BCUW / MADELINE
PARTNERSHIP
Special homes for very special people

IDENTIFYING INFORMATION (please print legibly)

Individual's Name: DOB:

Last 4 Digits of Social Security #:

CIRCLE APPLICABLE CODES

PRIMARY ICD-9 CODES	ICD-9 CODE	ICD-10 DIAGNOSTIC CODE	PRIMARY ICD-9 CODES	ICD-9 CODE	ICD-10 DIAGNOSTIC CODE
Abetalipoproteinemia	272.5	E78.6	Hallervorden-Spatz Syndrome	333.0	G23.0
Acrocephalosyndactyly (Apert's Syndrome)	755.55	Q87.0	Head Injury, unspecified – Age of onset:	959.01	S09.90XA
Adrenaleukodystrophy	277.86	E71.529	Hemiplegia, unspecified	342.9	G81.90
Arginase Deficiency	270.6	E72.21	Holoprosencephaly	742.2	Q04.2
Agenesis of the Corpus Callosum	742.2	Q04.3	Homocystinuria	270.4	E72.11
Agenesis of Septum Pellucidum	742.2	Q04.3	Huntington's Chorea	333.4	G10
Argyria/Pachygyria/Microgyria	742.2 or 758.33	Q04.3	Hurler's Syndrome	277.5	E76.01
Aicardi Syndrome	333	G23.8	Hyperammonemia Syndrome	270.6	E72.4
Alcohol Embryo and Fetopathy	760.71	F84.5	I-Cell Disease	272.2	E77.0
Anencephaly	655.0	Q00.0	Idiopathic Torsion Dystonia	333.6	G24.1
Angelman Syndrome	759.89	Q93.5	Incontinentia Pigmenti	757.33	Q82.3
Asperger Syndrome	299.8	F84.5	Infantile Cerebral Palsy, unspecified	343.9	G80.9
Ataxia-Telangiectasia	334.8	G11.3	Intractable Seizure Disorder	345.1	G40.309
Autistic Disorder (Childhood Autism, Infantile Psychosis, Kanner's Syndrome)	299.0	F84.0	Klinefelter's Syndrome	758.7	Q98.4
Biotinidase Deficiency	277.6	D84.1	Krabbe Disease	333.0	E75.23
Canavan Disease	330.0	E75.29	Kugelberg-Welander Disease	335.11	G12.1
Carpenter Syndrome	759.89	Q87.0	Larsen's Syndrome	755.8	Q74.8
Cerebral Palsy, unspecified	343.69	G80.9	Leigh Disease	330.8	G31.82
Cerebral Palsy, Hemiplegic, Congenital	343.1	G80.2	Lesch-Nyhan Syndrome	277.2	E79.1
Cerebral Palsy, Paraplegic, Congenital	343	G80.1	Lissencephaly	742.2	Q04.3
Cerebral Palsy, Quadriplegic	343.2	G80.0	Lowe (Terrey MacLachlan) Syndrome (Oculocerebrorenal Dystrophy)	270.8	E72.03
Charcot Marie Tooth Disease	356.1	G60.0	Marfan Syndrome	759.82	Q87.40
CHARGE Association	759.89	Q89.8	Megalencephaly	742.4	Q04.5
Cockayne Syndrome	759.89	Q89.8	Menkes Disease (X-Linked)	275.1	E83.09
Coffin-Lowry Syndrome	759.89	Q89.8	Metachromatic Leukodystrophy	330.0	E75.25
Congenital Defects of Glycosylation	279.03	D80.3	Methylmalonic Aciduria (Acidemia)	270.3 or 270.7	E71.120
Cornelia de Lange Syndrome	759.89	Q89.8	Microencephaly	742.1	Q02
Cri-du-chat Syndrome	758.31	Q93.4	Mild Intellectual Disability	317.0	F70
Crouzon Syndrome	756.0	Q75.1	Mixed Conductive and Sensorineural Hearing Loss	389.2	H90.8
DiGeorge Syndrome	279.11	D82.1	Moderate Intellectual Disability	318.0	F71
Down Syndrome	758.0	Q90.9	Moderate or Severe Impairment, Better Eye, Profound Impairment Lesser Eye	369.1	H54.10

