International Classification of Disease, 10th Revision (ICD-10)

Diagnostic Codes Related to Family Infant Toddler (FIT) Program

Revised October 2015

The following relevant diagnosis codes under the four Family Infant Toddler Program eligibility categories are taken directly from the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM).

Diagnostic Codes Related to Family Infant Toddler (FIT) Program

Introduction:

The following is a guide for Family Infant Toddler (FIT) Program providers to determine the whether the child's medical diagnosis meets eligibility under the "Established Condition" or "Medical / Biological Risk", in accordance with the Requirements for Family Infant Toddler Early Intervention Services (NMAC 7.30.8). This guide is also to be used to determine the diagnostic billing codes (ICD-10 International Classification of Diseases) to be entered into FIT-KIDS (Key Information Data System) for any the FIT eligibility criteria.

This guide is categorized by four main categories of Developmental Delay, Established Condition, Medical/Biological Risk Condition, and Environmental Risk Factors. All codes have been converted from the ICD-9 to ICD-10 mapping.

The categories of Established Condition, and Medical/Biological Risk Condition have their own subcategories of the following: Genetic Disorders; Mental/Psychosocial Disorders (DC: 0-3 diagnosis are indicated in bold); Neurologic/Musculoskeletal; Perinatal Factors; Physical Impairments; and Sensory Abnormalities.

Reminder: Eligibility determination in the categories of Medical/Biological Risk Condition and Established Condition shall be made by a license Medical Practitioner.

The category of Environmental Risk Factors have subcategories of the following: Other personal history presenting hazards to health; Family history of certain chronic disabling diseases; family history of certain other specific conditions; family history of other conditions; health supervision of a child; housing, household, and economic circumstances; other family circumstances; and other psychosocial circumstances.

Reminder: Eligibility determination in the category of Environmental Risk Factors shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program.

Any questions regarding conditions not on this guide must be directed to your **FIT Manager** who will work with a New Mexico Department of Health Medical Director/Physician to determine whether a proposed condition will be recognized. Reminder that there are 100,000 more codes in the ICD-10 conversion which makes the mapping process take longer.

Family Infant Toddler (FIT) Providers may keep this document electronically and in print. Note, that the electronic version has the option to find sections by clicking the table of contents (hyperlinks), diagnosis and ICD-10 Codes can also be found by clicking CTRL+F.

Note: for billing purposes through FIT-KIDS all codes that are not listed on the drop-down list may be entered in the Field/Box of "Other ICD-10 Code".

Please contact your FIT Program Manager if you have any questions.

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Developmental Delay

A delay of 25% or more, after correction for prematurity, in one or more of the following areas of development: cognitive; communication; physical/motor; social or emotional; or adaptive of the following:
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Development	F70	Mild Intellectual Disabilities
Development	F71	Moderate Intellectual Disabilities
Development	F72	Severe Intellectual Disabilities
Development	F73	Profound Intellectual Disabilities
Development	F79	Unspecified Intellectual Disabilities
Development	F80.1	Expressive Language Disorder
Development	F80.2	Mixed Receptive-Expressive Language Disorder
Development	F80.81	Childhood Onset Fluency Disorder
Development	F80.89	Other developmental disorders of speech and language
Development	F82	Specific Developmental Disorder of Motor Functions (Mixed Development Disorder)
Development	F84.0	Autistic Disorder **(DC 0-3 R: 710 Multisystem Development Disorder)**
Development	F84.9	Pervasive Development Disorder (PDD), Unspecified
Development	F88	Other disorders of psychological development Mixed Development Disorder
Development	F89	Other disorders of psychological development Other Specified Delays in Development

ESTABLISHED CONDITION: Genetic Disorders

A diagnosed physical, mental, or neurobiological condition that has a high probability of resulting in developmental delay. The established condition shall be diagnosed by a health care provider and documentation shall be kept on file. Established conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	A81.1	Subacute Sclerosing Panencephalitis
Genetic Disorders	D33.2	Benign Neoplasm of Brain, Unspecified
Genetic Disorders	E70.20	Disorder of tyrosine metabolism, unspecified
Genetic Disorders	E70.21	Tyrosinemia, Type I (Tyrosinosis) and Type II (Richner-Hanhart Syndrome)
Genetic Disorders	E70.29	Disorder of tyrosine metabolism, unspecified
Genetic Disorders	E70.8	Other disorders of aromatic amino-acid metabolism (Waardenburg's Syndrome, Type I)
Genetic Disorders	E70.9	Disorder of aromatic amino-acid metabolism, unspecified
Genetic Disorders	E72.03	Lowe's Syndrome or (Oculocerebrorenal Syndrome)
Genetic Disorders	E75.23	Krabbe's Disease
Genetic Disorders	E75.29	Other sphingolipidosis (Canavan Disease)
Genetic Disorders	E75.3	Sphingolipidosis, unspecified
Genetic Disorders	E760.1	Hurler Syndrome
Genetic Disorders	E760.2	Hurler-Scheie syndrome
Genetic Disorders	E760.3	Scheie's syndrome
Genetic Disorders	E77.1	Defects in Glycoprotein Degradation (Mannosidosis)
Genetic Disorders	E77.8	Other disorders of glycoprotein metabolism
Genetic Disorders	E77.9	Disorder of glycoprotein metabolism, unspecified
Genetic Disorders	E79.1	Lesch-Nyhan Syndrome
Genetic Disorders	E83.00	Disorder of copper metabolism, unspecified Menkes' Syndrome (Kinky Hair Disease)
Genetic Disorders	E83.01	Wilson's Disease
Genetic Disorders	F84.2	Rett's Syndrome
Genetic Disorders	G31.81	Alper's Disease

