Part C of the IDEA and Family Education and Support Programs:
Use of the Established Condition List for Type I Eligibility
May 23, 2019

When determining Type I Eligibility for Part C of the IDEA and Family Education and Support Programs:

1. The Program providers use the Established Condition Statement as completed and signed by a physician or psychologist.
2. After receipt of the Statement, the Program providers' designated personnel will refer to this Established Condition List to determine if the child's condition is one that is identified as meeting Type I Established Condition criteria in the Montana.
3. If yes, the Program provider completes the multidisciplinary assessment (with written consent from the family) of the child's needs and family's concerns, priorities, and resources to enhance the development of the child for the development of the IFSP.
4. If no, the Program provider (with written consent from the family) moves toward establishing Type II Measured Delay eligibility (50% delay in one developmental domain or two or more 25% delays in developmental domains) using a multidisciplinary evaluation team that includes the family, the service coordinator, and at least one other discipline.
5. If yes, the child meets Type II Measured Delay eligibility, the Program provider completes (with written consent from the family) the multidisciplinary assessment of the child's needs and family's concerns, priorities, and resources to enhance the development of the child for the development of the IFSP.
6. If no, the family receives written notification of the eligibility decision and the Program provider shares more appropriate referrals with the family.
7. Eligibility is determined annually, and child assessments are completed at least annually to monitor development and appropriately develop IFSPs based upon the child's assessed developmental needs and the family's concerns, priorities, and resources to enhance the development of the child.

Montana DPHHS Part C of the IDEA Program and FES Program Established Condition List 6 12 2019

The following conditions meet the Type I Established Condition criteria in Montana

Medical Diagnosis Name	**The following conditions <u>meet</u> the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19.** Diagnosis Details
Acrocallosal Syndrome, Schinzel	Acrocallosal Syndrome, Schinzel Type (aka: Schinzel Acrocallosal Syndrome; Acrocallosal Syndrome, Schinzel Type; ACLS; ACS; Hallux duplication, postaxial polydactyly, absence of the corpus callosum, severe mental retardation syndrome)
Adrenoleukodystrophy	Adrenoleukodystrophy (aka: ALD; AMN; X-ALD; Addison disease and cerebral sclerosis; Adrenomyeloneuropathy; Siemerling-creutzfeldt disease; Bronze schilder disease; Schilder Disease; Melanodermic Leukodystrophy; sudanophilic leukodystrophy; Pelizaeus Merz...
Agenesis of Corpus Callosum	Agenesis of the Corpus Callosum (aka: Absence of the corpus callosum; Hypogenesis of the corpus callosum; Dysplastic corpus callosum)
Aicardi Syndrome	Aicardi Syndrome (aka: Agenesis of Corpus Callosum and Chorioretinal Abnormality; Agenesis of Corpus Callosum With Chorioretinitis Abnormality; Agenesis of Corpus Callosum With Infantile Spasms and Ocular Anomalies; Chorioretinal Anomalies with Agenesis ...
Alexander Disease	Alexander Disease
Allan Herndon Syndrome	Allan Herndon Syndrome (aka: Allan-Herndon-Dudley Syndrome (AHDS); Allan-Herndon-Dudley Mental Retardation)
Alper Disease	Alper Disease (aka: Alper's Diffuse Degeneration of Cerebral Gray Matter with Hepatic Cirrhosis; Alpers Progressive Infantile Poliodystrophy; Christensen's disease; Christensen-Krabbe disease; Diffuse Cerebral Degeneration in Infancy; Poliodystrophia Cer...
Anencephaly	Anencephaly
Angelman Syndrome	Angelman Syndrome (aka: AS)
Aniridia Cerebellar Ataxia MD	Aniridia Cerebellar Ataxia Mental Deficiency (aka: Gillespie syndrome; Aniridia, Partial-Cerebellar Ataxia-Oligophrenia; Aniridia-Cerebellar Ataxia-Mental Retardation)hrenia)
Anophthalmia, bilateral	Anophthalmia, bilateral
Aphasia w brain damage	Aphasia (with evidence of brain damage) (aka: aphemia with evidence of brain damage)
Argininosuccinic aciduria	Argininosuccinic aciduria (aka: Arginino Succinase Deficiency; ASA Deficiency; Argininosuccinate Lyase Deficiency; ASL Deficiency)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Asphyxia w brain damage	Asphyxia (with evidence of brain damage within athe first couple days of the event)
Autism Spectrum Disorder	Autism Spectrum Disorder (aka: Infantile Autism; Kanner Syndrome; Autistic Disorder; Asperger Syndrome; Asperger's Disorder; Pervasive Developmental Disorder; PDD; Pervasive Developmental Disorder-Not Otherwise Specified; PDD-NOS)
Baller Gerold Syndrome	Baller Gerold Syndrome (aka: Craniosynostosis with Radial Defects, Craniosynostosis-Radial Aplasia Syndrome)
Bannayan Riley Ruvalcaba	Bannayan Riley Ruvalcaba (aka: Bannayan Riley Ruvalcaba Syndrome; BRRS, Bannayan-Zonana syndrome (BZS); Macrocephaly, multiple lipomas, and hemangiomata; Macrocephaly, pseudopapilledema, and multiple hemangiomata; Riley-Smith syndrome; Ruvalcaba-Myhre-Sm...
Bardet-Biedl Syndrome	Bardet-Biedl Syndrome (aka: Biedl-Bardet Syndrome; Laurence Moon-Biedl)
Batten Disease	Batten Disease (aka: CLN3, JNCL, Neuronal Ceroid Lipofuscinosis, Juvenile, Spielmeyer-Sjogren Disease, Vogt-Spielmeyer Disease, or Vogt-Spielmeyer-Sjogren Disease)
Borjeson Syndrome	Borjeson Syndrome (aka: BORJ; Borjeson-Forssman-Lehmann Syndrome; BFLS)
Brain Tumor	Brain Tumor (aka: brain cancer; brain teratoma; glioma; astrocytoma; glioblastoma multiforme; ependymoma; oligodendroglioma; medulloblastoma; meningioma; Schwannoma; acoustic neuroma; craniopharyngioma; germ cell tumor of the brain; germinoma; pineal reg...
C Syndrome	C Syndrome (aka: Opitz Trigonocephaly Syndrome, Trigonocephaly "C" Syndrome, Trigonocephaly Syndrome)
Cardiofaciocutaneous Syndrome	Cardiofaciocutaneous syndrome (aka: CFC syndrome, Cardio-facial-cutaneous syndrome, Facio-cardio-cutaneous syndrome)
Cerebellar agenesis	Cerebellar Agenesis (aka: Cerebellar Aplasia; Cerebellar Hemiagenesis; Cerebellar Hypoplasia; Cerebellar Atrophy)
Cerebral atrophy	Cerebral atrophy
Cerebral Dysgenesis	Cerebral Dysgenesiias
Cerebro Oculo Facio Skel Syndrome	Cerebro Oculo Facio Skeletal Syndrome (aka: COFS Syndrome, Cerebrooculofacioskeletal Syndrome, Cockayne Syndrome type II, Pena Shokeir II Syndrome, Pena Shokeir Syndrome Type II)
Cerebromalacia	Cerebromalacia (aka: Encephalomalacia)
CHAMP 1	Champ 1 gene mutation
CHARGE Syndrome	CHARGE Syndrome (aka: CHARGE Association)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Chromosome 10, Monosomy 10p	Chromosome 10, Monosomy 10p (aka: 10p Deletion Syndrome (Partial); Chromosome 10, 10p- Partial; Chromosome 10, Partial Deletion (short arm))
Chromosome 11, Partial Monosomy 11q	Chromosome 11, Partial Monosomy 11q (aka: 11q- Syndrome; Partial, Deletion 11q Syndrome, Partial; Distal 11q Monosomy; Distal 11q- Syndrome; JBS; Jacobsen Syndrome; Monosomy 11q, Partial; Partial Monosomy of Long Arm of Chromosome 11; 11q14 deletion; 11q21 deletion; 11q14-21 deletion)
Chromosome 11, Partial Trisomy 11p14.3	Chromosome 11, Partial Trisomy 11p14.3
Chromosome 11, Partial Trisomy 11q	Chromosome 11, Partial Trisomy 11q (aka: 11q Partial Trisomy, Chromosome 11; Partial Trisomy 11q13-qter, Chromosome 11; Partial Trisomy 11q21-qter;, Chromosome 11; Partial Trisomy 11q23-qter; Distal Trisomy 11q; Partial Trisomy 11q; Trisomy 11q, Partial)
Chromosome 12 Deletion	Chromosome 12 deletion
Chromosome 12 Partial Trisomy	Partial Trisomy 12
Chromosome 13, Partial Monosomy 13q	Chromosome 13, Partial Monosomy 13q (aka: 13q- Syndrome, Partial; Deletion 13q Syndrome, Partial; Monosomy 13q, Partial; Partial Monosomy of the Long Arm of Chromosome 13)
Chromosome 14 Deletion	Chromosome 14 deletion
Chromosome 14 Ring	Chromosome 14 Ring (aka: Ring 14; Ring Chromosome 14; r14)
Chromosome 14, Trisomy Mosaic	Chromosome 14, Trisomy Mosaic (aka: Trisomy 14 Mosaic; Trisomy 14 Mosaicism Syndrome; Trisomy 14 Syndrome)
Chromosome 15 Ring	Chromosome 15 Ring (aka: Ring 15; Ring 15 Chromosome; Ring 15 Chromosome (mosaic pattern); r15)
Chromosome 15, Distal Trisomy 15q	Chromosome 15, Distal Trisomy 15q (aka: Chromosome 15, Trisomy 15q2; Distal Duplication 15q; Partial Duplication 15q Syndrome)
Chromosome 15q11-q13 Dup	Chromosome 15q11-q13 duplication (aka: Isodicentric 15; Inverted duplication 15)
Chromosome 15q24 microdeletion syndrome	Chromosome 15q24 Microdeletion Syndrome
Chromosome 16 Duplication	Chromosome 16 Duplication
Chromosome 17q12 Duplication	Chromosome 17q12 Duplication
Chromosome 18 Ring	Chromosome 18 Ring (aka: Ring 18; Ring Chromosome 18; r18)
Chromosome 18, Tetrasomy 18p	Chromosome 18, Tetrasomy 18p (aka: Tetrasomy, Short Arm of Chromosome 18)
Chromosome 19p duplication	Chromosome 19p duplication