Dubowitz Syndrome	742.8	Q07.8	Mucolipidosis Type IV	330.1	E75.11
Duchenne Muscular Dystrophy	359.1	G71.0	Mucopolysaccharidosis (Hunter's Syndrome, Hurler's Syndrome, Scheie's Syndrome)	277.5	E76.01
			Multiple Sclerosis	340	G35-37
Dystonia Musculoram Deformans	333.6	G24.1	Neuroaxonal Dystrophy	333	G23.0
Encephalopathy, not elsewhere classified	348.3	G93.40	Neurofibromatosis (von Recklinghausen's Disease)	237.71	Q85.01
Epilepsy, unspecified	345.9	G40.90	Neuronal Heterotopia	742.8	Q07.8
Fetal Alcohol Syndrome	760.71	Q86.0	Niemann-Pick Disease	272.7	E75.249
Fragile X Syndrome	759.83	Q99.2	Noonan Syndrome	759.81	Q87.1
Friedreich's Ataxia	334.0	G11.1	Other Cerebral Degeneration	331.8 or 349.89	G32.89 (nonspecified)
Fucosidosis	271.8	E77.1	Other Chromosomal Abnormalities, not elsewhere classified	758.89	Q99.8
Gaucher's Disease	272.7	E75.22	Other Disorders of Purine and Pyrimidine Metabolism (Lesch-Nyhan Syndrome)	277.2	E79.1
Generalized Convulsive Epilepsy	345.1	G40.309	Other Specified Anomalies (Cornelia de Lange Syndrome, Seckel Syndrome)	759.9	Q87.1
Generalized Non-Convulsive Epilepsy	345.0	G40.401	Other Specified Anomalies of Nervous System (Familial Dysautonomia; Riley-Day Syndrome)	742.8	G90.1
Gonadal Dysgenesis (Turner's Syndrome)	758.6	Q96.9	Other Specified Cerebral Degenerations in Childhood (Alper's Disease or Gray-Matter Degeneration; Infantile Necrotizing Encephalomyelopathy; Leigh's Disease; Subaute Necrotizing Encephalopathy or Encephamyelopathy, Rett's Syndrome)	330.8	G31.81
Grand Mal Status	345.3	G40.409	Other Specified Pervasive Developmental Disorders (Asperger's Disorder, Atypical Childhood Psychosis; Borderline Psychosis of Childhood)	299.8	F84.5
Other Spinocerebellar Diseases (Ataxia-Telangiectasia [Louis-Bar Syndrome])	334.8	G11.3	Spina Bifida without mention of Hydrocephalus	741.9	Q05.8
Paraplegia (Paralysis of Both Lower Limbs)	344.1	G82.20	Spinal Cord Injury (Initial Encounter)	952.9	S14.109A
Partial Epilepsy, with Impairment of Consciousness (Psychomotor Epilepsy)	345.4	G40.201	Spinal Muscular Atrophy, Unspecified	335.1	G12.1
Patau's Syndrome	758.1	Q91.7	Sturge-Weber Syndrome	759.6	Q85.8
Pervasive Developmental Disorder- NOS	299.9	F84.9	Symptomatic Torsion Dystonia (Athetoid Cerebral Palsy)	333.7	G80.3
Pick's Disease	331.11	G31.01	Tay-Sachs Disease	330.1	E75.02
Propionic Acidemia	270.3	E71.121	Torch Syndrome	760.02	P00.2
Prader-Willi syndrome	759.81	Q87.1	Trisomy 13	758.1	Q91.13
Profound Intellectual Disabilty	318.2	F73	Trisomy 18 (Edwards' Syndrome)	758.2	Q91.3
Pyruvate Dehydrogenase Deficiency (lactic, pyruvic)	271.8	E74.4	Tuberous Sclerosis	759.5	Q85.1
Quadriplegia and Quadriparesis	344.00	G82.5	Unspecified (Traumatic Blindness NOS)	950.9	S04.019A
Refsum's Disease	356.3	G60.1	Unspecified Anomaly of Brain, Spinal Cord, and Nervous System	742.9	Q07.9
Rett's Syndrome	330.8	F84.2	Unspecified Cause of Encephalitis	323.9	G04.90
Rubinstien-Taybi Syndrome	759.89	Q87.2	Unspecified Delay in Development (Developmental Disorder NOS)	315.9	F89
Sandhoff Disease	330.1	E75.01	Unspecified Disease of Spinal Cord	336.9	G95.9
Sanfillippo Syndrome	277.5	E76.22	Unspecified Intellectual Disability	319	F79
Schindler Disease Type 1	271.8	E77.1	Unspecified Pervasive Developmental Disorder (Pervasive Developmental Disorder NOS)	299.9	F84.9
Schizencephaly	742.4	Q04.6	Untreated Phenylketonuria	270.1	E70.0
Seckel Syndrome	759.89	Q87.1	Urea Cycle Defects	270.6	E72.20
Septo-optic Dysplasia	742.4	Q04.4	Usher Syndrome Type II	694.4	L10.4

Severe Hypoxic Ischemis CNS Injury	768.73	P91.63	Vater Association	759.89	Q87.2
Severe Intellectual Disability	318.1	F72	Werdnig-Hoffman	335.0	G12.0
Sjogren-Larsson Syndrome	757.1	Q80.9	Williams-Beauren Syndrome	758.9	Q87.8
Spastic Hemiplegia	342.1	G80.2	Wilson Disease	275.1	E83.01
Spielmeyer-Vogt Disease	330.1	E75.4	Zellwager Syndrome	277.86	E71.510
Spina Bifida	741	Q05	Psychiatric Disorder or Problem		F99

My signature on this document certifies that the diagnosis identified is based on medical evaluation and documentation and/or established medical evaluation and documentation.

Physician's Name: _____

License Number: _____

Address: _____

Telephone Number: _____

Physician's Signature/Date: _____