ESTABLISHED CONDITION: Genetic Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	G31.82	Leigh's Disease
Genetic Disorders	G70.00	Myasthenia Gravis, without (acute) exacerbation (Familial Infantile)
Genetic Disorders	G71.2	Congenital Myopathies
Genetic Disorders	G91.1	Obstructive Hydrocephalus
Genetic Disorders	M21.511	Acquired clawhand, right hand
Genetic Disorders	M21.512	Acquired clawhand, left hand
Genetic Disorders	M21.519	Acquired clawhand, unspecified hand
Genetic Disorders	M21.521	Acquired clubhand, right hand
Genetic Disorders	M21.522	Acquired clubhand, left hand
Genetic Disorders	M21.529	Acquired clubhand, unspecified hand
Genetic Disorders	M21.531	Acquired clawfoot, right foot
Genetic Disorders	M21.532	Acquired clawfoot, left foot
Genetic Disorders	M21.539	Acquired clawfoot, unspecified foot
Genetic Disorders	M21.541	Acquired Clubfoot, Right Foot
Genetic Disorders	M21.542	Acquired Clubfoot, Left Foot
Genetic Disorders	M21.549	Acquired Clubfoot, Unspecified Foot (Paralytic Clubbed Foot)
Genetic Disorders	M21.511	Acquired clawhand, right hand
Genetic Disorders	Q03.1	Atresia of foramina of Magendie and Luschka Dandy Walker Syndrome
Genetic Disorders	Q03.9	Congenital hydrocephalus, unspecified
Genetic Disorders	Q04.0	Congenital malformations of corpus callosum
Genetic Disorders	Q04.3	Other Reduction Deformities of Brain (Polymicrogyria)
Genetic Disorders	Q72.10	Congenital Absence of Unspecified limbs
Genetic Disorders	Q73.1	Phocomelia, unspecified limb(s)
Genetic Disorders	Q73.2	Other Reduction Defects of Unspecified Limb(s)
Genetic Disorders	Q75.9	Congenital Anomalies of Skull and Face Bone, unspecified (Goldenhar)
Genetic Disorders	Q87.89	Spondyloepiphyseal dysplasia Dyggve-Melchior-Clausen Syndrome (D-M-C Dwarfism)
Genetic Disorders	Q78.0	Osteogenesis Imperfecta

ESTABLISHED CONDITION: Genetic Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	Q80.8	Other congenital ichthyosis Sjogren-Larsson Syndrome
Genetic Disorders	Q85.1	Tuberous Sclerosis
Genetic Disorders	Q85.8	Other Phakomatosis, not elsewhere classified (Sturge-Weber Syndrome)
Genetic Disorders	E78.72	Congenital malformation syndromes predominantly associated with short stature: (Prader-Willi Syndrome)
Genetic Disorders	Q87.89	Congenital malformation syndromes predominantly associated with short stature: (DeLange's Syndrome)
Genetic Disorders	Q87.89	Congenital malformation syndromes predominantly associated with short stature: (Dubowitz Syndrome)
Genetic Disorders	Q87.1	Congenital malformation syndromes predominantly associated with short stature: (Smith-Lemli-Opitz Syndrome)
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs: (Caudal Regression Syndrome-Sirenomelia)
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs: (Rubinstein-Taybi Syndrome)
Genetic Disorders	Q87.89	Other specified congenital malformation syndromes, not elsewhere classified Zellweger/Cerebrohepatorenal Syndrome
Genetic Disorders	Q90.0	Trisomy 21, nonmosaicism (meiotic nondisjunction)
Genetic Disorders	Q90.1	Trisomy 21, mosaicism (mitotic nondisjunction)
Genetic Disorders	Q90.2	Trisomy 21, translocation
Genetic Disorders	Q90.9	Down Syndrome, Unspecified
Genetic Disorders	Q91.0	Trisomy 18, nonmosaicism (meiotic nondisjunction)
Genetic Disorders	Q91.1	Trisomy 18, mosaicism (mitotic nondisjunction)
Genetic Disorders	Q91.2	Trisomy 18, translocation
Genetic Disorders	Q91.3	Trisomy 18, Unspecified (Edward's, E3)
Genetic Disorders	Q91.4	Trisomy 13, nonmosaicism (meiotic nondisjunction)
Genetic Disorders	Q91.5	Trisomy 13, mosaicism (mitotic nondisjunction)

ESTABLISHED CONDITION: Genetic Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	Q91.6	Trisomy 13, translocation
Genetic Disorders	Q91.7	Trisomy 13, Unspecified (Patau's, D1)
Genetic Disorders	Q93.4	Deletion of short arm of chromosome 5(Cri Du Chat or Cat's Cry)
Genetic Disorders	Q93.5	Other deletions of part of a chromosome (Angelman's Syndrome)
Genetic Disorders	Q99.9	Other deletions from the autosomes Williams Syndrome
Genetic Disorders	Q99.2	Fragile X Syndrome For Boys
Genetic Disorders	R47.01	Aphasia

ESTABLISHED CONDITION: Neurologic/Musculoskeletal

A diagnosed physical, mental, or neurobiological condition that has a high probability of resulting in developmental delay. The established condition shall be diagnosed by a health care provider and documentation shall be kept on file. Established conditions include the following:
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	A80.39	Paralysis, Other Specified
Neurologic/Musculoskeletal	G06.2	Extradural and Subdural Abscess, Unspecified (Hemiplegia (Hemiparesis)
Neurologic/Musculoskeletal	G11.9	Hereditary Ataxia, Unspecified (Spinocerebellar Disorders)
Neurologic/Musculoskeletal	G24.1	Genetic Torsion Dystonia (Dystonia Musculorum Deformans)
Neurologic/Musculoskeletal	G25.3	Myoclonus
Neurologic/Musculoskeletal	G31.89	Other Specified Degenerative Diseases of Nervous System
Neurologic/Musculoskeletal	G60.9	Hereditary and Idiopathic Neuropathy, Unspecified
Neurologic/Musculoskeletal	G71.0	Muscular Dystrophy
Neurologic/Musculoskeletal	G71.11	Myotonic Muscular Dystrophy
Neurologic/Musculoskeletal	G71.12	Myotonia Congenita (Thomsen's Disease)
Neurologic/Musculoskeletal	G71.2	Congenital Myopathies
Neurologic/Musculoskeletal	G80.0	Spastic quadriplegic cerebral palsy
Neurologic/Musculoskeletal	G80.1	Spastic Diplegic Cerebral Palsy (Congenital Diplegia, Paraplegia)
Neurologic/Musculoskeletal	G80.2	Spastic hemiplegic cerebral palsy (Congenital Hemiplegia) (Infantile Postnatal NOS) (Spastic Infantile Paralysis)
Neurologic/Musculoskeletal	G80.3	Athetoid cerebral palsy
Neurologic/Musculoskeletal	G80.4	Ataxic cerebral palsy
Neurologic/Musculoskeletal	G80.8	Other cerebral palsy
Neurologic/Musculoskeltal	G80.9	Cerebral palsy, unspecified
Neurologic/Musculoskeletal	G82.20	Paraplegia, unspecified Paralysis of Both Lower Limbs (Paraplegia)
Neurologic/Musculoskeletal	G82.50	Quadriplegia, unspecified Paralysis of All Four Limbs (Quadriplegia, Quadriparesis,)
Neurologic/Musculoskeletal	G83.0	Diplegia of upper limbs Paralysis of Both Upper Limbs (Diplegia)

