

Possibilities Unlimited International

Specialty Challenged Disabilities Code Sheet

<u>Code</u>	<u>Description</u>	<u>Code</u>	<u>Description</u>
A1	Aarskog Syndrome	L1	Lesch-Nyhan Syndrome
A2	Achondroplasia	L2	Leukodystrophy
A3	Agenesis of Corpus Callosum	L3	Loew's Syndrome
A4	Albinism		
A5	Apraxia		
A6	Arthritis, Juvenile	M1	Marfan Syndrome
A7	Arthrogyposis, Multiple Congenita	M2	MELAS Syndrome
A8	Asperger's Syndrome	M3	Meningitis
A9	taxia, Friedrich's	M4	Mental Retardation
A10	Ataxia, Telangiectasia	M5	Microcephaly
A11	Attention Deficit Hyperactivity Disorder	M6	Mucopolysaccharidosis
A12	Autism	M7	Multiple Sclerosis
		M8	Muscular Dystrophy
B1	Batten Disease	M9	Muscular Dystrophy, Duchenne
B2	Bell's Palsy	M10	Muscular Dystrophy, Fukuyama Type
B3	Brain Tumors, General		
B4	Branchio-Oto Renal Syndrome	N1	Neurofibromatosis
		N2	Noonan Syndrome
C1	Cerebral Palsy		
C2	Charcot-Marie-Tooth Disease	O1	Osteogenesis Imperfecta
C3	CHARGE Association		
C4	Chromosome 9, Trisomy Mosaic	P1	Penta X Syndrome
C5	Cleft Lip and Palate	P2	Phenylketonuria
C6	Cockayne Syndrome	P3	Pierre Robin Syndrome
C7	Conradi-Hunermann Syndrome	P4	Post-Polio Syndrome
C8	Cornelia de Lange Syndrome	P5	Prader-Willi Syndrome
C9	Cri du Chat Syndrome		
C10	Crouzon Disease	R1	Retinosis Pigmentosa
		R2	Rett Syndrome
D1	Dandy-Walker Malformation	R3	Rubenstein-Taybi Syndrome
D2	Deafness and Hearing Loss		
D3	Dejerine Sottas Disease	S1	Schizophrenia
D4	Diabetes, Insipidus	S2	Seizures
D5	Diabetes, Insulin-Dependent	S3	Sickle Cell Anemia
D6	Down Syndrome	S4	Smith Lemli Optiz Syndrome
D7	Dysautonomia, Familial	S5	Smith Magnesium Syndrome
D8	Dyslexia	S6	Sotos Syndrome
D9	Dystonia, Torsion	S7	Spina Bifida
		S8	Spondylolisthesis/Spondylolysis
E1	Encephalopathy, Hypoxic-Hypotensive	S9	Stroke
E2	Encephalopathy, Toxic		
E3	Epilepsy	T1	Tay-Sachs Disease
E4	Erb's Palsy	T2	Teratomas
		T3	Tetralogy of Fallot
F1	Fetal Alcohol Syndrome	T4	Thalassemia
F2	Fragile X Syndrome	T5	Thirteen Q Syndrome (13Q)
F3	Freeman Sheldon Syndrome	T6	Tourette Syndrome
		T7	Trichorhinophalangeal Syndrome
G1	Galactosemia	T8	Trisomy 13 (Patau Syndrome)
G2	Glutaricaciduria I	T9	Trisomy 18 (Edwards Syndrome)
		T10	Tuberous Sclerosis
H1	Hurler Syndrome	T11	Turner Syndrome
H2	Hydrocephalus		
H3	Hypothyroidism	W1	Werdnig Hoffman Disease
		W2	Williams Syndrome
J1	Joubert Syndrome	W3	Wiscott Aldrich Syndrome
K1	Kabuki Make-up Syndrome	Z	Other: _____
K2	Klinefelter Syndrome		
K3	Klippel Trenaunay Syndrome		