism without goiter
P94.2	Congenital hypotonia
Q04.9	Congenital malformation of brain, unspecified
Q24.9	Congenital malformation of heart, unspecified
Q07.9	Congenital malformation of nervous system, unspecified
Q75.9	Congenital malformation of skull and face bones, unspecified
Q87.3	Congenital malformation syndromes involving early overgrowth
Q87.0	Congenital malformation syndromes predominantly affecting facial appearance
Q87.19	Congenital malformation syndromes predominantly associated with short stature
Q87.2	Congenital malformation syndromes predominantly involving limbs
Q89.9	Congenital malformation, unspecified
Q89.1	Congenital malformations of adrenal gland
Q04.0	Congenital malformations of corpus callosum
G71.2	Congenital myopathies
P96.0	Congenital renal failure
P35.0	Congenital rubella syndrome/rubella pneumonitis
A50.9	Congenital syphilis, unspecified
P37.1	Congenital toxoplasmosis
D61.01	Constitutional (pure) red blood cell aplasia
Z77.011	Contact with and (suspected) exposure to lead
P90	Convulsions of newborn
H47.619	Cortical blindness, unspecified side of brain
Q75.0	Craniosynostosis/Acrocephaly/Imperfect fusion of skull/Oxycephaly/Trigonocephaly

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DIAGNOSIS CODE	DESCRIPTION
E84.9	Cystic fibrosis, unspecified
E77.1	Defects in glycoprotein degradation
E77.0	Defects in post-translational modification of lysosomal enzymes
Q93.9	Deletion from autosomes, unspecified
Q93.3	Deletion of short arm of chromosome 4
Q93.4	Deletion of short arm of chromosome 5
D82.1	Di George's syndrome
G90.9	Disorder of the autonomic nervous system, unspecified
E72.3	Disorders of lysine and hydroxylysine metabolism
E74.4	Disorders of pyruvate metabolism and gluconeogenesis
Q20.1	Double outlet right ventricle
Q90.9	Down syndrome, unspecified
E07.1	Dyshormogenetics goiter
Q82.4	Ectodermal dysplasis
Q79.6	Ehlers-Danlos syndrome
Q01.9	Encephalocele, unspecified
Q78.4	Enchondromatosis
A85.0	Enteroviral Encephalitis
G40.823	Epileptic spasms, intractable with status epilepticus
G40.824	Epileptic spasms, intractable without status epilepticus
G40.821	Epileptic spasms, not intractable with status epilepticus
G40.822	Epileptic spasms, not intractable,without status epilepticus
P14.0	Erb's paralysis due to birth injury
P07.22	Extreme immaturity of newborn, gestational age 23 completed weeks
P07.23	Extreme immaturity of newborn, gestational age 24 completed weeks
P07.24	Extreme immaturity of newborn, gestational age 25 completed weeks
P07.25	Extreme immaturity of newborn, gestational age 26 completed weeks
P07.26	Extreme immaturity of newborn, gestational age 27 completed weeks
P07.14	Extremely low birth weight newborn, 1,000 - 1,249 grams
P07.15	Extremely low birth weight newborn, 1,250 -1,499 grams
P07.02	Extremely low birth weight newborn, 500 - 749 grams
P07.03	Extremely low birth weight newborn, 750 - 999 grams
P07.01	Extremely low birth weight newborn, less than 500 grams
E75.21	Fabry (-Anderson) disease
R62.51	Failure to thrive (child)
G90.1	Familial dysautonomia [Riley-Day]
Q97.1	Female with more than three X chromosomes
Q86.0	Fetal alcohol syndrome (dysmorphic)
Q86.1	Fetal hydantoin syndrome
M21.41	Flat foot [pes planus] (acquired), right foot
Q99.2	Fragile X chromosome
E74.21	Galactosemia