ESTABLISHED CONDITION: Neurologic/Musculoskeletal (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	G83.10	Monoplegia of lower limb affecting unspecified side Paralysis of One Lower Limb (Monoplegia)
Neurologic/Musculoskeletal	G83.20	Monoplegia of upper limb affecting unspecified side Paralysis of One Upper Limb (Monoplegia)
Neurologic/Musculoskeletal	G83.9	Paralytic syndrome, unspecified
Neurologic/Musculoskeletal	G93.0	Cerebral Cysts
Neurologic/Musculoskeletal	G93.5	Compression of Brain
Neurologic/Musculoskeletal	G93.40	Encephalopathy, unspecified
Neurologic/Musculoskeletal	G93.49	Other encephalopathy
Neurologic/Musculoskeletal	P11.5	Birth injury to spine and spinal cord
Neurologic/Musculoskeletal	P90	Convulsions in newborn
Neurologic/Musculoskeletal	P91.60	Hypoxic ischemic encephalopathy [HIE], unspecified
Neurologic/Musculoskeletal	P94.2	Congenital hypotonia
Neurologic/Musculoskeletal	Q00.0	Anencephaly
Neurologic/Musculoskeletal	Q01.9	Encephalocele, Unspecified
Neurologic/Musculoskeletal	Q02	Microcephaly
Neurologic/Musculoskeletal	Q04.2	Holoprosencephaly
Neurologic/Musculoskeletal	Q04.3	Other reduction deformities of brain Lissencephaly
Neurologic/Musculoskeletal	Q04.6	Congenital cerebral cysts Cortical Dysplasia
Neurologic/Musculoskeletal	Q05.8	Sacral spina bifida without hydrocephalus (Meningomyelocele, Meningocele)
Neurologic/Musculoskeletal	Q05.9	Spina Bifida, Unspecified
Neurologic/Musculoskeletal	Q07.00	Sacral Spina Bifida Without Hydrocephalus
Neurologic/Musculoskeletal	Q07.01	Arnold-Chiari Syndrome with Spina Bifida
Neurologic/Musculoskeletal	Q07.03	Arnold-Chiari syndrome with spina bifida and hydrocephalus
Neurologic/Musculoskeletal	Q76.49	Other congenital malformations of spine, not associated with scoliosis (Sacral Agenesis)
Neurologic/Musculoskeletal	Please refer to your medical professional	Traumatic Brain Injury

ESTABLISHED CONDITION: **Perinatal Factors**

A diagnosed physical, mental, or neurobiological condition that has a high probability of resulting in developmental delay. The established condition shall be diagnosed by a health care provider and documentation shall be kept on file. Established conditions include the following:
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Perinatal Factors	P07.22	Extreme immaturity of newborn, gestational age 23 completed weeks
Perinatal Factors	P07.23	Extreme immaturity of newborn, gestational age 24 completed weeks
Perinatal Factors	P07.24	Extreme immaturity of newborn, gestational age 25 completed weeks
Perinatal Factors	P07.25	Extreme immaturity of newborn, gestational age 26 completed weeks
Perinatal Factors	P07.26	Extreme immaturity of newborn, gestational age 27 completed weeks
Perinatal Factors	P07.31	Preterm newborn, gestational age 28 completed weeks
Perinatal Factors	P35.0	Congenital rubella syndrome
Perinatal Factors	P39.9	Infection specific to the perinatal period, unspecified
Perinatal Factors	P84	Other Problems with Newborns (birth asphyxia NOS, Severe)
Perinatal Factors	P91.0	Neonatal cerebral ischemia
Perinatal Factors	P91.1	Acquired periventricular cysts of newborn
Perinatal Factors	P91.2	Neonatal cerebral leukomalacia
Perinatal Factors	P91.4	Neonatal Cerebral Depression
Perinatal Factors	P91.5	Neonatal Coma
Perinatal Factors	P96.1	Neonatal withdrawal symptoms from maternal use of drugs of addiction

ESTABLISHED CONDITION: **Physical Impairments**

A diagnosed physical, mental, or neurobiological condition that has a high probability of resulting in developmental delay. The established condition shall be diagnosed by a health care provider and documentation shall be kept on file. Established conditions include the following:
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Physical Impairments	C69.01	Malignant neoplasm of right conjunctiva (Retinoblastoma)
Physical Impairments	C69.02	Malignant neoplasm of left conjunctiva (Retinoblastoma)
Physical Impairments	H33.001	Unspecified retinal detachment with retinal break, right eye
Physical Impairments	Q11.1	Other Anophthalmos
Physical Impairments	Q11.2	Microphthalmia
Physical Impairments	Q13.1	Absence of Iris (Aniridia)
Physical Impairments	Q13.89	Other congenital malformations of anterior segment of eye
Physical Impairments	Q16.1	Congenital absence, atresia and stricture of auditory canal (external)
Physical Impairments	Q17.2	Microtia
Physical Impairments	Q17.8	Other specified congenital anomalies of ear
Physical Impairments	Q67.8	Other congenital deformities of chest
Physical Impairments	Q74.9	Unspecified Congenital Malformation of Limb(s)

ESTABLISHED CONDITION: **Sensory Abnormalities**

A diagnosed physical, mental, or neurobiological condition that has a high probability of resulting in developmental delay. The established condition shall be diagnosed by a health care provider and documentation shall be kept on file. Established conditions include the following:
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Sensory Abnormalities	H26.009	Unspecified Infantile and Juvenile Cataract, Unspecified Eye
Sensory Abnormalities	H27.03	Aphakia Bilateral
Sensory Abnormalities	H27.03	Aphakia. Bilateral
Sensory Abnormalities	H33.009	Unspecified Retinal Detachment with Retinal Break, Unspecified eye
Sensory Abnormalities	H35.029	Exudative Retinopathy, Unspecified Eye
Sensory Abnormalities	H35.119	Retinopathy of Prematurity, Stage 0, Unspecified Eye
Sensory Abnormalities	H35.129	Retinopathy of Prematurity, Stage 1, Unspecified Eye
Sensory Abnormalities	H35.139	Retinopathy of Prematurity, Stage 2, Unspecified Eye
Sensory Abnormalities	H35.149	Retinopathy of Prematurity, Stage 3, Unspecified Eye
Sensory Abnormalities	H35.159	Retinopathy of Prematurity, Stage 4, Unspecified Eye
Sensory Abnormalities	H35.169	Retinopathy of Prematurity, Stage 5, Unspecified Eye
Sensory Abnormalities	H35.179	Retrolental Fibroplasia, Unspecified Eye (ROP)
Sensory Abnormalities	H35.52	Pignentary Retinal Distrophy (RP)
Sensory Abnormalities	H35.54	Dystrophies Primarily Involving the Retinal Pigment Epithelium/Leber's Congenital Amaurosis
Sensory Abnormalities	H47.039	Optic Nerve Hypoplasia, Unspecified Eye
Sensory Abnormalities	H47.20	Unspecified Optic Atrophy
Sensory Abnormalities	H47.619	Cortical Blindness, Unspecified Side of the Brain (Disorders of visual cortex)
Sensory Abnormalities	H53.40	Unspecified Visual Field Defects
Sensory Abnormalities	H53.469	Homonymous Bilateral Field Defects, Unspecified Side