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Chromosome 20q Trisomy	Chromosome 20q Trisomy (aka: Chromosome 20q Duplication; Partial Trisomy 20q; Trisomy 20q11.2; Chromosome 20q11.2 Duplication; Trisomy 20)
Chromosome 22 Ring	Chromosome 22 Ring (aka: Ring 22; Ring 22, Chromosome; r22)
Chromosome 22, Trisomy Mosaic	Chromosome 22, Trisomy Mosaic (aka: Trisomy 22 Mosaic; Trisomy 22 Mosaicism Syndrome)
Chromosome 22q11.2 duplication	Chromosome 22q11.2 duplication
Chromosome 2q32 Deletion	Chromosome 2q32 deletion
Chromosome 3, Monosomy 3p2	Chromosome 3, Monosomy 3p2 (aka: Chromosome 3, Deletion of Distal 3p; Chromosome 3, Distal 3p Monosomy; Monosomy 3p2; Partial Deletion of Chromosome 3)
Chromosome 3, Trisomy 3q2	Chromosome 3, Trisomy 3q2 (aka: Chromosome 3, Distal 3q2 Duplication; Chromosome 3, Distal 3q2 Trisomy; Partial Duplication 3q Syndrome; Partial Trisomy 3q Syndrome)
Chromosome 4 Ring	Chromosome 4 Ring (aka: Ring 4; Ring 4, Chromosome; r4)
Chromosome 4, Monosomy 4q	Chromosome 4, Monosomy 4q (aka: Interstitial Deletion of 4q; Proximal Deletion of 4q; Terminal Deletion of 4q; Chromosome 4Q minus micro deletion)
Chromosome 4, Monosomy Distal 4q	Chromosome 4, Monosomy Distal 4q (aka: 4q Deletion Syndrome, Partial; Chromosome 4, 4q Terminal Deletion Syndrome; Chromosome 4, Partial Monosomy 4q; Del(4q) Syndrome, Partial; Distal 4q Monosomy; Distal 4q- Syndrome; Chromosome 4Q minus micro deletion)
Chromosome 4, Partial Trisomy Distal 4q	Chromosome 4, Partial Trisomy Distal 4q (aka: Chromosome 4, Partial Trisomy 4q (4q2 and 4q3); Chromosome 4, Partial Trisomy 4q (4q21-qter to 4q32-qter), Distal 4q Trisomy, Dup(4q) Syndrome; Partial Duplication 4q Syndrome; Partial Trisomy 4q Synd...
Chromosome 4, Trisomy 4p	Chromosome 4, Trisomy 4p (aka: Chromosome 4 (Partial Trisomy 4p); Dup(4p) Syndrome; Duplication 4p Syndrome)
Chromosome 5, Trisomy 5p	Chromosome 5, Trisomy 5p (aka: Chromosome 5, Trisomy 5p, (Complete (5p11-ter), Included); Chromosome 5, (Trisomy 5p, Partial, Included); Dup(5p) Syndrome; Duplication 5p Syndrome)
Chromosome 6 Ring	Chromosome 6 Ring (aka: Ring 6; Ring 6, Chromosome; r6)
Chromosome 6, Partial Trisomy 6q	Chromosome 6, Partial Trisomy 6q (aka: 6q+ Syndrome, Partial; Chromosome 6, Trisomy 6q2; Distal Duplication 6q; Distal Trisomy 6q; Duplication 6q, Partial; Trisomy 6q Syndrome, Partial; Trisomy 6q, Partial)
Chromosome 6p Partial Monosomy	Chromosome 6p, Partial Monosomy (aka Partial Deletion of Chromosome 6p)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
6q terminal deletion syndrome	6q terminal deletion syndrome
Chromosome 7, Partial Monosomy 7p	Chromosome 7, Partial Monosomy 7p (aka: Chromosome 7, 7p Deletion Syndrome, Partial; Chromosome 7, Partial Deletion of Short Arm; Del(7p) Syndrome, Partial; Interstitial 7p Monosomy; Partial 7p Monosomy; Terminal 7p Monosomy; Terminal 7p Monosomy)
Chromosome 7p Partial Dup Syndrome	Chromosome 7p Partial Duplication Syndrome (aka: 7p Duplication Syndrome)
Chromosome 7q Partial Monosomy	Chromosome 7q Partial Monosomy (aka: Chromosome 7q Deletion; Chromosome 7q Partial Deletion; Chromosome 7q11.22 deletion; Chromosome 7q34-36.1 deletion; Monosomy Chromosome 7q; Partial Monosomy Chromosome 7q; Monosomy Chromosome 7q34-36.1)
Chromosome 8, Monosomy 8p2	Chromosome 8, Monosomy 8p2 (aka: 8p- Syndrome, Partial; Chromosome 8, 8p Deletion Syndrome, Partial Chromosome 8, Partial Deletion of Short Arm; Chromosome 8, Partial Monosomy 8p2; Del(8p) Syndrome, Partial; Distal 8p Monosomy; Partial 8p Monosomy; Monosomy 8p23.1)
Chromosome 8, Monosomy 8p2	Chromosome 8, Monosomy 8p2
Chromosome 9 Ring	Chromosome 9 Ring (aka: r9; Ring 9; Ring 9, Chromosome)
Chromosome 9, Complete Trisomy 9P	Chromosome 9 Trisomy (aka: Complete Trisomy 9P; Partial Trisomy 9; Chromosome 9, Partial Trisomy 9P; Trisomy 9P Syndrome (Partial); Rethore Syndrome (obsolete); Duplication 9p Syndrome; Dup(9p) Syndrome; Chromosome 9, Trisomy 9pter-q11-13; Chromosome 9, ...
Chromosome 9, Tetrasomy 9p	Chromosome 9, Tetrasomy 9p (aka: Chromosome 9, Tetrasomy 9p Mosaicism; Mosaic Tetrasomy 9p; Tetrasomy 9p; Tetrasomy, Short Arm of Chromosome 9)
Chromosome 9, Trisomy Mosaic	Chromosome 9, Trisomy Mosaic (aka: Trisomy 9 Mosaic; Trisomy 9 Mosaicism; Trisomy 9 Mosaicism Syndrome)
Chromosome 9q Partial Monosomy	Chromosome 9q Partial Monosomy
Chromosome Xq26.2 duplication	Chromosome Xq26.2 duplication (aka: duplication of the distal portion of the long arm of the X chromosome; chromosome X duplication; chromosome Xq duplication)
Chromosome 10q25 deletion syndrome	
Chromosome 10q26 deletion syndrome	

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Chromosome 12p duplication	
Chromosome 17p 13.2 duplication	
Chromosome 17p13.1 and/or 17p13.2 microdeletion	Chromosome 17p13.1 and/or 17p13.2 microdeletion
Chromosome 1p36 Deletion Syndrome	Chromosome 1p36 deletion
Chromosome 21q Partial Deletion Syndrome	Chromosome 21q Partial Deletion Syndrome (aka: 21q22 Deletion), not including small deletions of only 21q22.3
Chromosome 7q duplication	aka 7q22.3-7q36.1 duplication
Chromosome 8p inverted duplication/deletion syndrome	
Chromosome Xp deletion	
Chromosome 2q37 Deletion	
Closed Head Injury	Closed Head Injury with neuroradiological evidence of intracranial injury (e.g., subarachnoid hemorrhage, or intracranial hemorrhage, or swelling) (aka: Traumatic Brain Injury; TBI)
Cockayne Syndrome	Cockayne Syndrome (aka: CS; Deafness-Dwarfism-Retinal Atrophy; Dwarfism with Renal Atrophy and Deafness; Neill-Dingwall Syndrome; Progeroid Nanism)
Coffin-Lowry Syndrome	Coffin-Lowry syndrome (aka: Coffin Syndrome; Mental Retardation with Osteocartilaginous Abnormalities)
Coffin-Siris Syndrome	Coffin-Siris syndrome (aka: Dwarfism-Onychodysplasia; Fifth Digit Syndrome; Mental Retardation with Hypoplastic 5th Fingernails and Toenails; Short Stature-Onychodysplasia)
Cohen Syndrome	Cohen Syndrome (aka: Pepper Syndrome)
Colpocephaly	Colpocephaly
Cornelia de Lange Syndrome	Cornelia de Lange Syndrome (aka: BDLS; Brachmann-de Lange Syndrome; CdLS; de Lange Syndrome)
Costello Syndrome	Costello Syndrome (aka: FCS Syndrome; Faciocutaneoskeletal Syndrome)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Dandy Walker Syndrome	Dandy Walker Syndrome (aka: Dandy-Walker Cyst; Dandy-Walker Deformity; Hydrocephalus, Internal, Dandy-Walker Type; Hydrocephalus, Noncommunicating, Dandy-Walker Type; DWM; Luschka-Magendie Foramina Atresia)
De Barsy Syndrome	De Barsy Syndrome (aka: Corneal Clouding-Cutis Laxa-Mental Retardation, Cutis Laxa-Growth Deficiency Syndrome; De Barsy-Moens-Diercks Syndrome; Progeroid Syndrome of De Barsy)
De Sanctis Cacchione Syndrome	De Sanctis Cacchione Syndrome (aka: Xerodermic Idiocy)
Deletion 18p Syndrome	Deletion 18p Syndrome (aka: Chromosome 18, Monosomy 18p; 18p Deletion Syndrome; 18p- Syndrome; Del(18p) Syndrome; Monosomy 18p Syndrome; Short Arm 18 Deletion Syndrome; Partial Deletion of Chromosome 18)
Deletion 18q Syndrome	Deletion 18q syndrome (aka: Chromosome 18q- Syndrome; 18q Deletion Syndrome; 18q- Syndrome; Chromosome 18 Long Arm Deletion Syndrome; Chromosome 18, Monosomy 18Q; Del(18q) Syndrome; Monosomy 18q Syndrome; Partial Deletion of Chromosome 18)
Deletion 22Q Syndrome	22q11.2 Deletion Syndrome (aka: DiGeorge syndrome; velocardiofacial syndrome; Deletion 22q11.2 syndrome; Shprintzen syndrome) 22q13 deletion syndrome (aka: Phelan-McDermid Syndrome)
Deletion 5p Syndrome	Deletion 5p Syndrome (aka: Cri du chat Syndrome; Chromosome 5p-syndrome; Cat's Cry Syndrome; Chromosome 5, Monosomy 5p; Chromosome 5p-Syndrome; Le Jeune Syndrome; Partial Deletion of the Short Arm of Chromosome 5 Syndrome)
Deletion 9p Syndrome	Deletion 9p Syndrome (aka: Chromosome 9, Partial Monosomy 9p; 9p Partial Monosomy; 9p- Syndrome, Partial; Chromosome 9, Partial Monosomy 9p22; Chromosome 9, Partial; Monosomy 9p22-pter; Del(9p) Syndrome, Partial; Deletion 9p Syndrome, Partial; Distal 9p-...
Diencephalic Syndrome	Diencephalic Syndrome (aka: Diencephalic Syndrome of Childhood; Diencephalic Syndrome of Emaciation; Paramedian Diencephalic Syndrome; Russell's Diencephalic Cachexia; Russell's Syndrome)
DOOR Syndrome	DOOR Syndrome (aka: DOOR(S) Syndrome; Deafness, Onychodystrophy, Osteodystrophy,and Mental Retardation)
Down Syndrome	Down Syndrome (aka: Trisomy 21 Syndrome; Chromosome 21, Mosaic 21 Syndrome; Chromosome 21, Translocation 21 Syndrome; Trisomy G Syndrome)
Dravet Syndrome	Dravet Syndrome (aka: severe myoclonic epilepsy of infancy; SMEI)
Dubowitz Syndrome	Dubowitz Syndrome (aka: Intrauterine Dwarfism)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Duchenne Muscular Dystrophy	Duchenne Muscular Dystrophy (aka: Childhood Muscular Dystrophy; DMD; Muscular Dystrophy (Classic X-linked Recessive); Progressive Muscular Dystrophy of Childhood; Pseudohypertrophic Muscular Dystrophy)
Duplication 10q Syndrome	Duplication 10q Syndrome (aka: Chromosome 10, distal trisomy 10q; Chromosome 10, Partial Trisomy 10q24-qter; Chromosome 10, Trisomy 10q2; Distal Duplication 10q; Distal Trisomy 10q Syndrome; Dup(10q) Syndrome)
Dyggve Melchior Clausen Syndrome	Dyggve Melchior Clausen Syndrome (aka: DMC Disease; DMC Syndrome; Smith-McCort Dysplasia)
Emanuel Syndrome	Emanuel Syndrome (aka: Emanual Syndrome; Derivative 22; der(22) chromosome; Supernumerary der(22) Syndrome)
Encephalitis, Herpes Simplex	Encephalitis, Herpes Simplex (aka: HSE; Herpes Encephalitis; Herpetic Brainstem Encephalitis; Herpetic Meningoencephalitis)
Encephalitis, Rasmussen's	Encephalitis, Rasmussen's (aka: Chronic Encephalitis and Epilepsy; Chronic Localized (Focal) Encephalitis; Epilepsy, Hemiplegia and Mental Retardation; Rasmussen's Syndrome)
Encephalocele	Encephalocele (aka: Bifid Cranium, Cephalocele, Cranial Meningoencephalocele, Craniocele, Cranium Bifidum)
Encephalopathy Congenital	Encephalopathy Congenital (aka: neonatal encephalopathy)
Encephalopathy, Hypoxic Ischemic	Encephalopathy, Hypoxic Ischemic (aka: HIE; subaccute hypoxic injury)
Encephalopathy, Static	Encephalopathy, Static
Epidermal Nevus Syndrome	Epidermal Nevus Syndrome (aka: Ichthyosis Hystrix Gravior; Inflammatory Linear Nevus Sebaceous Syndrome; Lambert Type Ichthyosis; Linear Nevus Sebacous Syndrome; Linear Sebaceous Nevus Sequence; Linear Sebaceous Nevus Syndrome; Nevus Sebaceous of Jadasso...
Fahr's Disease	Fahr's Disease (aka: Cerebrovascular Ferrocalcinosis; Fahr Disease; Idiopathic Basal Ganglia Calcificationl; IBGC; Nonarteriosclerotic Cerebral Calcifications; SPD Calcinosis; Striopallidodentate Calcinosis)
Familial MR Syndrome	Familial Mental Retardation Syndromes (incl: Familial Mental Retardation Syndrome ATR-16)
Fetal Alcohol Syndrome	Fetal Alcohol Syndrome (aka: FAS)
Fetal Hydantoin Syndrome	Fetal Hydantoin Syndrome (aka: Dilantin Embryopathy, Phenytoin Embryopathy)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
FG Syndrome	FG Syndrome (aka: Opitz-Kaveggia Syndrome)
Fiber Type Disproportion, Congenital	Fiber Type Disproportion, Congenital (aka: Atrophy of Type I Fibers; CFTD; CFTDM; Myopathy of Congenital Fiber Type Disproportion; Myopathy, Congenital, With Fiber-Type Disproportion)
Fibrodysplasia Ossificans Progressiva	Fibrodysplasia Ossificans Progressiva (aka: FOP; Myositis Ossificans Progressiva)
Filippi Syndrome	Filippi Syndrome (aka: Syndactyly Type I with Microcephaly and Mental Retardation)
Floating Harbor Syndrome	Floating Harbor Syndrome (aka: FHS; Pelletier-Leisti syndrome)
Fountain Syndrome	Fountain Syndrome (aka: Mental Retardation-Deafness-Skeletal Abnormalities-Coarse Face with Full Lips)
Fragile X Syndrome	Fragile X Syndrome (aka: Fragile Site, Folic Acid Type, Rare; Fra(X)(Q27.3); Fragile X Mental Retardation Protein; FMRP; Fragile X Mental Retardation Syndrome; Marker X Syndrome; Martin-Bell Syndrome; Mental Retardation, X-Linked, Associated With Mar Xq2...
Friedreich's ataxia	Friedreich's ataxia
Fryns Syndrome	Fryns Syndrome (aka: FRNS)
Fucosidosis	Fucosidosis (aka: Alpha-L-Fucosidase Deficiency)
Fukuyama Type Congenital Muscular Dystrophy	Fukuyama Type Congenital Muscular Dystrophy (aka: Cerebromuscular Dystrophy, Fukuyama Type; Congenital Muscular Dystrophy, Fukuyama Type; FCMD; Micropolygyria With Muscular Dystrophy; Muscular Dystrophy, Congenital Progressive with Mental Retardation; Mu...
Galactosemia	Galactosemia (aka: Classic Galactosemia; GALT Deficiency; Galactose-1-Phosphate Uridyl Transferase Deficiency)
Gaucher Disease Type III	Gaucher Disease Type III (Subacute/Chronic form, or "Atypical, due to Saposin C Deficiency") (aka: Gaucher Disease, Cardiovascular Form; Gaucher Disease, Type 3)
Gaucher Disease, Type II	Gaucher Disease, Type II ("Perinatal Lethal" or Colloidal Type) (aka: Gaucher Disease, Type 2; Glucocerebrosidase Deficiency; Glucosylceramidase Deficiency)
Glycinemia	Glycinemia (aka: glycine encephalopathy; nonketotic hyperglycinemia; transient neonatal hyperglycinemia)
Gm1Gangliosidosis	Gm1 Gangliosidosis (aka: Beta-Galactocidase-1 Deficiency; GLB1 Deficiency; Galactocidase, Beta-1; GLB1; Morquio Disease, Type B; Elastin-Binding Protein)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Griscelli Syndrome, type 1	Griscelli Syndrome, type 1
Hallervorden-Spatz Disease	Hallervorden-Spatz Disease (aka: Pantothenate kinase-associated neurodegeneration; PKAN)
Hallgren Syndrome	Hallgren Syndrome
Hearing Loss (bilateral loss)	Hearing Loss (diagnosis with bilateral hearing loss)
Hemimegalencephaly	Hemimegalencephaly
HIV (confirmed)	HIV (where the child's status of the HIV infection has been confirmed)
Holoporencephaly	Holoporencephaly (aka: Alobar Holoprosencephaly; Arhinencephaly; Familial Alobar Holoprosencephaly; HS; Holoprosencephaly Malformation Complex; Holoprosencephaly Sequence; Lobar Holoprosencephaly; Semilobar Holoprosencephaly)
Human HOXA1 Syndrome	Human HOXA1 Syndromes (aka: Athabaskan Brainstem Dysgenesis Syndrome (ABDS); Navajo Brainstem Syndrome; Bosley-Salih-Alorainy Syndrome; BSAS)
Hunter Syndrome	Hunter Syndrome (aka: MPSII, MPS Disorder II, Mucopolysaccharidosis Type II)
Hurler Syndrome	Hurler Syndrome (aka: Mucopolysaccharidosis Type I; MPS I-H; MPS1)
Hydranencephaly	Hydranencephaly
Hydrocephalus, Congenital	Hydrocephalus, Congenital (aka: Benign Hydrocephalus, Communicating Hydrocephalus, Internal Hydrocephalus, Non-Communicating Hydrocephalus, Normal Pressure Hydrocephalus, Obstructive Hydrocephalus)
Hydrocephalus, Post-hemorrhagic	
Hypomelanosis of Ito	Hypomelanosis of Ito (aka: Incontinentia pigmentosa acromians or Incontinentia pigmentosa)
I Cell Disease	I Cell Disease
Ichthyosis, Sjogren Larsson Syndrome	Ichthyosis, Sjogren Larsson Syndrome
Infantile Neuroaxonal Dystrophy	Infantile Neuroaxonal Dystrophy (aka: INAD or Seitelberger Disease)
IVH Grade III	Intraventricular Hemorrhage (aka: IVH, periventricular hemorrhage, PVH) Grade III
IVH Grade IV	Intraventricular Hemorrhage (aka: IVH, periventricular hemorrhage, PVH) Grade IV
Johanson-Blizzard Syndrome	Johanson-Blizzard Syndrome
Joubert Syndrome	Joubert Syndrome
Juberg-Marsidi Syndrome	Juberg-Marsidi Syndrome
KBG Syndrome	KBG Syndrome