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DIAGNOSIS CODE	DESCRIPTION
K21.9	Gastro-esophageal reflux disease without esophagitis
~~Q79.3~~	~~Gastroschisis~~
E75.22	Gaucher disease
Z15.01	Genetic susceptibility to malignant neoplasm of breast
G24.1	Genetic torsion dystonia
E74.00	Glycogen storage disease, unspecified
Z93.1	G-tube Dependence
I51.9	Heart disease, unspecified
G60.0	Hereditary motor and sensory neuropathy
Q04.2	Holoprosencephaly
E72.11	Homocystinuria
E76.01	Hurler's syndrome
Q23.4	Hypoplastic left heart syndrome
P91.60	Hypoxic ischemic encephalopathy [HIE], unspecified
M62.3	Immobility syndrome (paraplegic)
Q82.3	Incontinentia pigmenti
Q00.2	Iniencephaly
P52.21	Intraventricular (nontraumatic) hemorrhage, grade 3, of newborn
P52.22	Intraventricular (nontraumatic) hemorrhage, grade 4, of newborn
E71.110	Isovaleric acidemia
Q97.0	Karyotype 47, XXX
Q98.5	Karyotype 47, XYY
Q76.1	Klippel-Feil syndrome
E75.23	Krabbe disease
H54.8	Legal blindness, as defined in USA
G31.82	Leigh's disease
G40.813	Lennox-Gastaut syndrome, intractable, with status epilepticus
G40.814	Lennox-Gastaut syndrome, intractable, without status epilepticus
G40.811	Lennox-Gastaut syndrome, not intractable, with status epilepticus
G40.812	Lennox-Gastaut syndrome, not intractable, without status epilepticus
E79.1	Lesch-Nyhan syndrome
E78.81	Lipoid dermatoarthritis
I45.81	Long QT syndrome
Q71.42	Longitudinal reduction defect of left radius
Q71.43	Longitudinal reduction defect of radius bilateral
Q71.41	Longitudinal reduction defect of right radius
Q72.40	Longitudinal reduction defect of unspecified femur
Q71.40	Longitudinal reduction defect of unspecified radius
E72.03	Lowe's syndrome
Q75.3	Macrocephaly
Q98.7	Male with sex chromosome mosaicism
Q03.0	Malformations of aqueduct of Sylvius

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DIAGNOSIS CODE	DESCRIPTION
C71.5	Malignant neoplasm of cerebral ventricle
C69.20	Malignant neoplasm of unspecified retina
Q75.4	Mandibulofacial dysostosis
E71.0	Maple-syrup-urine disease
Q87.40	Marfan's syndrome, unspecified
E74.04	McArdle disease
Q04.5	Megalencephaly
E88.41	MELAS syndrome
G03.8	Meningitis due to other unspecified cases
G03.9	Meningitis, unspecified
E88.42	MERRF syndrome
E75.25	Metachromatic leukodystrophy
E71.120	Methylmalonic acidemia
Q02	Microcephaly
Q11.2	Microphthalmos/Cryptophthalmos NOS/Dyplasia of eye/Hypolasis of eye/Rudimentary eye
Q17.2	Microtia
E88.40	Mitochondrial metabolism disorder, unspecified
H90.8	Mixed hearing loss unspecified
H90.72	Mixed hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side
H90.71	Mixed hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side
E76.210	Morquio A mucopolysaccharidoses
E76.211	Morquio B mucopolysaccharidoses
E76.219	Morquio mucopolysaccharidoses, unspecified
Q97.2	Mosaicism, lines with various numbers of X chromosomes
I67.5	Moyamoya disease
E76.1	Mucopolysaccharidosis, type II
E76.3	Mucopolysaccharidosis, unspecified
Q89.7	Multiple congenital malformations, not elsewhere classified
G71.0	Muscular dystrophy
G70.00	Myasthenia gravis without (acute) exacerbation
G71.13	Myotonic chondrodystrophy
G71.11	Myotonic muscular dystrophy
E71.511	Neonatal adrenoleukodystrophy
P91.2	Neonatal cerebral leukomalacia
Q85.01	Neurofibromatosis, Type 1
E75.4	Neuronal Ceroid Lipofuscinosis
P04.3	Newborn (suspected to be) affected by maternal use of alcohol
P04.1	Newborn (suspected to be) affected by other maternal medication
P04.49	Newborn affected by maternal use of other drugs of addiction

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DIAGNOSIS CODE	DESCRIPTION
E75.249	Niemann-Pick disease, unspecified
E72.51	Non-ketotic hyperglycinemia
E70.319	Ocular albinism, unspecified
E70.329	Oculocutaneous albinism, unspecified
Q75.5	Oculomandibular dysostosis
H47.039	Optic nerve hypoplasia, unspecified eye
Q78.0	Osteogenesis imperfecta
Q78.2	Osteopetrosis
H93.299	Other abnormal auditory perceptions, unspecified ear (ANSD)
Q11.1	Other anopthalmos, anopthalmos NOS, agensis of eye, aplasia of eye
G00.8	Other Bacterial Meningitis
I63.8	Other cerebral infarction
Q13.4	Other congenital corneal malformations
Q03.8	Other congenital hydrocephalus
Q87.5	Other congenital malformation syndromes with other skeletal changes
Q79.8	Other congenital malformations of musculoskeletal system
P35.8	Other congenital viral diseases
D61.09	Other constitutional aplastic anemia
Q93.5	Other deletions of part of a chromosome
E77.8	Other disorders of glycoprotein metabolism
E83.39	Other disorders of phosphorus metabolism, unspecified
R13.19	Other dysphagia
G40.803	Other epilepsy, intractable, with status epilepticus
G40.804	Other epilepsy, intractable, without status epilepticus
G40.801	Other epilepsy, not intractable, with status epilepticus
G40.802	Other epilepsy, not intractable, without status epilepticus
E74.09	Other glycogen storage disease
E78.89	Other lipoprotein metabolism disorders
Q93.88	Other microdeletions
E88.49	Other mitochondrial metabolism disorders
E76.29	Other mucopolysaccharidoses
E70.318	Other ocular albinism
E70.328	Other oculocutaneous albinism
Q85.8	Other phakomatoses, not elsewhere classified
Q73.8	Other reduction defects of unspecified limb(s)
Q04.3	Other reduction deformities of brain
Q04.8	Other speciated congenital malformations of brain
M21.80	Other specified acquired deformities of unspecified limb
Q99.8	Other specified chromosome abnormalities
Q65.89	Other specified congenital deformities of hip
Q87.89	Other specified congenital malformation syndromes, not elsewhere classified
Q89.8	Other specified congenital malformations