ESTABLISHED CONDITION: Sensory Abnormalities (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Sensory Abnormalities	H53.47	Heteronymous Bilateral Field Defects
Sensory Abnormalities	H53.51	Achromatopsia
Sensory Abnormalities	H54.7	Cortical Visual Impairment (CVI) (Unspecified Visual Loss)
Sensory Abnormalities	H54.8	Blindness
Sensory Abnormalities	H55.00	Unspecified nystagmus
Sensory Abnormalities	H55.01	Congenital nystagmus
Sensory Abnormalities	H55.09	Other Forms of Nystagmus
Sensory Abnormalities	H90.3	Sensorineural Hearing Loss, Bilateral
Sensory Abnormalities	H90.41	Sensorineural Hearing Loss, Unilateral, Right Ear, with Unrestricted Hearing on the Contralateral Side
Sensory Abnormalities	H90.42	Sensorineural Hearing Loss, Unilateral, Left Ear, with Unrestricted Hearing on the Contralateral Side
Sensory Abnormalities	H90.5	Unspecified Sensorineural Hearing Loss
Sensory Abnormalities	H90.8	Mixed Conductive and Sensorineural Hearing Loss, Unspecified
Sensory Abnormalities	H90.6	Mixed conductive and sensorineural hearing loss, bilateral
Sensory Abnormalities	H90.71	Mixed conductive and sensorineural hearing loss, unilateral, right ear, with unrestricted hearing on the contralateral side
Sensory Abnormalities	H90.72	Mixed conductive and sensorineural hearing loss, unilateral, left ear, with unrestricted hearing on the contralateral side
Sensory Abnormalities	H91.90	Unspecified Hearing Loss, Unspecified Ear
Sensory Abnormalities	H91.91	Unspecified hearing loss, right ear
Sensory Abnormalities	H91.92	Unspecified hearing loss, left ear

ESTABLISHED CONDITION: Sensory Abnormalities (continued)

Sensory Abnormalities	H91.93	Unspecified hearing loss, bilateral
Sensory Abnormalities	Q15.0	Congenital Glaucoma
Sensory Abnormalities	Q16.9	Congenital Malformations of Ear Causing Impairment of hearing, Unspecified
Sensory Abnormalities	Q89.8	Other Specified Congenital Malformations

MEDICAL/BIOLOGICAL RISK CONDITION: Genetic Disorders

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

***Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.**

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	D82.1	Di George's Syndrome
Genetic Disorders	E00.9	Congenital iodine-deficiency syndrome, unspecified
Genetic Disorders	E70.0	Classical phenylketonuria
Genetic Disorders	E70.30	Albinism, unspecified
Genetic Disorders	E71.0	Maple Syrup Urine Disease
Genetic Disorders	E71.110	Isovaleric Acidemia
Genetic Disorders	E71.121	Methylmalonic Aciduria
Genetic Disorders	E71.522	Adrenomyeloneuropathy
Genetic Disorders	E72.09	Other disorders of amino-acid transport (Fanconi (-de Toni) (-Debre) Syndrome)
Genetic Disorders	E72.11	Homocystinuria
Genetic Disorders	E72.22	Argininosuccinic Aciduria
Genetic Disorders	E72.51	Non-ketotic hyperglycinemia
Genetic Disorders	E74.02	Pompe's Disease
Genetic Disorders	E74.21	Galactosemia
Genetic Disorders	E75.02	Tay-Sachs disease
Genetic Disorders	E77.0	Defects in Post Translational Modification of Lysosomal Enzymes

MEDICAL/BIOLOGICAL RISK CONDITION: Genetic Disorders (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	E75.19	Other gangliosidosis
Genetic Disorders	E75.22	Gaucher's Disease
Genetic Disorders	E75.240	Niemann-Pick disease type A
Genetic Disorders	E75.25	Metachromatic Leukodystrophy
Genetic Disorders	E75.4	Neuronal Ceroid Lipofuscinoses
Genetic Disorders	E75.01	Sandhoff disease
Genetic Disorders	F19.939	Other psychoactive substance use, unspecified with withdrawal, unspecified
Genetic Disorders	G11.3	Cerebellar ataxia with defective DNA repair
Genetic Disorders	G12.0	Infantile Spinal Muscular Atrophy Werdnig-Hoffman Disease,
Genetic Disorders	G60.9	Hypertrophic Interstitial Neuritis
Genetic Disorders	G90.1	Familial Dysautonomia (Riley-Day Syndrome, HSAN III)
Genetic Disorders	I67.9	Cerebrovascular Disease, Unspecified
Genetic Disorders	Q79.6	Cutis laxa senilis
Genetic Disorders	L94.0	Linear scleroderma
Genetic Disorders	P02.8	Newborn (suspected to be) affected by other abnormalities of membranes
Genetic Disorders	P35.1	Congenital cytomegalovirus infection
Genetic Disorders	P35.1	Congenital cytomegalovirus infection

MEDICAL/BIOLOGICAL RISK CONDITION: Genetic Disorders (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	G93.89	Other Conditions of the Brain
Genetic Disorders	Q06.2	Diastematomyelia
Genetic Disorders	Q26.2	Total anomalous pulmonary venous connection
Genetic Disorders	Q87.40	Other specified congenital musculoskeletal deformities (Beals-Hecht Syndrome)
Genetic Disorders	Q74.8	Other specified congenital malformations of limb(s) Larsen's Syndrome
Genetic Disorders	Q77.0	Achondrogenesis
Genetic Disorders	Q77.4	Achondroplasia
Genetic Disorders	Q77.5	Diastrophic Dysplasia
Genetic Disorders	Q79.9	Other osteochondrodysplasia with defects of growth of tubular bones and spine Kniest's Syndrome (Metatrophic Dwarfism II)
Genetic Disorders	Q78.8	Other Specified Osteochondrodysplasia (Acrodysplasia I)
Genetic Disorders	Q78.9	Osteochondrodysplasia, unspecified
Genetic Disorders	Q79.6	Ehlers-Danlos Syndrome
Genetic Disorders	Q80.9	Congenita Ichthyosis, Unspecified (Harlequin Fetus)
Genetic Disorders	Q82.1	Xeroderma Pigmentosum
Genetic Disorders	Q85.01	Other Neurofibromatosis
Genetic Disorders	Q87.0	Congenital malformation syndromes predominantly affecting facial appearance Apert's Syndrome (Acrocephalosyndactyly I, ACS I)
Genetic Disorders	Q87.0	Congenital malformation syndromes predominantly affecting facial appearance Saethre-Chotzen Syndrome (Chotzen's, Acrocephalosyndactyly III)