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Keratitis Ichthyosis Deafness Syndrome	Keratitis Ichthyosis Deafness Syndrome
Klinefelter Syndrome	Klinefelter Syndrome (aka: XXY Syndrome)
Kufs Disease	Kufs Disease (aka: Jansk)
Kugelberg Welander Syndrome	Kugelberg Welander Syndrome (aka: Wohlfart-Kugelberg-Welander syndrome or mild SMA)
L1 Syndrome	L1 Syndrome
Landau Kleffner Syndrome	Landau Kleffner Syndrome
Langer-Giedion Syndrome	Langer-Giedion Syndrome (aka: Tricho-Rhino-Phalangeal Syndrome Type II, TRP II)
Lead encephalopathy	Lead encephalopathy (aka: lead poisoning encephalopathy)
Leigh's Disease	Leigh's Disease
Lennox Gastaut Syndrome	Lennox Gastaut Syndrome
Lenz Microphthalmia Syndrome	Lenz Microphthalmia Syndrome
Lesch-Nyhan Syndrome (LNS)	Lesch-Nyhan syndrome (LNS)
Leukodystrophy	Leukodystrophy (aka: Metachromatic Leukodystrophy; Krabbe Disease; Krabbe Leukodystrophy; globoid cell leukodystrophy; galactosylceramide lipidosis; Canavan Disease; ACY2 Deficiency; ASP Deficiency; ASPA Deficiency; Aminoacylase-2 Deficiency; Aspartoacyl...
Levy-Yeboa Syndrome	Levy-Yeboa Syndrome
Ligase IV syndrome	Ligase IV Syndrome (aka: Ligase IV Deficiency; LIG4 Syndrome)
Lipodystrophy, generalized	Lipodystrophy, generalized (aka: Berardinelli Lipodystrophy; Berardinelli Lipodystrophy Syndrome; Congenital Generalized Lipodystrophy)
Lissencephaly	Lissencephaly
Locked In Syndrome	Locked In Syndrome
Lowe Syndrome	Lowe Syndrome (aka: Oculocerebrorenal Syndrome)
Lysosomal Storage Disorders	Lysosomal Storage Disorders (aka: Cystinosis)
Macrocephaly Cutis Marmorata Telangiectatica	Macrocephaly, Cutis Marmorata Telangiectatica Congenita Syndrome (aka: M-CMTC)
Maple Syrup Urine Disease, untreated	Maple Syrup Urine Disease (where the diagnosis is late, or there is no or inadequate treatment) (aka: BCKD Deficiency, Branched Chain Alpha-Ketoacid Dehydrogenase Deficiency, Branched Chain Ketonuria I, Classical Maple Syrup Urine Disease)
Marden Walker Syndrome	Marden Walker Syndrome
Marinesco Sjogren Syndrome	Marinesco Sjogren Syndrome

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Marshall Smith Syndrome	Marshall Smith Syndrome
Meckel-Gruber Syndrome	Meckel-Gruber Syndrome (aka: Meckel Syndrome (w/ skull defect); Dysencephalia Splanchnocystica)
Megalocornea MR Syndrome	Megalocornea Mental Retardation Syndrome (aka: MMR Syndrome; Neuhauser Syndrome)
MELAS Syndrome	MELAS Syndrome
Menkes Syndrome	Menkes Syndrome (aka: Menkes Disease; Kinky Hair Disease)
MERRF Syndrome	MERRF Syndrome
MHBD Deficiency	MHBD Deficiency (a.k.a. 2-methyl-3-hydroxybutyryl-CoA Dehydrogenase Deficiency)
Microcephaly	Microcephaly
Microdeletion 15q13.3 Syndrome	
Motor Neuron Disease	Motor Neuron Disease
Mowat-Wilson Syndrome	Mowat-Wilson Syndrome
Mucolipidosis IV	Mucolipidosis IV
Mucopolysaccharidosis (not IVB)	Mucopolysaccharidosis (except for type IVB)
Mucopolysaccharidosis VII	Mucopolysaccharidosis VII (aka: Sly disease; Sly Syndrome; beta-glucuronidase deficiency)
Multiple Sulfatase Deficiency	Multiple Sulfatase Deficiency
Multisystem Developmental Disorder	Multisystem Developmental Disorder (aka: MSDD) (as defined within DC:0-5) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, psychiatrist, licensed clinical social worker...
Myasthenia Gravis	Myasthenia Gravis (familial infantile type)
Myhre Syndrome	Myhre Syndrome
Myoclonic Encephalopathy of Childhood	Myoclonic Encephalopathy of Childhood (Kinsbourne Syndrome)
Myopathy, Congenital	Myopathy, Congenital
Myopathy, Desmin Storage	Myopathy, Desmin Storage (aka: DSM)
Myopathy, Scapuloperoneal	Myopathy, Scapuloperoneal
Myotonic dystrophy type 1	Myotonic dystrophy type 1 (aka: DM; DM1; Curschmann-Batten-Steinert syndrome; Steinert disease; dystrophia myotonia; myotonia atrophica)
Myotubular Myopathy	Myotubular Myopathy
Neu Laxova Syndrome	Neu Laxova Syndrome
Neuropathy, Ataxia & Retinitis	Neuropathy, Ataxia and Retinitis Pigmentosa

