Child Health Programs Eligibility Conditions
2011-2012

NOTE: Diagnosis with an (X) are automatically eligible for the associated program.

CATEGORIES, CODES, DIAGNOSES:		DIAGNOSES BY PROGRAM				
		CMS	BCW	FIRST CARE	NEWBORN SCREENING	HEARING
001-139: INFECTIOUS AND PARASITIC DISEASES						
042	HIV					X
052.9	Varicella, congenital		X			X
079.6	Respiratory syncytial virus (RSV)			X		
090.9	Syphilis, congenital-unspecified		X			
140-239: NEOPLASMS						
190.5	Malignant neoplasm of retina (includes **Retinoblastoma**)		X			
216	Benign neoplasm of skin	X				X
237.70	Neurofibromatosis	X				
240-279: ENDOCRINE, NUTRITIONAL AND METABOLIC DISEASES, AND IMMUNITY DISORDERS						
242.0	Toxic diffuse goiter (**Hyperthyroidism**)	X				
243	Congenital hypothyroidism	X	X		X	
244	Acquired hypothyroidism	X				
250	Diabetes mellitus	X				X
252	Disorders of the parathyroid gland	X				
253	Disorders of the pituitary gland	X				
253.3	Pituitary dwarfism	X				
253.5	Diabetes insipidus	X				
255	Disorders of adrenal glands	X				
255.0	Cushing's syndrome	X				
255.2	Adrenogenital disorders (includes **Congenital adrenal hyperplasia**)	X			X	
259.0	Delay in sexual development and puberty	X				
259.1	Precocious sexual development and puberty	X				
259.8	Other specified endocrine disorders	X				
270.0	Disturbances of amino-acid transport (includes **Cystinosis**, **Hartnup disease**)		X			
270.1	Phenylketonuria (PKU), Hyperphenylalaninemia	X	X		X	
270.2	Other disturbances of aromatic amino-acid metabolism (includes **Albinism**, **Tyrosinosis**, **Tyrosinuria/Tyrosemia-types I, II, III**)	X	X		X	X (Albinism)
270.3	Disturbances of branched-chain amino-acid metabolism (includes **Maple syrup urine disease**, **Methylmalonic acidemia**, **Propionic acidemia**, **Isovaleric acidemia**, **Isobutyryl-CoA dehydrogenase**)	X	X		X	
270.4	Disturbances of sulphur-bearing amino-acid metabolism (includes **Homocystinuria**)	X	X		X	
270.6	Disorders of urea cycle metabolism (includes **Argininosuccinic aciduria**, **Citrullinemia**, **Carbamoyl phosphate synthetase I deficiency**, **Hyperargininemia**, **Ornithine transcarbamylase deficiency**)	X	X		X	
270.7	Other disturbances of straight-chain amino-acid metabolism (includes **Glutaric aciduria, Type I**)	X	X		X	
270.8	Other specified disorders of amino-acid metabolism (includes **2-Hydroxyglutaric aciduria**, **Lowe syndrome**, **Molybdenum cofactor deficiency**, **Pediatric neurotransmitter disease**, **2-Methyl-3-hydroxybutaric aciduria**, **2-Methylbutyryl-CoA dehydrogenase**)	X	X		X	
270.9	Unspecified disorder of amino-acid metabolism (includes **3-Methylglutaconic aciduria**)	X	X		X	
271.0	Glycogenosis (includes **Pompe disease-Type II**, **McArdle/myophosphorlase deficiency-Type V**)	X	X			X (Type II)
271.1	Galactosemia (includes Galactokinase defiency,	X	X		X	

Child Health Programs Eligibility Conditions
2011-2012

NOTE: Diagnosis with an (X) are automatically eligible for the associated program.