MEDICAL/BIOLOGICAL RISK CONDITION: Genetic Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	Q87.0	Congenital malformation syndromes predominantly affecting facial appearance Robin's Syndrome
Genetic Disorders	Q87.5	Congenital malformation syndromes predominantly affecting facial appearance Oral-Facial-Digital Syndrome
Genetic Disorders	Q87.89	Congenital malformation syndromes predominantly associated with short stature Noonan's Syndrome
Genetic Disorders	Q87.1	Congenital malformation syndromes predominantly associated with short stature Russell (-Silver) Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs Holt-Oram (Cardiac-Limb) Syndrome (Atriodigital Dysplasia) (Klippel-Trenaunay-Weber Syndrome) Otopalatodigital Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs Klippel-Trenaunay-Weber Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs Otopalatodigital Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs Otopalatodigital Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs VATER Syndrome
Genetic Disorders	Q87.2	Congenital malformation syndromes predominantly involving limbs VATER Syndrome
Genetic Disorders	Q87.3	Congenital malformation syndromes involving early overgrowth Cerebral Gigantism (Soto's Syndrome)
Genetic Disorders	Q87.40	Marfan's Syndrome, Unspecified
Genetic Disorders	Q87.89	Other specified congenital malformation syndromes, not elsewhere classified Langer-Giedion Syndrome (Acrodysplasia V, Klingmuller's)
Genetic Disorders	Q89.8	Other Specified Congenital Malformations CHARGE Syndrome (CHARGE Association)

MEDICAL/BIOLOGICAL RISK CONDITION: Genetic Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Genetic Disorders	Q92.8	Other specified trisomies and partial trisomies of autosomes
Genetic Disorders	Q93.7	Deletions with other complex rearrangements
Genetic Disorders	Q93.81	Velo-cardio-facial Syndrome
Genetic Disorders	Q96.9	Turner's syndrome, unspecified
Genetic Disorders	Q97.8	Other specified sex chromosome abnormalities, female phenotype
Genetic Disorders	Q98.4	Klinefelter syndrome, unspecified
Genetic Disorders	Q98.8	Other specified sex chromosome abnormalities, male phenotype
Genetic Disorders	Q99.9	Chromosomal Abnormality, Unspecified, (Coffin-Lowry Syndrome)
Genetic Disorders	R62.7	Failure to thrive (child)

MEDICAL/BIOLOGICAL RISK CONDITION: Mental/Psychosocial Disorders

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

***Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.**

Condition Type	ICD-10 Code	Condition/Diagnosis
Mental/Psychosocial Disorders	F32.0	Major Depressive Affective Disorder Single Episode Unspecified **(DC 0-3 R: 231 Type I Major Depression)**
Mental/Psychosocial Disorders	F39	Episodic Mood Disorder Unspecified **(DC 0-3 R: 231 Type II Depressive Disorder)**
Mental/Psychosocial Disorders	F40.10	Social Phobia, Unspecified **(DC 0-3 R: 220's Anxiety Disorder Disorder)**
Mental/Psychosocial Disorders	F40.8	Other Phobic Anxiety Disorders **(DC 0-3 R: 220's Anxiety Disorder Disorder)**
Mental/Psychosocial Disorders	F41.1	Generalized Anxiety Disorder **(DC 0-3 R: 220's Anxiety Disorder Disorder)**
Mental/Psychosocial Disorders	F41.9	Anxiety Disorder, unspecified **(DC 0-3 R: 225 Anxiety Disorder NOS)**
Mental/Psychosocial Disorders	F43.10	Post-traumatic stress disorder, unspecified (DC 0-3 R: 100 Post Traumatic Stress Disorder)
Mental/Psychosocial Disorders	F43.20	Adjustment disorder, unspecified (DC 0-3 R: 300 Adjustment Disorder)
Mental/Psychosocial Disorders	F43.21	Adjustment disorder with depressed mood (DC 0-3 R: 210 Prolonged Bereavement/Grief Reaction)
Mental/Psychosocial Disorders	F43.22	Adjustment disorder with anxiety (DC 0-3 R: 300 Adjustment Disorder)
Mental/Psychosocial Disorders	F43.23	Adjustment disorder with mixed anxiety and depressed mood (DC 0-3 R: 300 Adjustment Disorder)
Mental/Psychosocial Disorders	F43.24	Adjustment disorder with disturbance of conduct (DC 0-3 R: 400's Regulation Disorder of Sensory Processing)

MEDICAL/BIOLOGICAL RISK CONDITION: Mental/Psychosocial Disorders (Continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Mental/Psychosocial Disorders	F43.25	Adjustment disorder with mixed disturbance of emotions and conduct (DC 0-3 R: 150 Deprivation /Maltreatment Disorder or DC 0-3 R: 900 Relationship Disorder If PIR-GAS of 40 or below)
Mental/Psychosocial Disorders	F50.00	Anorexia Nervosa, Unspecified **(DC 0-3 R: 600's Feeding Behavior Disorder)**
Mental/Psychosocial Disorders	F50.2	Bulimia Nervosa (DC 0-3 R: 600's Feeding Behavior Disorder)
Mental/Psychosocial Disorders	F50.9	Eating disorder, unspecified (DC 0-3 R: 600's Feeding Behavior Disorder)
Mental/Psychosocial Disorders	F51.02	Adjustiment Insominia **(DC 0-3 R: 500's Sleep Behavior Disorder)**
Mental/Psychosocial Disorders	F51.09	Other insomnia not due to a substance or known physiological condition **(DC 0-3 R: 500's Sleep Behavior Disorder)**
Mental/Psychosocial Disorders	F84.8	Other Specified Pervasive Developmental Disorder **(DC 0-3 R: 710 Multisystem Development Disorder)**
Mental/Psychosocial Disorders	F93.0	Separation anxiety disorder of childhood (DC 0-3 R: 220's Anxiety Disorder Disorder)
Mental/Psychosocial Disorders	F93.9	Childhood emotional disorder, unspecified (DC 0-3 R: 240 Mixed Disorder of Emotional Expressiveness)
Mental/Psychosocial Disorders	F43.29	Other childhood disorders of social functioning (DC 0-3 R: 300 Adjustment Disorder)
Mental/Psychosocial Disorders	F98.21	Rumination Disorder of Infancy (DC 0-3 R: 600's Feeding Behavior Disorder)
Mental/Psychosocial Disorders	F98.29	Other feeding disorders of infancy and early childhood (DC 0-3 R: 600's Feeding Behavior Disorder)
Mental/Psychosocial Disorders	F98.3	PICA of Infancy and Childhood (DC 0-3 R: 600's Feeding Behavior Disorder)