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Neuropathy, Congenital Hypomyelination	Neuropathy, Congenital Hypomyelination (aka: Charcot-Marie-Tooth Type 4E; CHN; CMT4E; Congenital Dysmyelinating Neuropathy; Congenital Hypomyelinating Polyneuropathy; Congenital Hypomyelination; Congenital Hypomyelination Neuropathy; Congenital Hypomyeli...
Neuropathy, Giant Axonal	Neuropathy, Giant Axonal
Neuropathy, Hereditary Sensory Type I	Neuropathy, Hereditary Sensory, Type I (aka: HSAN1; HSN1; Hereditary Sensory and Autonomic Neuropathy Type 1)
Neuropathy, Hereditary Sensory Type II	Neuropathy, Hereditary Sensory, Type II
Neuropathy, Hereditary Sensory Type IV	Neuropathy, Hereditary Sensory, Type IV (aka: Familial Dysautonomia, Type II; Hereditary Sensory and Autonomic Neuropathy IV; HSAN IV; HSN IV; Insensitivity to Pain, Congenital, with Anhydrosis; CIPA; Neuropathy, Congenital Sensory, with Anhydrosis)
Neuropathy, Peripheral	Neuropathy, Peripheral
Niemann-Pick Disease	Niemann-Pick Disease (Classic Infantile and Juvenile)
Nonketotic Hyperglycinemia	Nonketotic Hyperglycinemia
Norrie's Syndrome	Norrie's Syndrome (aka: Anderson-Warburg Syndrome; Atrophia Bulborum Hereditaria; Episkopi Blindness; Fetal Iritis Syndrome; ND; NDP; Norrie Syndrome; Whitnall-Norman Syndrome)
Oculocerebral Syndrome with Hypopigmentation	Oculocerebral Syndrome with Hypopigmentation (aka: Cross Syndrome; Kramer Syndrome)
Oculocerebrocutaneous Syndrome	Oculocerebrocutaneous Syndrome (aka: Delleman Syndrome; Delleman-Oorthuys Syndrome; OCC Syndrome; OCCS; Orbital Cyst with Cerebral and Focal Dermal Malformations)
Ohtahara Syndrome	Ohtahara syndrome (aka: early infantile epileptic encephalopathy 1)
Olivopontocerbellar Atrophy	Olivopontocerebellar Atrophy, Hereditary (aka: Hereditary OPCA)
Opsoclonus-Myoclonus Syndrome	Opsoclonus-Myoclonus Syndrome (aka: Dancing Eyes-Dancing Feet; Kinsbourne Syndrome; Myoclonic Encephalopathy, Kinsbourne Type; Opsoclonic Encephalopathy)
Optico-Cochleo-Dentate Degen	Optico-Cochleo-Dentate Degeneration
Ornithine Transcarbamylase Deficiency	Ornithine Transcarbamylase Deficiency (aka: OTC)
Orocraniodigital Syndrome	Orocraniodigital Syndrome (aka: Juberg Hayward Syndrome; Cleft Lip/Palate with Abnormal Thumbs and Microcephaly; Cranio-Oro-Digital Syndrome; Digital-Oro-Cranio Syndrome)
Otopalatodgtl Syndrome Type I & II	Otopalatodigital Syndrome Type I and II (aka: Taybi Syndrome; Cranioorodigital Syndrome; FPO; Faciopalatoosseous Syndrome; OPD Syndrome)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Pachygyria	Pachygyria
Pallister Killian Mosaic Syndrome	Pallister Killian Mosaic Syndrome (aka: tetrasomy 12p; Killian/Teschler-Nicola Syndrome)
Pallister W Syndrome	Pallister W Syndrome
Paraplegia, Hereditary Spastic	Paraplegia, Hereditary Spastic
Partial Deletion of Chromosome 16p	Partial monosomy 16p
Partial Deletion of Chromosome 16q	partial monosomy 16q
Partial-Cerebellar Ataxia-MR	Partial-Cerebellar Ataxia-Mental Retardation
Peripheral Dys-Nasal Hypopl-MR	Peripheral Dysostosis-Nasal Hypoplasia-Mental Retardation (aka: PNP; Acrodysostosis)
Perisylvian Syndrome, Cong Bilat	Perisylvian Syndrome, Congenital Bilateral
Periventricular Leukomalacia	Periventricular Leukomalacia (PVL)
Pervasive Dev Disorder	Pervasive Developmental Disorder (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist...
Phenylketonuria (Untreated)	Phenylketonuria (Untreated)
Phocomelia Syndrome	Phocomelia Syndrome (aka: Roberts SC-Phocomelia Syndrome; Roberts Tetraphocomelia Syndrome; SC Phocomelia Syndrome; Pseudo-thalidomide Syndrome)
Phosphoglycerate Kinase Deficiency	Phosphoglycerate Kinase Deficiency (aka: Anemia, Hemolytic with PGK Deficiency; Erythrocyte Phosphoglycerate Kinase Deficiency; PGK; Phosphoglycerokinase)
Polymicrogyria, bilateral	Polymicrogyria, bilateral (aka: Congenital Bilateral Perisylvian Syndrome)
Pompe Disease	Pompe Disease
Porencephaly	Porencephaly
Potocki-Lupski Syndrome	Potocki-Lupski Syndrome (aka Chromosome 17p Duplication)
Prader-Willi Syndrome	Prader-Willi Syndrome
Progressive Cystic Encephalomalacia	
Progressive Multifocal Leukoencephalopathy	Progressive Multifocal Leukoencephalopathy
Progressive Myoclonus Epilepsy	Progressive Myoclonus Epilepsy
Pseudo Hurler Polydystrophy	Pseudo Hurler Polydystrophy (aka: mucolipidosis type III)

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Pyruvate Carboxylase Def	Pyruvate Carboxylase Deficiency
Pyruvate Dehydrogenase Def	Pyruvate Dehydrogenase Deficiency (PDCD)
Recombinant Chromosome 8 Syndrome	Recombinant Chromosome 8 Syndrome (aka: Rec8 Syndrome; San Luis Valley Syndrome)
Refsum Syndrome	Refsum Syndrome (aka: DOC 11 (Phytanic Acid Type); Disorder of Cornification 11 (Phytanic Acid Type); Heredopathia Atactica Polyneuritiformis; Hypertrophic Neuropathy of Refsum; Phytanic Acid Storage Disease)
Retinoic Acid Embryopathy	Retinoic Acid Embryopathy (aka: Accutane Embryopathy; Accutane (Fetal Effects of); Isotretinoin Embryopathy; Isotretinoin Teratogen Syndrome; Isotretinoin (Fetal Effects of); Fetal Retinoid Syndrome)
Rett Syndrome	Rett Syndrome (aka: Rett Disorder)
Rhombencephalosynapsis	(aka: RES)
Roberts Syndrome	Roberts Syndrome
Rosenberg Chutorian Syndrome	Rosenberg Chutorian Syndrome
Roussy Levy Syndrome	Roussy Levy Syndrome (aka: Charcot-Marie-Tooth Disease (Variant); Charcot-Marie-Tooth-Roussy-Levy Disease; Hereditary Areflexic Dystasia; Hereditary Motor Sensory Neuropathy; Hereditary Motor Sensory Neuropathy I; HMSN I)
Rubella, Congenital	Rubella, Congenital
Rubinstein Taybi Syndrome	Rubinstein Taybi Syndrome (aka: Broad Thumbs and Great Toes, Characteristic Facies, and Mental Retardation; Michail-Matsoukas-Theodorou-Rubinstein-Taybi Syndrome; RSTS; Rubinstein Taybi (RTS) Broad Thumb-Hallux syndrome; Rubinstein syndrome)
Sandhoff Disease	Sandhoff Disease
Sanfilippo Syndrome	Sanfilippo Syndrome (aka: Mucopolysaccharidosis Type III)
Santavuori Disease	Santavuori Disease (aka: CLN1; INCL; Infantile Finnish Type Neuronal Ceroid Lipofuscinosis; Balkan Disease; Infantile Neuronal Ceroid Lipofuscinosis; Infantile Type Neuronal Ceroid Lipofuscinosis; Neuronal Ceroid Lipofuscinosis Type 1; Santavuori-Haltia ...
Schindler Disease	Schindler Disease (aka: Alpha-N-Acetylgalactosaminidase Deficiency, Schindler Type; Alpha-NAGA Deficiency, Schindler Type; Lysosomal Alpha-N-Acetylgalactosaminidase Deficiency, Schindler Type; Alpha-Galactosidase B Deficiency; GALB Deficiency; Alpha-GalN...
Schinzel Giedion Syndrome	Schinzel Giedion Syndrome (aka: Schinzel-Giedion Midface-Retraction Syndrome)
Schizencephaly	Schizencephaly

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Schwartz Jampel Syndrome	Schwartz Jampel Syndrome
Scott Craniodigital Syndrome	Scott Craniodigital Syndrome
Seckel Syndrome	Seckel Syndrome
Shaken Baby Syndrome	Shaken Baby Syndrome (aka: Shaken Impact Syndrome; Shaken Infant Syndrome)
Simpson Dysmorphia Syndrome	Simpson Dysmorphia Syndrome (aka: Bulldog Syndrome; DGSX Golabi-Rosen Syndrome; Dysplasia Gigantism Syndrome, X-Linked; SDYS; SGB Syndrome; Simpson-Golabi-Behmel Syndrome)
Sirenomelia Sequence	Sirenomelia Sequence
Smith-Lemli-Opitz Syndrome	Smith-Lemli-Opitz Syndrome (aka: SLO)
Smith-Magenis Syndrome	Smith-Magenis syndrome
Sotos Syndrome	Sotos Syndrome
Spina Bifida (except occulta)	Spina Bifida (except for spina bifida occulta, in which the spinal cord is not exposed, but the vertebral bones aren't completely closed) (aka: meningomyelocele; myelomeningocele; MMC)
Spinal Muscular Atrophy	Spinal Muscular Atrophy
Stroke	Stroke (aka: acute neurologic syndrome; congenital stroke syndrome; neonatal stroke syndrome; brain attack; basal ganglia hemorrhage - changes in ischemia; Vein of Galen ischemia; perinatal stroke; Cerebral Venous Sinus Thrombosis; Cerebral Venous Thrombosis)
Subacute Sclerosing Panencephalitis	Subacute Sclerosing Panencephalitis
Succinic Semialdehyde Dehydrogenase	Succinic Semialdehyde Dehydrogenase Deficiency
Tay Sachs Disease	Tay Sachs Disease (aka: Cerebromacular Degeneration; GM2 Gangliosidosis, Type 1; Hexoaminidase Alpha-Subunit Deficiency (Variant B); Infantile Cerebral Ganglioside; Infantile Sipoidosis GM-2 Gangliosideosis (Type S); Lipidosis, ganglioside, infantile; Sp...
Timothy Syndrome	Timothy Syndrome
Triphosphate Isomerase Def	Triphosphate Isomerase Deficiency
Triple X Syndrome	Triple X syndrome (aka: Trisomy X; 47,XXX; Triplo X Syndrome)
Triploidy Syndrome	Triploidy Syndrome (aka: triploid syndrome)
Trisomy 13	Trisomy 13 (aka: Trisomy 13 - 15; Patau Syndrome)
Trisomy 18	Trisomy 18 (aka: Edwards Syndrome)
Trisomy 8	Trisomy 8 (aka: Trisomy 8 Syndrome; Trisomy 8 mosaic; Trisomy 8 mosaicism)
TTF-1 deletion	aka NKX2 deletion