	CATEGORIES, CODES, DIAGNOSES:	DIAGNOSES BY PROGRAM: CMS	BCW	FIRST CARE	NEWBORN SCREENING	HEARING
271.8	Other specified disorders of carbohydrate transport and metabolism (includes **Carbohydrate deficiency glycoprotein syndrome-Type I**, **Fucosidosis**, **Pyruvate carboxylase deficiency**, **Pyruvate dehydrogenase deficiency**)	X	X			
271.9	Unspecified disorder of carbohydrate transport and metabolism (Includes **Schindler's Disease**)		X			
272.2	Mixed hyperlipidemia (includes **Xanthomatosis**)	X	X			
272.5	Lipoprotein deficiencies (includes **Abetalipoproteinemia/Bassen-Kornzweig syndrome**, **Hypobetalipoproteinemia**)	X	X			
272.7	Lipidoses (includeds **Infantile Gaucher disease**, **Cerebrotendinous Xanthomatosis, Niemann-Pick syndrome**)	X	X			
275.1	Disorders of copper metabolism (includes **Wilson's disease**)	X	X			
275.3	Disorders of phosphorus metabolism (includes **Vitamin D-resistant rickets**)	X				
275.8	Other specified disorders of mineral metabolism (includes **Menkes kinky hair syndrome xq12-q13**)	X	X			
276.2	Acidosis (includes **3-Hydroxy-3-Methylglutaryl-CoA Lyase deficiency**, **3-Methylcrotonyl-CoA Carboxylase deficiency**, **B-Ketothiolase deficiency**, **Multiple-CoA-Carboxylase deficiency**)	X			X	
277	Other and unspecifieid disorders of metabolism (includes **Fumerase deficiency**)		X			
277.0	Cystic Fibrosis	X			X	X
277.2	Other disorders of purine and pyrimidine metabolism (includes **Lesch-Nyhan disease**)	X	X			
277.5	Mucopolysacharidosis I (**Hunter Syndrome**), II (**Hurler Syndrome**), III (**Sanfilippo Syndrome**)	X	X			
277.6	Other deficiencies of circulating enzymes (includes **Biotinidase deficiency**)	X	X		X	
277.81	Primary carnitine deficiency	X	X			
277.82	Carnitine deficiency due to inborn errors of metabolism	X			X	
277.84	Other secondary carnitine deficiency	X			X	
277.85	Disorders of fatty acid oxidation (includes the following deficiencies listed below*: **LCHAD, CUD, CPT1/CPT2, MAD, LCAD/VLCAD, MCADD, SCAD, TFP**)	X			X	
	*3-Hydroxy Long Chain Acyl-CoA Dehydrogenase (**LCHAD**) Deficiency	X			X	
	*Carnitine Uptake Defect (**CUD**)	X				
	*Carnitine palmitoyltranserase deficiencies (**CPT1, CPT2**)	X	X		X	
	*Glutaric Aciduria type II, Multiple Acyl-CoA Dehydrogenase (**MAD**)	X	X		X	
	*Long Chain/Very Long Chain Acyl-CoA-Dehydrogenase Deficiency (**LCAD, VLCAD**)	X	**X (VLCAD only)**		X	
	*Medium-Chain Acyl CoA Dehydrogenase Deficiency (**MCADD**)	X	X		X	
	*Short chain acyl CoA dehydrogenase deficency (**SCAD**)	X			X	
	*Trifunctioinal Protein (**TFP**) Deficiency	X			X	
277.86	Peroxisomal disorders (includes **Adrenoleukodystrophy**, **Mevalonate kinase deficiency**, **Zellweger syndrome**, **Rhizomelic chrondrodysplasia punctata**)	X	X			**X (Zellweger)**
277.87	Disorders of mitochondrial metabolism (includes **deficiencies, syndromes and disorders listed below****)	X	X			X
	****2-Ketoglutarate Dehydrogenase Deficiency**	X	X			
	****Complex I, II, III, IV, V deficiencies**	X	X			
	****Dihydrolipoyl Dehydrogenase deficiency**	X	X			
	****Kearns-Sayre syndrome**	X	X			X