Diagnostic Codes Related to Family Infant Toddler (FIT) Program

MEDICAL/BIOLOGICAL RISK CONDITION: Neurologic/Musculoskeletal

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

***Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.**

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	A87.9	Viral meningitis, unspecified
Neurologic/Musculoskeletal	C72.9	Malignant neoplasm of central nervous system, unspecified
Neurologic/Musculoskeletal	G03.9	Meningitis, Unspecified
Neurologic/Musculoskeletal	G40.001	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.009	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.011	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.019	Localization-related (focal) (partial) idiopathic epilepsy and epileptic syndromes with seizures of localized onset, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.111	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, not intractable, with status epilepticus

MEDICAL/BIOLOGICAL RISK CONDITION: Neurologic/Musculoskeletal (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	G40.109	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.111	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with simple partial seizures, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.011	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.009	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.019	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.119	Localization-related (focal) (partial) symptomatic epilepsy and epileptic syndromes with complex partial seizures, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.301	Generalized idiopathic epilepsy and epileptic syndromes, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.309	Generalized idiopathic epilepsy and epileptic syndromes, not intractable, without status epilepticus

MEDICAL/BIOLOGICAL RISK CONDITION: Neurologic/Musculoskeletal (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	G40.A11	Generalized idiopathic epilepsy and epileptic syndromes, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.A19	Generalized idiopathic epilepsy and epileptic syndromes, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.821	Other generalized epilepsy and epileptic syndromes, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.822	Other generalized epilepsy and epileptic syndromes, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.823	Other generalized epilepsy and epileptic syndromes, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.824	Other generalized epilepsy and epileptic syndromes, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.801	Other epilepsy, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.802	Other epilepsy, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.803	Other epilepsy, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.804	Other epilepsy, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.812	Lennox-Gastaut Syndrome (Gastaut's Syndrome)
Neurologic/Musculoskeletal	G40.901	Epilepsy, unspecified, not intractable, with status epilepticus

MEDICAL/BIOLOGICAL RISK CONDITION: Neurologic/Musculoskeletal (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	G40.909	Epilepsy, unspecified, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.911	Epilepsy, unspecified, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.919	Epilepsy, unspecified, intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.B01	Juvenile myoclonic epilepsy, not intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.B09	Juvenile myoclonic epilepsy, not intractable, without status epilepticus
Neurologic/Musculoskeletal	G40.B11	Juvenile myoclonic epilepsy, intractable, with status epilepticus
Neurologic/Musculoskeletal	G40.B19	Juvenile myoclonic epilepsy, intractable, without status epilepticus
Neurologic/Musculoskeletal	G72.9	Myopathy Unspecified
Neurologic/Musculoskeletal	G83.89	Other Specified Paralytic Syndromes
Neurologic/Musculoskeletal	G95.0	Syringomyelia and syringobulbia
Neurologic/Musculoskeletal	G95.89	Other specified diseases of spinal cord
Neurologic/Musculoskeletal	H55.89	Other irregular eye movements (Kinsbourne Syndrome)
Neurologic/Musculoskeletal	I61.9	Nontraumatic intracerebral hemorrhage, unspecified
Neurologic/Musculoskeletal	P14.0	Erb's Paralysis due to Birth Injury
Neurologic/Musculoskeletal	Q03.8	Other congenital hydrocephalus
Neurologic/Musculoskeletal	Q04.6	Congenital cerebral cysts
Neurologic/Musculoskeletal	Q04.8	Other Specified Congenital malformations of brain
Neurologic/Musculoskeletal	R56.9	Unspecified convulsions

MEDICAL/BIOLOGICAL RISK CONDITION: Neurologic/Musculoskeletal (continued)

Condition Type	ICD-10 Code	Condition/Diagnosis
Neurologic/Musculoskeletal	S14.3XXA	Injury of Brachial Plexus, Initial Encounter
Neurologic/Musculoskeletal	S14.3XXD	Injury of Brachial Plexus, Subsequent Encounter
Neurologic/Musculoskeletal	S14.3XXS	Injury of Brachial Plexus, Sequela Encounter
Neurologic/Musculoskeletal	T56.0X1A	Toxic Effects of Lead and Its Compounds, accidental (Sequela), Initial Encounter
Neurologic/Musculoskeletal	T56.0X1A	Toxic Effects of Lead and Its Compounds, accidental (unintentional), Initial Encounter
Neurologic/Musculoskeletal	T74.4XXA	Shaken infant syndrome, initial encounter
Neurologic/Musculoskeletal	T74.4XXD	Shaken infant syndrome, subsequent encounter
Neurologic/Musculoskeletal	T74.4XXS	Shaken infant syndrome, sequela

MEDICAL/BIOLOGICAL RISK CONDITION: **Perinatal Factors**

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

***Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.**

Condition Type	ICD-10 Code	Condition/Diagnosis
Perinatal Factors	A33	Tetanus neonatorum
Perinatal Factors	E80.7	Disorder of bilirubin metabolism, unspecified
Perinatal Factors	O30.109	Triplet pregnancy, unspecified number of placenta and unspecified number of amniotic sacs, unspecified trimester
Perinatal Factors	O30.129	Triplet pregnancy with two or more monoamniotic fetuses, unspecified trimester
Perinatal Factors	O30.199	Triplet pregnancy, unable to determine number of placenta and number of amniotic sacs, unspecified trimester
Perinatal Factors	P00.9	Newborn (suspected to be) affected by unspecified maternal condition
Perinatal Factors	P04.3	Newborn (suspected to be) affected by maternal use of alcohol
Perinatal Factors	P04.41	Newborn (suspected to be) affected by maternal use of cocaine
Perinatal Factors	P04.49	Newborn (suspected to be) affected by maternal use of other drugs of addiction
Perinatal Factors	P04.9	Newborn (suspected to be) affected by maternal noxious substance, unspecified
Perinatal Factors	P05.00	Newborn light for gestational age, unspecified weight
Perinatal Factors	P07.01	Extremely low birth weight newborn, less than 500 grams
Perinatal Factors	P07.02	Extremely low birth weight newborn, 500-749 grams
Perinatal Factors	P07.03	Extremely low birth weight newborn, 750-999 grams
Perinatal Factors	P07.10	Other low birth weight newborn, unspecified weight
Perinatal Factors	P07.14	Other low birth weight newborn, 1000-1249 grams