Medical Diagnosis Name	The following conditions meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Turner Syndrome	Turner Syndrome (aka: 45X Syndrome; XO Syndrome)
Unbalanced Chromosome Translocation	Unbalanced Chromosome Translocation
Urea Cycle Defects (untreated)	Urea Cycle Defects (where the diagnosis is late, or there is no or inadequate treatment)
VACTERL w Hydrocephalus	VACTERL with Hydrocephalus (aka: VACTERL Association with Hydrocephalus; VACTERL-H Association; VATER Association with Hydrocephalus; VACTERL-H Association; VATER Association with Hydrocephalus)
Van der Knapp Syndrome	Van der Knapp Syndrome (aka: Megalencephalic leukoencephalopathy with subcortical cysts)
Visual Impair (not correctable)	Visual Impairment, Bilateral (not correctable with treatment, surgery, glasses, or contact lenses) (aka bilateral vision impairment bilateral vision loss, bilateral blindness)
WAGR Syndrome	WAGR Syndrome
Walker Warburg Syndrome	Walker Warburg Syndrome (aka: WWS)
Watson Syndrome	Watson Syndrome (aka: Pulmonic Stenosis with Cafe-Au-Lait Spots; Cafe-Au-Lait Spots with Pulmonic Stenosis)
Weaver Syndrome	Weaver Syndrome
Werdnig Hoffman Disease	Werdnig Hoffman Disease
West Syndrome	West Syndrome
Wieacker Syndrome	Wieacker Syndrome
Wiedemann Rautenstrauch Syndrome	Wiedemann Rautenstrauch Syndrome
Williams Syndrome	Williams Syndrome (aka: Williams-Beuren syndrome)
Wolf-Hirchhorn Syndrome	Wolf-Hirchhorn Syndrome (aka: Partial Monsomy 4p)
X-linked creatine deficiency	
XXXXX Syndrome	XXXXX Syndrome (aka: Penta X Syndrome)
XXYY Syndrome	XXYY Syndrome
XYY Syndrome	XYY Syndrome
Zellweger Syndrome	Zellweger Syndrome (aka: Bowen Syndrome; Cerebrohepatorenal Syndrome)

The following conditions do not meet the Type I Established Condition criteria in Montana

Medical Diagnosis Name	**The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19.** Diagnosis Details
Griscelli Syndrome, type 2	
3-Methylcrotonyl-Coenzyme A (CoA) Carboxylase Deficiency	
3rd Nerve Palsy	(aka: Third Nerve Palsy; Third Cranial Nerve Palsy; 3rd Cranial NervePalsy; Oculomotor Palsy; Oculomotor Nerve Palsy; Cranial Nerve III Palsy)
6th Nerve Palsy	(aka: Abducens palsy; Lateral rectus palsy; Cranial mononeuropathy VI)
Aarskog-Scott syndrome	
Abnormal Neurological Exam at Discharge from NICU	
Absent Septum Pellucidum	(aka: Absence Septum Pellucidum, Absent Cavum Septum Pellucidum)
Achondrogenesis I	Achondrogenesis I (Parenti-Fraccaro)
Achondrogenesis II	Achondrogenesis II (Langer-Saldino)
Achondroplasia	
Acrodysostosis	Acrodysostosis (aka: Arkless-Graham; Acrodysplasia; Maroteaux-Malamut; Peripheral Dysostosis-Nasal Hypoplasia-Mental Retardation Syndrome; PNM Syndrome)
Acute Lymphoid Leukemia	Acute Lymphoid Leukemia (aka: ALL; Leukemia)
Adams Oliver Syndrome	(aka: AOS; Absence Defect of Limbs, Scalp, and Skull; Congenital Scalp Defects with Distal Limb Reduction Anomalies; Aplasia Cutis Congenita with Terminal Transverse Limb Defects)
Adjustment Disorder	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Adrenal Hyperplasia, Congenital	AKA: 21 Hydroxylase Deficiency
Adrenal Hypoplasia	
Alagille Syndrome	(aka: AHD; Arteriohepatic Dysplasia; Cholestasis with Peripheral Pulmonary Stenosis; Syndromatic Hepatic Ductular Hypoplasia)
Albinism	(aka Ocular Cutaneous Albinism, Oculocutaneous Albinism, Ocular Albinism)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Alcohol-Related Neurodevelopmental Disorder	(aka: ARND; Fetal Alcohol Spectrum Disorder; FASD)
Alport Syndrome	(aka: Hematuria-Nephropathy Deafness; Hemorrhagic Familial Nephritis; Hereditary Deafness and Nephropathy; Hereditary Nephritis with Sensory Deafness; Hereditary Nephritis and Nerve Deafness)
Alstrom Syndrome	(aka: ALMS)
Amniotic Band Syndrome (affecting fetus or newborns)	
Amputation of leg at hip	Amputation of the leg at the hip
Anemia	
Aniridia	
Anophthalmia, unilateral	
Anoxic Insult to Brain	(aka anoxic brain injury)
Anterior Jaw Displacement	(aka: mandibular enlargement; prognathism)
Anterior segment dysgenesis	
Anxiety Disorder Not Otherwise Specified (NOS)	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Apert Syndrome	Apert Syndrome (aka: ACS 1; ACS I; Acrocephalosyndactyly, Type I; Syndactylic Oxycephaly)
APGAR score 3 or less @ 20 min	APGAR score of 3 or less at 20 minutes
APGAR score of five or less at five minutes	
Arachnoid Cyst	
Arnold-Chiari Malformation	AKA: Chiari 1 malformation
Arthrogryposis	Arthrogryposis (aka: arthrogryposis multiplex congenita; AMC; multiple congenital contractures)
Ataxia Telangiectasia	Ataxia Telangiectasia (aka: AT; Cerebello-Oculocutaneous Telangiectasia; Immunodeficiency with Ataxia Telangiectasia)
Auditory Neuropathy	(aka: Auditory neuropathy/auditory dyssynchrony; Auditory neuropathy/auditory dys-synchrony; AN/AD)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Aural Atresia, Bilateral or Unilateral	(aka: congenital aural atresia; CAA)
Bartter Syndrome	Bartter's syndrome (aka: Hypokalemic Alkalosis with Hypercalciuria)
Beals Syndrome	(aka: Beals-Hecht Syndrome; Arachnodactyly, Contractural Beals Type; Contractural Arachnodactyly, Congenital; CCA)
Beckwith Wiedemann Syndrome	(aka: BWS; Beckwith-Syndrome;Exomphalos-Macroglossia-Gigantism Syndrome; EMG Syndrome; Hypoglycemia with Macroglossia; Macroglossia-Omphalocele-Visceromegaly Syndrome; Omphalocele-Visceromegaly-Macroglossia Syndrome; Visceromegaly-Umbilical Hernia-Macrog...
Benign Enlargement of the Subarachnoid Space in infancy (aka: BESS)	
Biliary Atresia	
Bjornstad Syndrome	(aka: Deafness and Pili Torti, Bjornstad Type; Pili Torti and Nerve Deafness; Pili Torti-Sensorineural Hearing Loss)
Blue Diaper Syndrome	(aka: Drummond's Syndrome: Hypercalcemia, Familial, with Nephrocalcinosis and Indicanuria)
Bowen Hutterite Syndrome	(aka:Bowen-Conradi Hutterite Syndrome, Bowen-Conradi Syndrome, Hutterite Syndrome, Bowen-Conradi Type)
Brachial Plexus Palsy	(aka: Duchenne's Paralysis, Duchenne-Erb Paralysis, Duchenne-Erb Syndrome, Erb's Paralysis, Erb-Duchenne Palsy, Erb-Duchenne Paralysis, Upper Brachial Plexus Palsy, Erb-Duchenne Type, or Upper Brachial Plexus Paralysis, Erb-Duchenne Type; Brachial Plexus...
Branchio-oculo-facial syndrome	(aka:BOFS, Branchiooculofacial Syndrome, Hemangiomatous Branchial Clefts-Lip Pseudocleft Syndrome, Imperforate Nasolacrimal Duct, and Premature Aging Syndrome, Lip Pseudocleft-hemangiomatous Branchial Cyst Syndrome)
Bronchopulmonary Dysplasia (BPD)	
Carboxylase Deficiency, Multiple	(aka: Biotinidase Deficiency, Carboxylase Deficiency, Multiple, Holocarboxylase Synthetase Deficiency, MCD)
Carpenter Syndrome	(aka: ACPS II, Acrocephalopolysyndactyly Type II)
Cataracts, Congenital	

Medical Diagnosis Name	The following conditions <u>do not meet</u> the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Catel-Manzke Syndrome	(aka: Catel-Manzke Type Palatodigital Syndrome, Hyperphalangy-Clinodactyly of Index Finger with Pierre Robin Syndrome, Index Finger Anomaly with Pierre Robin Syndrome, Pierre Robin Syndrome with Hyperphalangy and Clinodactyly)
Caudal Regression Syndrome	(aka: Caudal Dysplasia; Caudal Dysplasia Sequence; Sacral Agenesis, Congenital; Sacral Regression)
Cerebral Bleed	(aka: intracerebral bleed, cerebral hemorrhage, intracerebral hemorrhage; parenchymal brain hemorrhage; periventricular hemorrhage)
Cerebral Palsy	Cerebral Palsy (aka: Ataxia Cerebral Palsy,Athetoid Cerebral Palsy, Congenital Cerebral Palsy, Diplegia of Cerebral Palsy,Hemiparesis of Cerebral Palsy, Hemiplegia of Cerebral Palsy, Hemiplegia, Hemiparesis, Postnatal Cerebral Palsy, Quadriparesis, Quadr...
Cerebrocostomandibular Syndrome	(aka: CCM Syndrome, CCMS, Rib Gap Defects with Micrognathia)
Chandler's Syndrome	(aka: Dystrophia Endothelialis Cornea, Iris Atrophy with Corneal Edema and Glaucoma)
Choanal Atresia (unilateral or bilateral)	
Choroid Plexus Cyst	
Chromosome 17q12 Deletion	
Chromosome 3q26.1-3q26.2 deletion	Chromosome 3q26.1-3q26.2 deletion
Chromosome 5q minus syndrome	Chromosome 5q minus syndrome
Chronic Ear Infections	
Citrullinemia	(aka: Citrullinuria)
Cleft Lip	
Cleft Palate and Cleft Lip	
Clinodactyly	
Clonus	(aka Clonospasm)
Club Foot (includes bilateral)	
Cogan Syndrome	
Cogan-Reese Syndrome	(aka: ICE Syndrome, Cogan-Reese Type; Iridocorneal Endothelial (ICE) Syndrome, Cogan-Resse Type; Iris Naevus Syndrome; Iris Nevus Syndrome)
Coloboma	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Complex congenital heart disease	
Cone Dystrophy	(aka: Combined Cone-Rod Degeneration; Cone-Rod Degeneration; Cone-Rod Degeneration, Progressive; Cone-Rod Dystrophy; Retinal Cone Degeneration; Retinal Cone Dystrophy; Retinal Cone-Rod Dystrophy)
Cong/acquired absence of limb	Congenital or acquired absence of limb (aka: severe limb deficiency; severe deficiency of extremity; severe form of Fibular Hemimelia; bilateral foot amputations at the ankle; complete absence of the fibula; tibular fibular hemimelia)
Congenital hypopituitarism and congenital disc pigmentation syndrome	
Congenital Nystagmus	
Congenital Syphilis	
Congenital Toxoplasmosis	
Congestive Heart Failure	(aka: CHF; Heart Failure)
Connexin 26	Connexin 26 gene mutation (with mutation on both copies and a diagnosed unilateral or bilateral hearing loss)
Connexin 30 gene mutation	
Conradi-Hunermann syndrome	(aka: Chondrodysplasia Punctata, X-linked Dominant Type; Chondrodystrophia Calcificans Congenita; Conradi Disease; Dysplasia Epiphysialis Punctata)
Cortical Dysplasia	Cortical Dysplasia
Cortical Visual Impairment	Cortical Visual Impairment (aka CVI and others)
Craniosynostosis without Radial Defects	
Crigler-Najjar Syndrome	(aka: Bilirubin Glucuronosyltransferase Deficiency Type I; Congenital Familial Nonhemolytic Jaundice Type I; Uridine Diphosphate Glucuronosyltransferase, Severe Def. Type I)
Crouzon Syndrome	aka: Craniofacial Dysotosis; Craniostenosis (Crouzon Type); Crouzon Craniofacial Dysostosis)
Cystic Fibrosis	
Cytomegalovirus	Cytomegalovirus (aka: CMV; Cytomegalic Inclusion Disease; Giant Cell Inclusion Disease (CID); Human Cytomegalovirus Infection; Salivary Gland Disease, CMV Type)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Dejerine Sottas Disease	Dejerine Sottas Disease (aka: Hereditary Motor Sensory Neuropathy Type III; HSMN Type III; Hypertrophic Interstitial Neuritis; Hypertrophic Interstitial Neuropathy; Hypertrophic Interstitial Radiculoneuropathy, Onion-Bulb Neuropathy)
Depression of Infant and Early Childhood: Type II - Depressive Disorder NOS	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Depression: Type I-Major Dep	Depression of Infant and Early Childhood: Type I - Major Depression (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professi...
Deprivation/Maltreatment Dis	Deprivation/Maltreatment Disorder (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologis...
Diaphragmatic Hernia	
Diaphragmatic Paralysis	aka: Bilateral Diaphragmatic Paralysis)
Diastematomyelia	
Diastrophic Dysplasia	Diastrophic Dysplasia (aka: DD; DTD; Diastrophic Dwarfism; Diastrophic Nanism Syndrome)
Disorders of Relating and Communicating	(as defined within DC:0-5, and diagnosed by specially-qualified professional as noted)
Drug Exposure, Prenatal	(aka neonatal abstinence syndrome)
Duane Syndrome	
Dysphagia	Dysphagia
Dystonia Musculorum Deformans	Dystonia Musculorum Deformans (aka: Torsion Dystonia)
Eagle-Barrett syndrome	(aka: Prune Belly Syndrome)
Eales Disease	(aka: Eales Retinopathy, Idiopathic Peripheral Periphlebitis)
Ear, Patella, Short Stature Syndrome	(aka: EPS; Meier-Gorlin Syndrome; Microtia, Absent Patellae, Micrognathia Syndrome)
Ectrodactyly Ectodermal Dysplasia Cleft Lip/Palate	(aka: EEC Syndrome; Ectrodactyly-Ectodermal Dysplasia-Clefting Syndrome)
Ehlers Danlos Syndrome	(aka: E-D Syndrome; ED Syndrome)
Ellis Van Creveld Syndrome	(aka: Chondroectodermal Dysplasia; Mesoectodermal Dysplasia)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Empty Sella Syndrome	(aka: Empty Sella Turcica)
Encephalitis, Japanese	(aka: JE, Japanese B Encephalitis, Russian Autumnal Encephalitis, Summer Encephalitis)
Epidermolysis Bullosa	
Epidermolytic Hyperkeratosis	
Epilepsy	Epilepsy
Esophageal Atresia	
Exostoses, Multiple	
Extraocular Fibrosis Syndrome	(aka: Congenital Fibrosis of the Extraocular Muscles; CFEOM)
Fabry Disease	(aka:Alpha-Galactosidase A Deficiency; Anderson-Fabry Disease; Angiokeratoma Corporis Diffusum; Angiokeratoma Diffuse; Ceramide Trihexosidase Deficiency; GLA Deficiency; Hereditary Dystopic Lipidosis)
Facial Palsy	(aka: Bell's Palsy; Facial Nerve Palsy; 7th Nerve Palsy)
Facioscapulohumeral Muscular Dystrophy	(aka: FMD; FSH; FSHD;Facio-Scapulo-Humeral Dystrophy; Muscular Dystrophy, Facioscapulohumeral;Muscular Dystrophy, Landouzy Dejerine)
Failure to Thrive	
Familial Dysautonomia	(aka: Riley-Day Syndrome; FD; Hereditary Sensory and Autonomic Neuropathy, Type III (HSAN, Type III); Hereditary Sensory Neuropathy Type III; HSAN III; HSN III)
Familial exudative vitreoretinopathy	(aka: FEVR)
Fanconi Syndrome	(aka: de Toni-Fanconi syndrome)
Farber Disease	Farber Disease (aka: Farber's lipogranulomatosis; ceramidase deficiency)
Feeding Disorder of CG-Inf Rec	Feeding Disorder of Caregiver-Infant Reciprocity (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physici...
Feeding Disorder of State Reg	Feeding Disorder of State Regulation (as defined within DC:0-5 and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psycholo...
Feeding Disorder w/ GI Tract	Feeding Disorder Associated with Insults to the Gastrointestinal Tract (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health profe...