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
	****Oxidative Phosphorylation Disorders**	X	X			
	****MELAS: Mitochondrial encephalomyopathy/lactic acidosis & strokelike episodes**	X	X			
	****MERRF: Myoclonic epilepsy with ragged red fibers**	X	X			X
	****NARP: Neurogenic weakness, ataxia, retinitis pigmentose**	X	X			
277.9	Unspecified disorder of metabolism (includes **Enzymopathy** NOS)	X				
279.11	DiGeorge's syndrome	X	X		**to be added**	X
280-289: DISEASES OF THE BLOOD AND BLOOD-FORMING ORGANS						
282	Herediatary hemolytic anemias	X				
282.3	Other hemolytic anemias due to enzyme deficiency (includes **Triosephosphate Isomerase Deficiency/TPI**)	X	X			
282.4	Thalessemia	X			X	
282.6	Sickle-cell disease	X			X	X
282.7	Other hemoglobinopathies	X			X	
286	Coagulation defects	X				
290-319: MENTAL DISORDERS						
299.00	Autistic disorder current or active state		X			
299.10	Childhood disintegrative disorder current or active state		X			
299.80	Other specified pervasive developmental disorders current or active state (includes **Asperger's disorder**)		X			
299.90	Unspecified pervasive developmental disorder current or active state (Includes **Pervasive Developmental Disorder NOS/PDD-NOS**)		X			
320-389: DISEASES OF THE NERVOUS SYSTEM AND SENSE ORGANS						
320	Bacterial Meningitis (**for BCW: with Neurological sequelae/Severe neurological insult)**		X			X
321	Meningitis due to other organisms					X
323.9	Unspecified cause of encephalitis					X
330.0	Leukodystrophy (Includes **Krabbe's disease**, **Pelizaeus-Merzbacher disease**, **Canavan disease**)	X	X			
330.1	Cerebral lipidoses (includes S**andhoff disease**, **Batten disease**, **Tay-Sachs**, **Gangliosidosis/GM1, GM2**)	X	X			X
330.3	Cerebral degeneration of childhood in other diseases classified elsewhere		X			
330.8	Other specified cerebral degenerations in childhood (includes **Leigh's disease**, **Rett's syndrome**)	X	X			X
330.9	Unspecified cerebral degeneration in childhood		X			
331.3	Communicating hydrocephalus	X	X			X
331.4	Obstructive hydrocephalus	X	X			X
331.9	Cerebral degeneration unspecified (includes **Cortical atrophy**)	X	X			X
333.0	Other extrapyramidal disease and abnormal movement disorders (includes **Hallervorden-Spatz syndrome**, **Huntington's Chorea**)	X	**X (Hallevorden-Spatz)**			
335.0	Werdnig-Hoffmann Disease (includes **Infantile Spinal Muscular Atrophy**)	X	X			
335.11	Kugelberg-Welander disease	X	X			
342.00	Flaccid hemiplegia and hemiparesis affecting unspecified side		X			
342.01	Flaccid hemiplegia and hemiparesis affecting dominant side		X			
342.02	Flaccid hemiplegia and hemiparesis affecting nondominant side		X			
342.1	Spastic hemiplegia and hemiparesis affecting unspecified side		X			
342.11	Spastic hemiplegia and hemiparesis affecting dominant side		X			

Child Health Programs Eligibility Conditions
2011-2012

NOTE: Diagnosis with an (X) are automatically eligible for the associated program.