MEDICAL/BIOLOGICAL RISK CONDITION: **Perinatal Factors (continued)**

Condition Type	ICD-10 Code	Condition/Diagnosis
Perinatal Factors	P07.15	Other low birth weight newborn, 1250-1499 grams
Perinatal Factors	P07.16	Other low birth weight newborn, 1500-1749 grams
Perinatal Factors	P07.32	Preterm newborn, gestational age 29 completed weeks
Perinatal Factors	P07.33	Preterm newborn, gestational age 30 completed weeks
Perinatal Factors	P07.34	Preterm newborn, gestational age 31 completed weeks
Perinatal Factors	P07.35	Preterm newborn, gestational age 32 completed weeks
Perinatal Factors	P07.36	Preterm newborn, gestational age 33 completed weeks
Perinatal Factors	P07.37	Preterm newborn, gestational age 34 completed weeks
Perinatal Factors	P09	Abnormal findings on neonatal screening
Perinatal Factors	P29.81	Cardiac arrest of newborn
Perinatal Factors	P52.0	Intraventricular (nontraumatic) hemorrhage, grade 1, of newborn
Perinatal Factors	P52.1	Intraventricular (nontraumatic) hemorrhage, grade 2, of newborn
Perinatal Factors	P52.21	Intraventricular (nontraumatic) hemorrhage, grade 3, of newborn
Perinatal Factors	P52.22	Intraventricular (nontraumatic) hemorrhage, grade 4, of newborn
Perinatal Factors	P52.3	Unspecified intraventricular (nontraumatic) hemorrhage of newborn
Perinatal Factors	P52.5	Subarachnoid (nontraumatic) hemorrhage of newborn
Perinatal Factors	P61.9	Perinatal hematological disorder, unspecified
Perinatal Factors	P70.4	Other neonatal hypoglycemia
Perinatal Factors	P84	Other problems with newborn Intrauterine hypoxia
Perinatal Factors	P92.9	Feeding problem of newborn, unspecified
Perinatal Factors	P93.8	Other reactions and intoxications due to drugs administered to newborn
Perinatal Factors	P04.8	Newborn (Suspected to be) affected by other maternal noxious substances

MEDICAL/BIOLOGICAL RISK CONDITION: **Physical Impairments**

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.) *Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.

Condition Type	ICD-10 Code	Condition/Diagnosis
Physical Impairments	M06.9	Rheumatoid arthritis, unspecified
Physical Impairments	M41.20	Other idiopathic scoliosis, site unspecified
Physical Impairments	Q68.0	Congenital Torticollis
Physical Impairments	Q35.9	Cleft palate, unspecified
Physical Impairments	Q36.0	Cleft Lip, Bilateral
Physical Impairments	Q36.9	Cleft Lip, unilateral
Physical Impairments	Q37.9	Cleft Palate with Cleft Lip
Physical Impairments	Q71.60	Lobster-claw hand, unspecified hand
Physical Impairments	Q71.61	Lobster-Claw Right Hand
Physical Impairments	Q71.62	Lobster-Claw Left Hand
Physical Impairments	Q71.63	Lobster-Claw Hand, Bilateral
Physical Impairments	Q73.8	Other Reduction Defects of Unspecified Limbs
Physical Impairments	Q74.3	Arthrogryposis multiplex congenita
Physical Impairments	Q74.9	Unspecified Congenital Malformation of Limb(s)
Physical Impairments	Q75.0	Craniosynostosis
Physical Impairments	Q75.4	Mandibulofacial Dysostosis (Franschetti-Klein Syndrome)
Physical Impairments	Q79.0	Congenital Diaphragmatic Hernia
Physical Impairments	Q79.3	Gastronschisis
Physical Impairments	Q79.4	Prune Belly Syndrome
Physical Impairments	Q79.8	Other Congenital Malformations of Musculoskeletal System Poland's Syndrome
Physical Impairments	Q87.0	Congenital Malformation Syndrome Predominantly Affecting Facial Appearance Goldenhar's Syndrome (Oculouriculoverterbral Dysplasia)0

MEDICAL/BIOLOGICAL RISK CONDITION: **Sensory Abnormalities**

A diagnosed physical, mental, or neurobiological condition. The biological or medical risk condition shall be diagnosed by a health care provider and documentation shall be kept on file. Biological and medical risk conditions include the following: (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

***Questions Regarding conditions not on this list shall be directed to your FIT Manager who will work with a Department of Health Physician to determine whether a proposed condition will be recognized.**

Condition Type	ICD-10 Code	Condition/Diagnosis
Sensory Abnormalities	F98.4	Stereotyped Movement Disorder (Spasmus Nutans)
Sensory Abnormalities	H02.401	Unspecified ptosis of right eyelid
Sensory Abnormalities	H02.402	Unspecified ptosis of left eyelid
Sensory Abnormalities	H02.403	Unspecified ptosis of bilateral eyelids
Sensory Abnormalities	H50.00	Esotropia Unspecified
Sensory Abnormalities	H50.10	Exotropia Unspecified
Sensory Abnormalities	H50.811	Duane's Syndrome, Right Eye
Sensory Abnormalities	H50.812	Duane's Syndrome, Left Eye
Sensory Abnormalities	H50.89	Other Specified Strabismus
Sensory Abnormalities	H51.9	Unspecified disorder of binocular movement
Sensory Abnormalities	H52.00	Hypermetropia, unspecified eye
Sensory Abnormalities	H52.01	Hypermetropia, right eye
Sensory Abnormalities	H52.02	Hypermetropia, left eye
Sensory Abnormalities	H52.03	Hypermetropia, bilateral
Sensory Abnormalities	H52.13	Myopia, Bilateral
Sensory Abnormalities	H52.10	Myopia, unspecified eye