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Feeding Disorder w/ Med Cond	Feeding Disorder Associated with Concurrent Medical Condition (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (...
Feeding Disorder, Sensory Food Aversions	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Femoral Facial Syndrome	(aka: Femoral Dysgenesis, Bilateral; Femoral Dysgenesis, Bilateral-Robin Anomaly; Femoral Hypoplasia-Unusual Facies Syndrome)
Fetal Alcohol Spectrum Disorder	(aka: Alcohol-related neurodevelopmental disorder; alcohol-related neurobehavioral disorder; fetal alcohol effects)
Fetal Valproate Syndrome	(aka: Dalpro, (Fetal Effects From); Depakene, (Fetal Effects From); Depakote Sprinkle (Fetal Effects From); Depakote, (Fetal Effects From); Divalproex, (Fetal Effects From); Epival (Fetal Effects From); Fetal Anti-Convulsive Syndrome; Myproic Acid (Fetal...
Fibromatosis, Congenital Generalized	(aka: CGF; Infantile Myofibromatosis; IM)
Forbes Disease	(aka: Amylo-1,6-Glucosidase Deficiency; Cori Disease; Debrancher Deficiency; Glycogen Storage Disease III; Glycogenosis Type III; Limit Dextrinosis)
Fracture of Vertebral Column with Spinal Cord Injury	
Fraser Syndrome	(aka: Cryptophthalmos Syndrome; Cryptophthalmos-Syndactyly Syndrome; Cyclopism Fraser-Francois Syndrome; Meyer-Schwickerath's Syndrome; Ulrich-Feichtiger Syndrome)
Freeman Sheldon Syndrome	(aka: Craniocarpotarsal dystrophy (dysplasia); DA2A; Distal arthrogryposis type 2A; FSS; Whistling face syndrome; Whistling face-windmill vane hand syndrome)
Frontofacionasal Dysplasia	(aka: Cleft Lip-Palate, Blepharophimosis, Lagophthalmos, and Hypertelorism; Facio-Fronto-Nasal Dysplasia; Frontofacionasal Dysostosis; Nasal-fronto-faciodysplasia)
Frontonasal Dysplasia	(aka: FND; Median Cleft Face Syndrome)
Galloway-Mowat Syndrome	(aka: Galloway Syndrome; Hiatal Hernia-Microcephaly-Nephrosis, Galloway Type; Microcephaly-Hiatal Hernia-Nephrosis, Galloway Type; Microcephaly-Hiatal Hernia-Nephrotic Syndrome; Nephrosis-Microcephaly Syndrome; Nephrosis-Neuronal Dysmigration Syndrome)
Gastroschisis	(aka: Abdominal Wall Defect; Aparoschisis)
Gaucher Disease, Type I	(aka: Sphingolipidosis 1)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Generalized Anxiety Disorder	Generalized Anxiety Disorder (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, ps...
Glutaricaciduria I	(aka: Dicarboxylic Aminoaciduria; GA I; Glutaric Acidemia I; Glutaric Aciduria I; Glutaricacidemia I; Glutaryl-CoA Dehydrogenase Deficiency; Glutaurate-Aspartate Transport Defect)
Glutaricaciduria II	(aka: Electron Transfer Flavoprotein, Deficiency of; Electron Transfer Flavoprotein: Ubiquinone Oxidoreductase, Deficiency of; GA II; Glutaric Acidemia II; Glutaric Aciduria II; Glutaricacidemia II; MADD; Multiple Acyl-Co-A Dehydrogenation Deficiency)
Goldenhar Syndrome	(aka: Oculo-Auriculo-Vertebral/OAV syndrome)
Goltz Syndrome	(aka: Gorlin Syndrome; nevoid basal cell carcinoma syndrome)
Goodman Syndrome	(aka: ACPS IV; Acrocephalopolysyndactyly Type IV)
Gordon Syndrome	(not the other "Gordon Syndrome with hypertension, hyperkalaemia, and normal glomerular filtration rate") (aka: Arthrogryposis Multiplex Congenita, Distal, Type IIA; Camptodactyly-Cleft Palate-Clubfoot; Distal Arthrogryposis, Type IIA)
Gorlin-Chaudhry-Moss Syndrome	(aka: Craniofacial Dysostosis-PD Arteriosus-Hypertrichosis-Hypoplasia of Labia; Craniosynostosis-Hypertrichosis-Facial and Other Anomalies; GCM Syndrome)
Gottron Syndrome	(aka: H. Gottron's syndrome; acrogeria, Gottron type; familial acrogeria; familial acromicria)
Greig Cephalopolysyndactyly Syndrome	(aka: Frontodigital Syndrome (obsolete); GCPS; Hootnick-Holmes Syndrome (obsolete); Polysyndactyly with Peculiar Skull Shape; Polysyndactyly-Dysmorphic Craniofacies, Greig Type)
Griscelli Syndrome, type 3	
Guillain Barre Syndrome	(aka: Acute Autoimmune Peripheral Neuropathy; Acute Immune-Mediated Polyneuropathy ; Acute Inflammatory Demyelinating Polyneuropathy; Acute Inflammatory Demyelinating Polyradiculoneuropathy; Acute Inflammatory Neuropathy; Acute Inflammatory Polyneuropath...
Hajdu Cheney Syndrome	(aka: HCS)
Hallermann Streiff Syndrome	
Hand-Schuller-Christian	(aka: Histiocytosis X; Hemophagocytic Lymphohistiocytosis, Familial, 1)
Hanhart Syndrome	(aka: aglossia-adactylia; hypoglossia-hypodactylia syndrome; peromelia with micrognathia)
Hartnup Disease	
Hay-Wells Syndrome	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Hearing Loss (diagnosis with unilateral permanent hearing loss)	
Heart transplant	Heart transplant
Hemihyperplasia	(aka: hemihypertrophy)
Hemophilia	
Hepatoblastoma	
Hereditary Hyperphosphatasia	
Hermansky Pudlak Syndrome	
Herpes-Zoster	(aka: Hunt Syndrome)
Heterotopia	(aka Subependymal Nodular Heterotopia)
Hirschsprung Disease	AKA: Hirschsprung's Disease; intestinal failure; near total agangliosis
Holt Oram Syndrome	(aka: HOS)
Homocystinuria	Homocystinuria
Horner's Syndrome	(aka: Horner Syndrome)
Huntington's Disease	
Hydrops	(aka: hydrops fetalis; erythroblastosis fetalis)
Hyperbilirubinemia	(aka: Jaundice of the newborn; Neonatal hyperbilirubinemia)
Hyperexplexia	
Hypertonia	
Hypophosphatasia	
Hypophosphatemia, Familial	
Hypoplastic Left Heart Syndrome	
Hypothyroidism, Congenital	
Hypotonia, Congenital, Non-Benign Form	
Ichthyosis Congenita	(aka: Ichthyosis Lamellar; Autosomal Recessive Congenital Ichthyosis, Ichthyosis, Harlequin Type)
Ichthyosis, Chanarin Dorfman Syndrome	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Infantile Anorexia	Infantile Anorexia (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, psychiatrist...
Infantile Botulism	
Infantile Spasms, Epilepsy	Infantile Spasms, Epilepsy (aka: Infantile Myoclonic Seizures, Infantile Spasm; Hypsarrhythmia)
Intestinal malrotation	(aka: malrotation of the intestine)
Intraventricular Cyst	
Intraventricular Hemorrhage (IVH) Grade I	(aka: Germinal Matrix Hemorrhage; Grade I IVH; Grade I periventricular hemorrhage, Grade I PVH)
Intraventricular Hemorrhage, Grade II	(aka: IVH, periventricular hemorrhage, PVH) Grade 2
IPEX Syndrome	(aka: immune dysregulation, polyendocrinopathy, enteropathy, X-linked syndrome)
IRF6-Related Disorders	
Isovaleric Acidemia	
IUGR/SGA 33 wks & < 1,325g	IUGR (qualifies children under two years only - evidence shows the child was born at 33 weeks gestational age and weighing 1,325 grams (2 pounds, 15 ounces) or less at birth or shortly after birth) (aka: for the purposes of determining EI Colorado eligibi...
IUGR/SGA 34 wks & < 1,500 g	IUGR (qualifies children under two years only - evidence shows the child was born at 34 weeks gestational age and weighing 1,500 grams (3 pounds, 5 ounces) or less at birth or shortly after birth) (aka: for the purposes of determining EI Colorado eligibil...
IUGR/SGA 35 wks & < 1,700 g	IUGR (qualifies children under two years only - evidence shows the child was born at 35 weeks gestational age and weighing 1,700 grams (3 pounds, 12 ounces) or less at birth or shortly after birth) (aka: for the purposes of determining EI Colorado eligibi...
IUGR/SGA 36 wks & < 1,875 g	IUGR (qualifies children under two years only - evidence shows the child was born at 36 weeks gestational age and weighing 1875 grams (4 pounds, 2 ounces) or less at birth or shortly after birth) (aka: for the purposes of determining EI Colorado eligibili...
IUGR/SGA 37-40 wk & < 2,000 g	IUGR (qualifies children under two years only - evidence shows the child was born between 37 and 40 weeks gestational age and weighing less than 2,000 grams (4 pounds, 5 ounces) or less at birth or shortly after birth) (for the purposes of determining EI ...