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DIAGNOSIS CODE	DESCRIPTION
Q74.8	Other specified congenital malformations of limb(s)
Q75.8	Other specified congenital malformations of skull and face bones
Q68.8	Other specified congential musculoskeletal deformities
H83.8X3	Other specified diseases of inner ear, bilateral
E34.8	Other specified endocrine disorders
A88.8	Other specified infections of CNS
E88.89	Other specified metabolic disorders
Q78.8	Other specified osteochondrodysplasias
Q05.9	Other specified postprocedural states
Q97.8	Other specified sex chromosome abnormalities, female phenotype
Q98.8	Other specified sex chromosome abnormalities, male phenotype
Q92.8	Other specified trisomies and partial trisomies of autosomes
A85.8	Other specified viral encephalitis
E75.29	Other sphingolipidosis
A87.8	Other viral meningitis
Q73.1	Phocomelia, unspecified limb(s)
E74.02	Pompe Disease
Q87.11	Prader-Willi syndrome
P07.31	Preterm newborn, gestational age 28 completed weeks
P07.32	Preterm newborn, gestational age 29 completed weeks
P07.33	Preterm newborn, gestational age 30 completed weeks
P07.34	Preterm newborn, gestational age 31 completed weeks
P07.35	Preterm newborn, gestational age 32 completed weeks
E71.121	Propionic acidaemia
F94.1	Reactive attachment disorder of childhood
G60.1	Refsum's disease
H35.149	Retinopathy of prematurity, stage 3, unspecified eye
H35.159	Retinopathy of prematurity, stage 4, unspecified eye
F84.2	Rett's syndrome
C49.9	Rhabdoid tumor
E71.540	Rhizomelic chondrodysplasia punctata
E55.0	Rickets, active
Q13.81	Rieger's anomaly
E76.22	Sanfilippo mucopolysaccharidoses
H90.3	Sensorineural hearing loss, bilateral
H90.42	Sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side
H90.41	Sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side
Q04.4	Septo-optic dysplasia of brain
T74.4XXA	Shaken infant syndrome, initial encounter
D57.40	Sickle-cell thalassemia without crisis

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DIAGNOSIS CODE	DESCRIPTION
E78.72	Smith-Lemli-Opitz syndrome
G81.10	Spastic hemiplegia affecting unspecified side
Q76.0	Spina Bifida occulta
Q05.9	Spina bifida, unspecified
G12.9	Spinal muscular atrophy, unspecified
Q77.7	Spondyloepiphyseal dysplasia
Q25.6	Stenosis of pulmonary artery
M35.9	Systemic involvement of connective tissue, unspecified
Q21.3	Tetralogy of Fallot
Q77.1	Thanatophoric short stature
D69.6	Thrombocytopenia, unspecified
M43.6	Torticollis
S06.0X0A	Traumatic Brain Injury
Q91.7	Trisomy 13, unspecified
Q91.3	Trisomy 18, unspecified
Q85.1	Tuberous sclerosis
Q96.9	Turner's syndrome, unspecified
Q37.9	Unspecified cleft palate with unilateral cleft lip
H91.90	Unspecified hearing loss, unspecified ear
H35.50	Unspecified hereditary retinal dystrophy
S06.9X9S	Unspecified intracranial injury with loss of consciousness of unspecified duration, sequela
H47.20	Unspecified optic atrophy
H33.103	Unspecified retinoschisis, bilateral
H90.5	Unspecified sensorineural hearing loss
Q05.4	Unspecified Spina Bifidia with hydrocephalus
A86	Unspecified viral encephalitis
Q93.81	Velo-cardio-facial syndrome
Z99.11	Ventilator Dependence
Q21.0	Ventricular septal defect
A87.9	Viral meningitis, unspecified
E74.01	Von Gierke disease
Q92.1	Whole chromosome trisomy, mosaicism (mitotic nondisjunction)
Q82.1	Xeroderma pigmentosum
E71.529	X-linked adrenoleukodystrophy, unspecified type
E71.510	Zellweger syndrome