

List of Established Conditions that indicate automatic eligibility for *Early On*[®] supports and services. Conditions must be **diagnosed** by an appropriate health care or mental health provider and include, but are not limited to, the following:

<p>1. <u>Congenital Anomalies</u></p> <p>1.1. <u>Central Nervous System</u> Agenesis of the Corpus Callosum Holoprosencephaly Hydrocephalus w/o Spina Bifida Microcephalus Spina Bifida w/o Anencephaly</p> <p>1.2. <u>Eye, Ear, Face and Neck</u> Anophthalmos/Microphthalmos Anotia/Microtia CHARGE Syndrome Congenital Cataract Pierre Robin Sequence Treacher Collins</p> <p>1.3. <u>Heart and Circulatory System</u> Aortic Valve Atresia & Stenosis Coarctation of Aorta Hypoplastic Left Heart Patent Ductus Arteriosus (PDA) Tetralogy of Fallot</p> <p>1.4. <u>Respiratory System</u> Choanal Atresia Lung Agenesis/Hypoplasia</p> <p>1.5. <u>Cleft Lip & Palate</u> Cleft Palate w/o Cleft Lip Cleft Lip w/ and w/o Cleft Palate</p> <p>1.6. <u>Digestive System</u> Esophageal Atresia/Tracheoesophageal Fistula Hirschsprung's Disease Pyloric Stenosis</p> <p>1.7. <u>Genital & Urinary Organs</u> Hypospadias and Epispadias Renal Agenesis</p> <p>1.8. <u>Musculoskeletal System</u> Achondroplasia Arthrogryposis Congenital Hip Dislocation Lower Limb Reduction Deformities Upper Limb Reduction Deformities Other Congenital Anomalies of the Musculoskeletal system</p>	<p>1.9. <u>Other and Unspecified</u> Bardet-Beidl Syndrome Fragile X Syndrome</p> <p>2. <u>Chromosomal Anomalies</u> Angelman Syndrome Cri-du-Chat DiGeorge Syndrome (Velo-Cardial-Facial Syndrome) Klinefelter Syndrome Prader—Willi Syndrome Trisomy 21 (Down Syndrome) Trisomy 13 (Patau Syndrome) Trisomy 18 (Edwards Syndrome) Turner Syndrome Williams Syndrome</p> <p>3. <u>Infectious Conditions</u></p> <p>3.1. <u>Congenital Infections</u> HIV / AIDS Syphilis TORCH: Toxoplasmosis Rubella Cytomegalovirus Herpes</p> <p>3.2. <u>Acquired Infections</u> Bacterial Meningitis Encephalitis Poliomyelitis Viral Meningitis</p> <p>4. <u>Endocrine/Metabolic Disorders</u></p> <p>4.1. <u>Mucopolysaccharidosis</u> Hunter Syndrome Maroteaux-Lamy Syndrome Sanfilippo Syndrome Scheie Syndrome Sly Syndrome</p> <p>4.2. <u>Enzyme Deficiency</u> Biotinidase Deficiency Medium Chain Acyl-CoA Dehydrogenase Deficiency (MCADD) Oculocerebrorenal Syndrome (Lowe Syndrome)</p>	<p>4.3. <u>Abnormalities of Amino Acid Metabolism</u> Argininosuccinic Aciduria Citrullinemia Homocystinuria Infant Phenylketonuria (PKU) Maple Syrup Urine Disease Methylmalonic Acidemia (MMA) Ornithine Transcarbamylase Deficiency</p> <p>4.4. <u>Abnormalities of Carbohydrate Metabolism</u> Galactosemia Glycogen Storage Disease</p> <p>4.5. <u>Abnormalities of Lipid Metabolism</u> Gaucher Disease Niemann Pick Disease</p> <p>4.6. <u>Abnormalities of the Purine/Pyrimidine Metabolism</u> Lesch Nyhan Syndrome</p> <p>4.7. <u>Abnormalities of the Parathyroid</u> Untreated Hyperparathyroidism Untreated Hypoparathyroidism</p> <p>4.8. <u>Abnormalities of the Pituitary</u> Hyperpituitary Hypopituitary</p> <p>4.9. <u>Abnormalities of Adrenocortical Function</u> Congenital Adrenal Hyperplasia Hyperadrenocortical Function Hypoadrenocortical Function</p> <p>4.10. <u>Hemoglobinopathies</u> Sickle Cell Disease Thalassemia (major and minor)</p> <p>4.11. <u>Abnormalities of the Thyroid Hormone</u> Congenital Hypothyroidism</p> <p>4.12. <u>Peroxisomal Disorders</u> Adrenoleukodystrophy (ADL) Cerebrohepato renal Syndrome (Zellweger Syndrome) Rhizomelic Chondrodysplasia Punctata</p>
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Note: The Endocrine/Metabolic Disorders Category also includes all disorders tested for in the Michigan Newborn Screening Program.