Medical Diagnosis Name	The following conditions <u>do not meet</u> the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Ivemark Syndrome	(aka: Asplenia Syndrome, Bilateral Right-Sidedness Sequence, Splenic Agenesis Syndrome)
Jackson-Weiss Syndrome	
Jansen Type Metaphyseal Chondrodysplasia	
Jarcho-Levin Syndrome	
Jejunal atresia	
Jervell & Lange-Nielsen Syndrome	Jervell and Lange-Nielsen Syndrome
Job Syndrome	
Juvenile Myelomonocytic Leukemia	
Juvenile Rheumatoid Arthritis	(aka: JRA)
Kabuki Make-up Syndrome	Kabuki Make-up Syndrome
Kartagener Syndrome	
Kawasaki Disease	
Kearns Sayre Syndrome	
Kenny-Caffey Syndrome	
Keratosis Follicularis	
Kernicterus	Kernicterus
Klippel-Feil Syndrome	(aka: Klippel-Feil Sequence)
Klippel-Trenaunay Syndrome	(aka: Klippel-Trenaunay-Weber Syndrome)
Kniest Dysplasia	
Laband Syndrome	
LADD Syndrome	
Lambert-Eaton Myasthenic Syn	Lambert-Eaton Myasthenic Syndrome
Langerhans Cell Histiocytosis	
Laron Syndrome	
Larsen Syndrome	
Laryngomalacia	
Laryngotracheoesophageal cleft	(aka: LTEC)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Laurence Moon Syndrome	Laurence Moon Syndrome (aka: Adipogenital-Retinitis Pigmentosa Syndrome; Laurence Syndrome; LM Syndrome)
LBW < 400 grams	LBW < 400 grams at birth or shortly after birth (qualifies children under two years only - infant born at any gestational age (for the purposes of determining eligibility for EI Colorado LBW can also be referred to as SGA or IUGR)
LBW 1,000 - 1,199 grams	LBW 1,000 - 1,199 grams at birth or shortly after birth (qualifies children under two years only - infant born at any gestational age (for the purposes of determining eligibility for EI Colorado LBW can also be referred to as SGA or IUGR)
LBW 400 - 599 grams	LBW 400 - 599 grams at birth or shortly after birth (qualifies children under two years only - infant born at any gestational age (for the purposes of determining eligibility for EI Colorado LBW can also be referred to as SGA or IUGR)
LBW 600 - 799 grams	LBW 600 - 799 grams at birth or shortly after birth (qualifies children under two years only - infant born at any gestational age (for the purposes of determining eligibility for EI Colorado LBW can also be referred to as SGA or IUGR)
LBW 800 - 999 grams	LBW 800 - 999 grams at birth or shortly after birth (qualifies children under two years only - infant born at any gestational age (for the purposes of determining eligibility for EI Colorado LBW can also be referred to as SGA or IUGR)
Lead level >10 ug/dL	
Leber Congenital Amaurosis	Leber Congenital Amaurosis
Legg Calve Perthes Disease	
LEOPARD Syndrome Leprechaunism	
Leri Pleonosteosis	
Leukemia	
Leukocytosis	
Linear Sebaceous Nevus Sequence	Linear Sebaceous Nevus Sequence (aka: Linear Sebaceous Nevus Syndrome; Sebaceous Nevus Syndrome; Linear Epidermal Nevus Syndrome; LEN Syndrome; Jadassohn nevus phakomatosis; JNP)
Loeys-Dietz Syndrome	
Loken Senior Syndrome	
Lumbosacral Agenesis	
Lyme Disease	
Lymphadenopathy	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Macrocephaly	
Macroglossia	
Macular lesion (of the eye)	(aka: lesion of the macula)
Maffucci Syndrome	
Marcus Gunn Phenomenon	
Marfan Syndrome	
Maroteaux Lamy Syndrome	
Marshall Syndrome	
Maxillofacial Dysostosis	Maxillofacial Dysostosis
Maxillonasal Dysplasia, Binder Type	
MCADD	aka: MCAD Deficiency
McCune Albright Syndrome	
McKusick Type Metaphyseal Chondrodysplasia	AKA cartilage hair hypoplasia syndrome
Megacystis Microcolon Intestinal Hypoperistalsis Syndrome	(aka: Berdon Syndrome; Megacystis Microcolon Intestinal Hydronephrosis Syndrome; Neonatal Hollow visceral myopathy)
Meige Syndrome	
Melkersson Rosenthal Syndrome	
Melnick Needles Syndrome	
Melorheostosis	
Meningitis	Meningitis (including but not limited to the following types: Bacterial, Infantile, Neonatal, Meningococcal, Pneumococcal, Tuberculous, Epidemic Cerebrospinal, Pyogenic; Waterhouse-Friederichsen Syndrome)
Metabolic Acidosis	
Metaphyseal Chondrodysplasia, Schmid Type	
Metatarsus adductus	(aka: Metatarsus varus; Forefoot varus)
Metatropic Dysplasia I	

Medical Diagnosis Name	The following conditions <u>do not meet</u> the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Methylmalonic Aciduria without Glycinemia, Group 5	(aka: Methylmalonic Acidemia; ketotic hyperglycinemia)
Micrognathia	
Micromelia	(aka: nanomelia)
Microphthalmia	
Microtia and Atresia	(aka: Microtia/Atresia; Microtia-Atresia; Microtia and Aural Atresia; Microtia)
Microvillus Inclusion Disease	
Miller Syndrome	(aka: Postaxial Acrofacial Dysostosis Syndrome)
Missing Fingers	(aka: missing thumbs; absent fingers; absent thumbs)
Mixed Disorder of Emotional Expressiveness	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Moebius Sequence	Moebius Sequence
Morquio Syndrome	
Motonia Congenita - Thomsen Disease	
Moyamoya Syndrome	
Mulibrey Nanism Syndrome (Perheentupa Syndrome)	
Mulvihill Smith Syndrome	
MURCS Association	
Murmur	
Muscular Dystrophies, Limb Girdle	
Muscular Dystrophy, Becker	
Muscular Dystrophy, Emery Dreifuss	
Muscular Dystrophy, Oculogastrointestinal	
Myoclonus, General	
Myopathy, Cong, Batten Turner	Myopathy, Congenital, Batten Turner Type
Myositis Ossificans Progressiva	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Myotonia Congenita	
Nager Syndrome	
Nail Patella Syndrome	
Necrotizing Enterocolitis (NEC)	(aka: short gut/bowel syndrome)
Nemaline Myopathy	Nemaline Myopathy
Neonatal Abstinence Syndrome	Neonatal Abstinence Syndrome
Neonatal Cerebral Depression	
Neonatal Hemochromatosis (NH)	
Neonatal Herpes Simplex (HSV)	
Neonatal Seizures	
Nephrotic Syndrome	(Aka: Membranous glomerulonephritis; Membranoproliferative glomerulonephritis; Mesangiocapillary glomerulonephritis)
Neuroblastoma	
Neuromyotonia	
Neutropenia	
Nezelof Syndrome	
NF1-Neurofibromatosis	(aka: NF-1; NF1; Von Recklinghausen Disease)
NF2-Bilateral Acoustic Neurofibromatosis	(aka Neurofibromatosis Type 2 (NF-2))
Night-Waking Disorder	(Night-Waking Protodyssomnia) (as defined within DC:0-5, and diagnosed by specially-qualified professional)
Noonan Syndrome	
Ochoa Syndrome	
Ocular Motor Apraxia, Cogan Type	(aka: Congenital Oculomotor Apraxia; oculomotor apraxia, Cogan type; COMA; saccade initiation failure, congenital)
Oculo-Dento-Digital Dysplasia	(aka: Dento-Oculo-Osseous Dysplasia ; ODD Syndrome ; ODDD ; ODOD ; Oculo Dento Digital Dysplasia ; Oculo-Dento-Osseous Dysplasia ; Oculodentodigital Dysplasia ; Osseous-Oculo-Dento Dysplasia)
Ollier Disease	(aka: Dyschondroplasia; Enchondromatosis; Multiple Cartilaginous Enchondroses; Multiple Enchondromatosis)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Omphalocele	
Opitz G/BBB Syndrome	Opitz G/BBB Syndrome (aka: BBBG Syndrome; Hypertelorism with Esophageal Abnormalities and Hypospadias; Hypertelorism-Hypospadias Syndrome; Hypospadias-Dysphagia Syndrome; Opitz BBB Syndrome; Opitz BBB/G Compound Syndrome; Opitz BBBG Syndrome; Opitz G Syn...
Optic atrophy	(aka: optic nerve atrophy)
Oral-Facial-Digital Syndrome	Oral-Facial-Digital Syndrome (aka: OFD Syndrome; Orofaciodigital Syndrome)
Organ Failure (other than brain/CNS)	(aka: end-organ failure; multiple organ failure; mulitple system organ failure; multiple organ system failure; multiple organ dysfunction syndrome (MODS))
Orotic Aciduria	
OSMED, Heterozygous	(aka: Arthro-Ophthalmopathy; Epiphyseal Changes and High Myopia; Ophthalmoarthropathy Weissenbacher-Zweymuller Syndrome; Hereditary Arthro-Ophthalmopathy; Stickler syndrome type I; Stickler syndrome type II)
OSMED, Homozygous	(aka: oto-spondylo-megaepiphyseal dysplasia, autosomal recessive; oto-spondylo-megaepiphyseal dysplasia, homozygous; Nance-Sweeney syndrome)
Osteodystrophy, Congenital	
Osteogenesis Imperfecta	
Pachydermoperiostosis	
Paget's Disease	
Pallister Hall Syndrome	
Panhypopituitarism	
Papillitis	
Papillon Lefevre Syndrome	
Paramyotonia Congenita	
Paraneoplastic Neurologic Syndromes	
ParentInfRel-GAS of 40 or less	Parent Infant Relationship-Global Assessment Scale (PIR-GAS) of 40 or less (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health p...
Parry Romberg Syndrome	
Pars Planitis	
Parsonage Turner Syndrome	

