

BabyNet Policy and Procedure Manual

Appendix 3:

Established Risk Conditions

Children with documentation that any condition on this list has been professionally diagnosed are eligible for BabyNet services based on “established risk”. Families, BabyNet Service Coordinators, and BabyNet Service Providers may link to a guide with a brief description of each condition developed by the Team for Early Childhood Solutions at http://uscm.med.sc.edu/tecs/babynet_covered_diagnoses.pdf for additional information.

| | |
|--|--|
| 10p13 Deletion | Birthweight \leq 1200 grams or \leq 28 weeks gestational age (until age 2 years) |
| 11q Deletion | Carpenter Syndrome |
| 13q Syndrome | Cataracts w/ Visual Impairment |
| 18q Deletion Syndrome | Caudal Regression Syndrome |
| 49xxxxxy syndrome (Multiple x Chromosome Syndrome) | Cerebral palsy (CP)/Static Encephalopathy |
| 1p Minus Syndrome | Charge Association/Syndrome |
| 4p Minus Syndrome | Citrullinemia |
| 6p Minus Syndrome | Cleft Hands Bilateral |
| 6q Minus Syndrome | Coffin- Lowry Syndrome |
| 7q Minus Syndrome | Cornelia de Lange |
| 8p Minus Syndrome. | Cortical Blindness |
| Agenesis of the Corpus Callosum | Cri du Chat |
| Albinism | Cystinosis |
| Amniotic Band Syndrome | Dandy Walker Malformation |
| Amyoplasia Congenita Disruptive Sequence | DiGeorge Syndrome |
| Anencephaly | Down Syndrome (Trisomy 21) |
| Angelman Syndrome | Duplication Short Arm Chromosome #20 |
| Anophthalmia | Encephalocele |
| Argininosuccinate lyase deficiency | Fazio-Londe disease |
| Argininosuccinic Aciduria | Fetal Alcohol Syndrome |
| Arthrogyriposis | Fragile X |
| Asphyxia | Glaucoma w/Visual Impairment |
| Athetoid Cerebral Palsy | Glutaric Acidemia Type 1 |
| Auditory Neuropathy | Grade IV Intraventricular Hemorrhage |
| Atresia of the External Auditory Canal | Harlequin Fetus Syndrome |
| Autism Spectrum Disorder (ASD) | Hearing Loss \geq 20 db |
| Bilateral Micromelia | Hemiparesis |
| Bilateral Optic Nerve Coloboma | Herpes Encephalitis |
| Bilateral Retinal Detachment w/Blindness | Holoprosencephaly |
| Bilateral Visual Acuity \leq 20/70 corrected vision best eye | Hydranencephaly |
| | Hydrocephaly |

Incontinentia Pigmenti Syndrome
Infantile Spasms
Isochrome 18p Syndrome
Jacobsen's Syndrome
Joubert Syndrome
Kabuki syndrome
Karsch-Neugebauer Syndrome
Klinefelter Syndrome
Krabbe Disease
Larsen syndrome
Lebers's Congenital Amaurosis
Lennox-Gastaut Syndrome
Lissencephaly Syndrome
Lowe Syndrome (oculo-cerebro-renal)
Marshal Smith Syndrome
Melnick-Frazier
Methylmalonic Acidemia
Microdactyly
Microtia
Midas Syndrome
Miller-Dieker Syndrome
Mobius sequence or Mobius Syndrome
MPS (Mucopolysaccharidosis)
MSUD (Maple Syrup Urine Disease)
Myelodysplasia
Myotonic Dystrophy
Myotubular Myopathy
Non-Ketotic Hyperglycemia
Neural Tube Defects
Opitz Syndrome
Optic Nerve Atrophy
Ornithine-Carbamyl-Transferase Deficiency
Osteogenesis Imperfecta
Pachygyria
Pallister-Killian syndrome
Pathologic Head Growth
Perinatal Asphyxia, severe
Pervasive Developmental Disorder (ASD)
Phocomelia
PKU
Pompe Disease
Prader-Willi syndrome

Propionic A acidemia
R.O.P. stage 4 & 5 Retrolental Fibroplasia
Retinitis Pigmentosa
Retinoblastoma
Rhizomelic Chondrodysplasia Punctata
Ring chromosome 9
Ring chromosome 13
Schizencephaly
Seckel Syndrome
Seizures w/Congenital Brain Malformation
Septo-Optic Dysplasia
Severe Attachment Disorder (ASD)
Shaken Baby Syndrome
Smith-Magenis Syndrome
Spastic Diplegia
Spastic Hemiplegia
Spastic Quadriplegia
Spina Bifida
Spinal Cord Injury
Spinal Muscular Atrophy
Stickler Syndrome
Syringohydromyelia
Tar syndrome
Tay- Sachs Disease
Tetrasomy 12p
Trisomy 1
Trisomy 5p
Trisomy 10
Trisomy 13
Trisomy 18
Trisomy 4
Trisomy 8 Mosaicism Syndrome
Trisomy 9
Tuberous Sclerosis
Turner's Syndrome
Vater Syndrome, with Limb Anomalies
Velo-Cardo-Facial Syndrome
Waardenberg Syndrome
Werdnig-Hoffman
William's Syndrome
Wolf-Hirschhorn Syndrome
Zellweger Spectrum Syndrome