MEDICAL/BIOLOGICAL RISK CONDITION: **Sensory Abnormalities (Continued)**

Condition Type	ICD-10 Code	Condition/Diagnosis
Sensory Abnormalities	H52.11	Myopia, right eye
Sensory Abnormalities	H52.12	Myopia, left eye
Sensory Abnormalities	H53.001	Unspecified amblyopia, right eye
Sensory Abnormalities	H53.002	Unspecified amblyopia, left eye
Sensory Abnormalities	H53.003	Unspecified amblyopia, bilateral
Sensory Abnormalities	H53.009	Unspecified Amblyopia
Sensory Abnormalities	H53.8	Other visual disturbances
Sensory Abnormalities	H53.9	Unspecified Visual Disturbance
Sensory Abnormalities	H53.9	unspecified visual loss (Delayed Visual Maturation)
Sensory Abnormalities	H65.20	Chronic serous otitis media, unspecified ear
Sensory Abnormalities	H65.21	Chronic serous otitis media, right ear
Sensory Abnormalities	H65.22	Chronic serous otitis media, left ear
Sensory Abnormalities	H65.23	Chronic serous otitis media, bilateral
Sensory Abnormalities	H90.2	Conductive hearing loss, unspecified
Sensory Abnormalities	Q10.0	Congenital ptosis
Sensory Abnormalities	Q17.9	Congenital malformation of ear, unspecified

ENVIRONMENTAL RISK FACTORS: Personal History Presenting Hazard to Health

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program.
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Personal History Presenting Hazard to Health	Z77.011	Contact with and (suspected) exposure to lead
Personal History Presenting Hazard to Health	Z77.9	Other contact with and (suspected) exposures hazardous to health
Personal History Presenting Hazard to Health	Z91.410	Personal history of adult physical and sexual abuse (e.g., sexual abuse or rape)
Personal History Presenting Hazard to Health	Z91.412	Personal history of adult neglect (e.g., neglect)
Personal History Presenting Hazard to Health	Z91.49	Other personal history of psychological trauma, not elsewhere classified
Personal History Presenting Hazard to Health	Z77.9	Other specified personal risk factors, not elsewhere classified

ENVIRONMENTAL RISK FACTORS: Family History of Certain Chronic Disabling Diseases

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Family History of Certain Chronic Disabling Diseases	Z81.8	Family history of other mental and behavioral disorders
Family History of Certain Chronic Disabling Diseases	Z82.0	Family history of epilepsy and other diseases of the nervous system
Family History of Certain Chronic Disabling Diseases	Z82.69	Family history of other diseases of the musculoskeletal system and connective tissue

ENVIRONMENTAL RISK FACTORS: Family History of Certain Other Specific Conditions

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Family History of Certain Other Specific Conditions	Z81.0	Family History of Intellectual Disabilities

ENVIRONMENTAL RISK FACTORS: Family History of Other Conditions

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Family History of Other Conditions	Z82.1	Family history of blindness and visual loss
Family History of Other Conditions	Z82.2	Family history of deafness and hearing loss
Family History of Other Conditions	Z82.79	Family history of other congenital malformations, deformations and chromosomal abnormalities
Family History of Other Conditions	Z84.89	Family history of other specified conditions

ENVIRONMENTAL RISK FACTORS: Health Supervision of Infant or Child

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Health Supervision of Infant or Child	Z76.2	Encounter for health supervision and care of other healthy infant and child (e.g., supervision of healthy infant in cases of socioeconomic adverse condition at home)

ENVIRONMENTAL RISK FACTORS: Housing, Household, and Economic Circumstances

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Housing, Household, and Economic Circumstances	Z59.0	Homelessness (e.g., Social migrants, Transients)
Housing, Household, and Economic Circumstances	Z59.1	Inadequate housing (e.g., Lack of heating, Restriction of space)
Housing, Household, and Economic Circumstances	Z59.6	Low income (e.g., Economic problem, Poverty)
Housing, Household, and Economic Circumstances	Z59.8	Other problems related to housing and economic circumstances
Housing, Household, and Economic Circumstances	Z59.9	Unspecified housing or economic circumstance
Housing, Household, and Economic Circumstances	Z74.2	Need for assistance at home and no other household member able to render care

ENVIRONMENTAL RISK FACTORS: Other Family Circumstances

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program. (NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Other Family Circumstances	Z62.898	Other specified problems related to upbringing (e.g., Problem concerning adopted or foster child)
Other Family Circumstances	Z63.6	Dependent relative needing care at home (e.g., Care of sick or handicapped person in family or household)
Other Family Circumstances	Z63.72	Alcoholism and drug addiction in family
Other Family Circumstances	Z62.891	Other specified problems related to primary support group
Other Family Circumstances	Z69.010	Encounter for mental health services for perpetrator of parental child abuse Counseling for parent-child problem, unspecified

ENVIRONMENTAL RISK FACTORS: **Other Family Circumstances (Continued)**

Condition Type	ICD-10 Code	Condition/Diagnosis
Other Family Circumstances	Z69.020	Encounter for mental health services for victim of parental child abuse Counseling for victim of child abuse
Other Family Circumstances	Z69.11	Encounter for mental health services for victim of spousal or partner abuse (Counseling for victim of spousal and partner abuse)
Other Family Circumstances	Z69.12	Encounter for mental health services for perpetrator of spousal or partner abuse (Counseling for perpetrator of spousal and partner abuse)
Other Family Circumstances	Z69.020	Other specified counseling (Counseling for marital and partner problems, unspecified)

ENVIRONMENTAL RISK FACTORS: Other Psychosocial Circumstances

A presence of adverse family factors in the child's environment that increases the risk for developmental delay in children. Eligibility determination shall be made using the Environmental Risk Assessment (ERA) tool approved by the FIT program.
(NMAC 7.30.8.10.9 Requirements for Family Infant Toddler Early Intervention Services.)

Condition Type	ICD-10 Code	Condition/Diagnosis
Other Psychosocial Circumstances	Z13.4	Encounter for screening for certain developmental disorders in childhood
Other Psychosocial Circumstances	Z60.3	Acculturation difficulty (e.g., Cultural deprivation, Social isolation or persecution)
Other Psychosocial Circumstances	Z63.9	Problem related to primary support group, unspecified
Other Psychosocial Circumstances	Z65.3	Problems related to other legal circumstances (e.g., Imprisonment)
Other Psychosocial Circumstances	Z65.8	Other specified problems related to psychosocial circumstances (e.g., Life circumstance problems)
Other Psychosocial Circumstances	Z64.4	Other specified problems related to psychosocial circumstances (Interpersonal problems, not elsewhere classified)
Other Psychosocial Circumstances	Z65.9	Problem related to unspecified psychosocial circumstances