Medical Diagnosis Name	The following conditions <u>do not meet</u> the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Pediatric Cardiomyopathy	
Pentalogy of Cantrell	
PEPCK Deficiency	
Peters Anomoly	(aka: bilateral anterior segment dysgenesis)
Peutz Jeghers Syndrome	
Pfeiffer Syndrome Type I	
PHACES Syndrome	(aka: PHACE syndrome)
Pharyngeal Dysphagia	
Pica	
Pick Disease	Pick's Disease
Pierre-Robin Sequence	(aka Perre-Robin Syndrome)
PIGA	PIGA (aka: phosphatidylinositol glycan anchor biosynthesis class A gene mutation)
Pineal Cysts, Symptomatic	
Plagiocephaly	
Pleuropulmonary Blastoma	
POEMS Syndrome	
Poland Syndrome	
Polychondritis	
Polycystic Kidney Disease	
Polydactyly (all types: postaxial, preaxial, and central)	
Polymicrogyria, unilateral focal	
Polymyalgia Rheumatica	
Polymyositis	
Porphyria, Acute Intermittent	
Porphyria, Congenital Erythropoietic	
Porphyria, Hereditary Coproporphyria	
Porphyria, Variegate	
Positive Toxicology Screen	
Posterior Uveitis	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Posttraumatic Stress Disorder	Posttraumatic Stress Disorder (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, p...
Potter sequence	Potter Sequence, Potter syndrome, and oligohydramnios sequence
Presumed septic thrombus	
Primary alveolar hypoventilation	(aka Ondine's Curse)
Progeria, Hutchinson Gilford	
Progressive Osseous Heteroplasia (POH)	
Prolonged Bereavement/Grief Reaction	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Proteus Syndrome	
Proximal Femoral Focal Deficiency	Proximal Femoral Focal Deficiency
Prune Belly Syndrome	(aka Eagle-Barrett syndrome)
Pseudo-achondroplasia	(aka: Pseudoachondroplastic Dysplasia)
Pseudohypoparathyroidism	
Pseudoxanthoma Elasticum (PXE)	
PTEN Hamartoma Tumor Syndrome	
Pterygium Syndrome, Multiple	
Ptosis	
Pulmonary Eventration	aka: Eventration of the Diaphragm
Pyknodysostosis	
Pyloric stenosis	
Pyridoxine-Dependent Seizures (PDS)	
Pyruvate Kinase Deficiency	
Rabson-Mendenhall Syndrome	
Ramsay-Hunt Syndrome	
Rapp Hodgkin Syndrome	(aka: Ectodermal dysplasia, Rapp-Hodgkin type; Ectodermal dysplasia, anhidrotic, with cleft lip and cleft palate; RHS; Rapp-Hodgkin (hypohidrotic) ectodermal dysplasia syndrome)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Reflex Sympathetic Dystrophy Syndrome	(aka: Algodystrophy ; Algoneurodystrophy ; Causalgia Syndrome (Major) ; Complex Regional Pain Syndrome; RSDS ; Reflex Neurovascular Dystrophy ; Sudeck's Atrophy)
Reg Dis Sens Proc: Hypersensitive	Regulation Disorders of Sensory Processing: Hypersensitive, Type A: Fearful/Cautious (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behaviora...
Reg Dis Sens Proc: Hypo/Under responsive	Regulation Disorders of Sensory Processing: Hyposensitive/Under responsive (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health pr...
Reg Dis Sens Proc: SSS/Imp	Regulation Disorders of Sensory Processing: Sensory Stimulation-Seeking/Impulsive (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral h...
Regulation Disorders of Sensory Processing: Hypersensitive, Type B: Negative	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Respiratory Distress Syndrome, Infant	(aka: Hyaline Membrane Disease; IRDS; Infantile Respiratory Distress Syndrome)
Retinal Hemorrhage	
Retinitis Pigmentosa	(aka: Progressive Pigmentary Retinopathy and RP)
Retinoblastoma	
Retinopathy of Prematurity	(aka: ROP and Retrolental Fibroplasia (obsolete))
Retinopathy, Arteriosclerotic	(aka: Arteriosclerosis, Retina)
Retinopathy, Diabetic	
Retinopathy, Hypertensive	
Retinoschisis	(aka: Congenital Retinal Cyst; Congenital Vascular Veils in the Retina; Giant Cyst of the Retina; Vitreoretinal Dystrophy) ((Disorder Subdivisions: Blessig Cysts; Congenital Retinoschisis; Familial Foveal Retinoschisis; Iwanoff Cysts; Peripheral Cystoi...
Reye Syndrome	(aka: Fatty Liver with Encephalopathy; RS; Reye's Syndrome)
Rh incompatibility	
Rieger Syndrome	(aka: Axenfeld-Rieger Syndrome; Goniodysgenesis-Hypodontia; Iridogoniodysgenesis With Somatic Anomalies; RGS)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Robinow Syndrome	(aka: Acral Dysostosis with Facial and Genital Abnormalities; Costovertebral segmentation defect with mesomelia (formerly); Fetal Face Syndrome; Robinow Dwarfism)
Romano Ward Syndrome	(aka: Autosomal Dominant Long QT Syndrome; LQTS1; Long QT Syndrome Type 1; Long QT Syndrome without Deafness; RWS; Romano-Ward Long QT Syndrome; Ward-Romano Syndrome)
Rosai-Dorfman Disease (RDD;SHML;Sinus Histiocytosis w/ Massive Lymphadenopathy)	
Roseola Infantum	(aka: Exanthem Subitum; Pseudorubella; Sixth Disease)
Rothmund Thomson Syndrome	(aka: Poikiloderma Atrophicans and Cataract; Poikiloderma Congenitale; RTS)
Rumination Syndrome	
Russell Silver Syndrome	(aka: RSS; Russell Syndrome; Russell-Silver Dwarfism; SRS; Silver Syndrome; Silver-Russell Dwarfism; Silver-Russell Syndrome)
Saethre Chotzen Syndrome	
Sakati Syndrome	
Sandifer Syndrome	
Schinzel Syndrome	
Scleroderma	
Scoliosis	
Seizures	AKA: Focal Seizures
Separation Anxiety Disorder	Separation Anxiety Disorder (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, psy...
Sepsis	
Septo-optic Dysplasia	(aka: Optic Nerve Hypoplasia (ONH); Pituitary Hypoplasia)
Setleis Syndrome	
Severe Combined Immunodeficiency (SCID)	
Short Chain Acyl CoA Dehydrogenase Deficiency (SCAD)	
SHORT Syndrome	SHORT Syndrome

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Shwachman Syndrome	
Simian B Virus Infection	
Singleton Merten Syndrome	Singleton Merten Syndrome (aka: Merten Singleton Syndrome)
Sjogren Syndrome	
Skeletal Dysplasia	Congenital skeletal dysplasia. (aka: Skeletal Dysplasia)
Sleep Behavior Disorder	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Sleep-Onset Disorder (Sleep-Onset Progodyssomnia)	(as defined within DC:0-5, and diagnosed by specially-qualified professional)
Sneddon Syndrome	
Social Anxiety Disorder	Social Anxiety Disorder (Social Phobia) (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psych...
Specific Phobia	Specific Phobia (as defined within DC:0-5, and diagnosed by specially-qualified professional as noted) [The person making the diagnosis must be a practicing medical or mental/behavioral health professional (i.e., physician, psychologist, psychiatrist, l...
Spinal Cord Injury	(aka: SCI) (due to the wide range of severity and outcomes, functional impairment will be used to determine eligibility)
Spinal Stenosis	(aka: Congenital spinal dislocation/kyphosis @T7-T8 with spinal cord compression)
Split Hand/Split Foot Malformation	
Spondyloepiphyseal Dysplasia, Congenital	Spondyloepiphyseal Dysplasia, Congenital
Spondyloepiphyseal Dysplasia Tarda	
Sprengel Deformity	
Stevens Johnson Syndrome	
Stickler Syndrome	
Sturge-Weber Syndrome	Sturge-Weber Syndrome
Stuve-Wiedemann Syndrome	
Subdural Hematoma	
Subgaleal Hemorrhage	
Summitt Syndrome	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Sydenham Chorea	Sydenham Chorea (aka: Sydenham's Chorea)
Syndactyly	(aka: polysyndactyly; webbing of the fingers and/or toes)
Syringobulbia	
Systemic Herpes Simplex Virus (HSV) Infection	(aka: disseminated herpes simplex infection)
Telecanthus with Associated Abnormalities	
Tethered Spinal Cord Syndrome	
Tetrahydrobiopterin Deficiency	(including the following subdivisions: Tetrahydrobiopterin Synthesis, GTP Cyclohydrolase I (GTPCH) Deficiency, 6-Pyruvoyl Tetrahydropterin Synthase (PTPS) Deficiency [aka: Hyperphenylalanemia with Biopterin Defect, PTPS Deficiency], Tetrahydrobiopterin R...
Tetralogy of Fallot	
Thalamic Syndrome (Dejerine Roussy)	
Three M Syndrome	
Thrombocytopenia	
Tietze Syndrome	
Tolosa Hunt Syndrome	
TORCH Syndrome	
Torticollis	
Tourette Syndrome	
Townes Brocks Syndrome	
Toxic Epidermal Necrolysis	
Toxic Optic Neuropathy	(aka: Toxic/Nutritional Optic Neuropathy)
Tracheoesophageal Fistula	(aka, Esophageal Atresia and Tracheoesophageal Fistula; Esophageal Fistula)
Tracheomalacia	
Tracheostomy	(aka: trach; trach tube; tracheostomy tube)
Transverse Myelitis	Transverse Myelitis (aka: Cervical Transverse Myelitis)
Treacher Collins Syndrome	
Tricho Dento Osseous Syndrome	

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Trichorhinophalangeal Syndrome Type I	
Trichorhinophalangeal Syndrome Type III	
Trigger Finger	
Trismus Pseudocamptodactyly Syndrome	
Truncus Arteriosus	
Tuberous Sclerosis	Tuberous Sclerosis
Turcot Syndrome	
Twin-Twin Transfusion Syndrome	(aka: TTTS)
Tyrosinemia, Hereditary	
Unilateral Blindness	
VACTERL Association	
Valinemia	(aka: Hypervalinemia; Valine Transaminase Deficiency)
Vascular Malformations of the Brain	(aka: Cerebral Malformations, Vascular; Intracranial Vascular Malformations; Occult Intracranial Vascular Malformations)
Vasculitis	(aka: angiitis)
Vasculitis, Cutaneous Necrotizing	(aka: CNV; Cutaneous Leukocytoclastic Angiitis; Dermal Necrotizing Angiitis; Hypersensitivity Vasculitis)
Ventricular Septal Defects	(aka: Hole in the Heart, VSD, Congenital Ventricular Defects)
Ventriculomegaly	
Vertebral anomalies	Vertebral anomalies (aka: fusion of vertebrae; hemi-vertebra; hemivertebra)
Very Long Chain Acyl CoA Dehydrogenase Deficiency	(aka: VLCAD; ACADL; LCAD)
Vocal Cord Nodules	
Vocal Cord Paresis	(aka: vocal cord paralysis), whether unilateral (left or right) or bilateral
Vogt Koyanagi Harada Syndrome	(aka: Alopecia-Poliosis-Uveitis-Vitiligo-Deafness-Cutaneous-Uveo-Oto Syndrome; Harada Syndrome; Uveomeningitis Syndrome; VKH Syndrome)

Medical Diagnosis Name	The following conditions do not meet the Type I Established Condition criteria for Montana DPHHS Part C of the IDEA Program and FES Program 6 12 19. Diagnosis Details
Von Hippel-Lindau Syndrome	(aka: Angiomatosis Retinae; Lindau Syndrome; Pheochromocytoma; Renal cell carcioma; Retinocerebellar Angiomatosis; VHL; VHL Syndrome; Von Hippel Lindau disease)
Waardenburg Syndrome	(aka: Waardenburg Syndrome Type I (WS1); Waardenburg Syndrome Type II (WS2); Waardenburg Syndrome Type IIA (WS2A); Waardenburg Syndrome Type IIB (WS2B); Waardenburg Syndrome Type III (WS3) ; Waardenburg Syndrome Type IV (WS4))
Waldmann Disease	
Weill Marchesani Syndrome	Weill Marchesani Syndrome
Weismann Netter Stuhl Syndrome	
Wernicke-Korsakoff Syndrome	Wernicke-Korsakoff Syndrome
Whipple Disease	
Wildervanck Syndrome	(aka cervicooculoacoustic syndrome)
Wilm's Tumor	
Wilson Disease	Wilson's Disease
Winchester Syndrome	
Wolff-Parkinson-White Syndrome	(aka: WPW)
Wolfram Syndrome	Wolfram Syndrome
Wyburn Mason Syndrome	
X linked Lymphoproliferative Syndrome	(aka: XLP)
Xeroderma Pigmentosum	Xeroderma Pigmentosum
Yunis Varon Syndrome	