Early On[®] Michigan

Established Conditions



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<p>5. <u>Other Disorders/Diseases</u></p> <p>5.1. <u>Neurological Disorders</u></p> <p><u>Neuromotor/Muscle Disorders</u></p> <ul style="list-style-type: none"> Cerebral Palsy Congenital Myasthenia Kernicterus Muscular Dystrophies Paralysis Periventricular Leukomalacia Torticollis Werdnig Hoffman Disease <p><u>Cerebrovascular Disease</u></p> <ul style="list-style-type: none"> Cerebral Arterial Thrombosis Cerebral Embolus Thrombosis Cerebral Venous Thrombosis <p><u>Brain Hemorrhages</u></p> <ul style="list-style-type: none"> Intracranial Hemorrhage Intraventricular Hemorrhage (grades III & IV) <p><u>Degenerative Disorders</u></p> <ul style="list-style-type: none"> Acute Disseminated Encephalomyelitis Cockayne Syndrome Friedreich's Ataxia Gangliosidosis Kugelberg-Welander Syndrome Leigh's Disease Leukodystrophy Schilder's Disease Tay Sachs Disease <p><u>Neurocutaneous Disorders</u></p> <ul style="list-style-type: none"> Block-Sulzberger Syndrome Neurofibromatosis Sturge Weber Syndrome Tuberous Sclerosis Xeroderma Pigmentosa <p><u>Malignancies</u></p> <ul style="list-style-type: none"> Intracranial Tumors and Other Malignancies of the CNS <p><u>Head and Spinal Cord Trauma</u></p> <ul style="list-style-type: none"> Fracture of vertebral column with or without spinal cord lesions Shaken Baby Syndrome Traumatic Brain Injury <p><u>Hypoxic/Anoxic Brain Injury</u></p> <ul style="list-style-type: none"> Hypoxic Ischemic Encephalopathy (Newborn Encephalopathy) Near Drowning 	<p>5.2. <u>Vision Impairment</u></p> <ul style="list-style-type: none"> Amblyopia Cortical Visual Impairment (CVI) Low Vision (20/700) Nystagmus Retinopathy of Prematurity (ROP) (Stage 3 - Stage 5) Visual Field Loss <p>6. <u>Hearing Deficiency</u></p> <ul style="list-style-type: none"> Auditory Neuropathy Bilateral or Unilateral hearing loss of ≥ 25 dB at 2+ frequencies between 500-4000 Hz. Mixed Hearing Loss Permanent Conductive Hearing Loss Sensorineural Hearing Loss Waardenburg Syndrome <p>7. <u>Other Fetal/Placental Anomalies</u></p> <ul style="list-style-type: none"> Twin to Twin Transfusion Syndrome Umbilical Cord Prolapse <p>8. <u>Exposures Affecting Fetus/Child</u></p> <p>8.1. <u>Prenatal</u></p> <ul style="list-style-type: none"> Fetal Alcohol Spectrum Disorders - Diagnosed Fetal Drug Exposure - Diagnosed Maternal PKU <p>8.2. <u>Postnatal</u></p> <ul style="list-style-type: none"> Lead – Blood Lead level at or above 10 micrograms per deciliter (10 μg/dL) Mercury – for recent exposure, blood level of more than 2 micrograms per deciliter (>2 μg/dL); for chronic exposure, urine level of more than 5 micrograms per deciliter (> 5 μg/dL) <p>9. <u>Chronic Illness</u></p> <p>9.1. <u>Medically Fragile</u></p> <ul style="list-style-type: none"> Renal Insufficiency 	<p>9.2. <u>Medical Illness</u></p> <ul style="list-style-type: none"> Bronchopulmonary Dysplasia Cancer Chronic Hepatitis Connective Tissue Disorders Cystic Fibrosis Diabetes Immune Disorders (ex. Juvenile Arthritis) Organic Failure to Thrive Renal Failure Very Low Birth Weight (<1500 grams or 3.3 lbs.) Chronic Asthma – moderate to severe Intrauterine Growth Retardation (IUGR) Small for Gestational Age (<10% weight for age) (SGA) <p>10. <u>Developmental Delay</u></p> <p>10.1. <u>Pervasive Developmental Disorders</u></p> <ul style="list-style-type: none"> Autism Spectrum Disorder Childhood Disintegrative Disorder Pervasive Developmental Disorders (NOS) <p>10.2. <u>Rett's Disorder</u></p> <p>10.3. <u>Regulatory Disorders of Sensory Processing</u></p> <ul style="list-style-type: none"> Hyposensitive / Hypersensitive Sensory-Seeking/Impulsive <p>11. <u>Mental Health Conditions</u></p> <ul style="list-style-type: none"> Adjustment Disorders Depression of Infancy and Early Childhood Maltreatment/Deprivation Disorder (A diagnosis of Reactive Attachment Disorder should be cross-walked to this diagnosis which is listed in the DC:0-3R) Disorders of Affect Mixed Disorders of Emotional Expressiveness Post Traumatic Stress Disorder (PTSD) Regulatory Disorders** <p>** Difficulties in regulating physiological, attentional, motor or affective processes, and in organizing a calm, alert or affectively positive state. These disorders affect the child's daily routines and interpersonal relationships. Must be diagnosed by a qualified professional. (Greenspan, 1992)</p>
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