		DIAGNOSES BY PROGRAM				
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
342.12	Spastic hemiplegia and hemiparesis affecting nondominant side		X			
342.80	Other specified hemiplegia and hemiparesis affecting unspecified side		X			
342.81	Other specified hemiplegia and hemiparesis affecting dominant side		X			
342.82	Other specified hemiplegia and hemiparesis affecting nondominant side		X			
342.90	Unspecified hemiplegia and hemiparesis affecting unspecified side		X			
342.91	Unspecified hemiplegia and hemiparesis affecting dominant side		X			
342.92	Unspecified hemiplegia and hemiparesis affecting nondominant side		X			
343.0	Congenital diplegia (includes **Congenital paraplegia**)	X	X			
343.1	Congenital hemiplegia	X	X			
343.2	Congential quadriplegia	X	X			
343.3	Congenital monoplegia	X	X			
343.4	Infantile hemiplegia	X	X			
343.8	Other specified infantile cerebral palsy	X	X			X
343.9	Infantile cerebral palsy unspecified	X	X			X
344.00	Quadriplegia unspecified	X				
344.01	Quadriplegia c1-c4 complete		X			
344.02	Quadriplegia c1-c4 incomplete		X			
344.03	Quadriplegia c5-c7 complete		X			
344.04	Quadriplegia c5-c7 incomplete		X			
344.09	Other quadriplegia		X			
344.1	Paraplegia	X	X			
344.2	Diplegia of upper limbs	X	X			
344.30	Monoplegia of lower limb affecting unspecified side	X	X			
344.31	Monoplegia of lower limb affecting dominant side	X	X			
344.32	Monoplegia of lower limp affecting nondominant side	X	X			
344.40	Monoplegia of upper limb affecting unspecified side	X	X			
344.41	Monoplegia of upper limb affecting dominant side	X	X			
344.42	Monoplegia of upper limp affecting nondominant side	X	X			
344.5	Unspecified monoplegia		X			
344.6	Cauda equina syndrome	X				
344.61	Cauda equina with neurogenic bladder	X				
344.9	Paralysis unspecified		X			
345	Epilepsy	X	X			X
347.00	Cataplexy and Narcolepsy (narcolepsy with/without cataplexy)	X				
348.0	Cerebral cysts	X				
348.1	Anoxic brain dmage		X			
348.3	Encephalopathy NEC (**for BCW: with Neurological sequelae**)	X	X			
348.31	Metabolic encephalopathy	X	X			
348.39	Other encephalopathy	X	X			
348.81	Temporal sclerosis	X	X			
348.89	Other conditions of brain		X			
348.9	Unspecified condition of brain		X			
349.9	Unspecified disorders of nervous system		X			
356.0	Heriditary peripheral neuropathy (includes **Dejerine-Sottas disease**)	X	X			
356.1	Peroneal muscular atrophy (includes **Charcot-Marie-Tooth disease**)	X	X			X
356.3	Refsum's disease	X	X			X
358.9	Myoneural disorders unspecified (includes **Neuromyopathy** NEC)	X	X			
359.0	Congenital hereditary muscular dystrophy (**includes diseases/disorders below*****)	X	X			X
	***Central Core Disease**	X	X			
	***Congenital muscular dystrophy (Fukuyama)**	X	X			X
	***Infantile Neuroaxonal dystrophy (Seitelberger's disease II)**	X	X			

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
	***Myotubular Myopathy (MTM)	X	X			
	***Nemaline Myopathy (NM), Nemaline Rod Myopathy	X	X			
359.1	Hereditary progressive muscular dystrophy (includes **Duchenne MD**, **Emery-Dreifuss MD**. **Facioscapulohumeral MD**, **Limb-girdle MD**)	X				
359.21	Myotonic muscular dystrophy (includes **Steinert's disease**)	X	X			X
359.22	Myotonia Congenital (includes **Thomsen's disease**)	X	X			
361.0	Retinal detachment with retinal defect	X				
362.0	Diabetic retinopathy	X				
362.12	Exudative retinopathy (includes **Coats' Syndrome/Disease**) retinopathy	X	X			
362.26	Retinopathy of Prematurity - Grade 4	X	X			
362.27	Retinopathy of Prematurity - Grade 5	X	X			
362.76	Dystrophies primarily involving the retinal pigment epithelium (includes **Leber's Congenital Amaurosis**)	X	X			
365	Glaucoma	X				
366	Cataracts	X				
368.00	Amblyopia, unspecified	X				
369.00	Blindness of both eyes impairment level not further specified	X	X			
369.01	Better eye total vision impairment; lesser eye not further specified	X	X			
369.02	Better eye near total vision impairment; lesser eye not further specified	X	X			
369.03	Better eye near total vision impairment; lesser eye total vision impairment	X	X			
369.04	Better eye near total vision impairment; lesser eye near total vision impairment	X	X			
369.05	Better eye profound vision impairment; lesser eye not further specified	X	X			
369.06	Better eye profound vision impairment; lesser eye total vision impairment	X	X			
369.07	Better eye profound vision impairment; lesser eye near total vision impairment	X	X			
369.08	Better eye profound vision impairment; lesser eye profound vision impairment	X	X			
369.11	Better eye severe vision impairment; lesser eye blind not further specified	X	X			
369.12	Better eye severe vision impairment; lesser eye total vision impairment	X	X			
369.13	Better eye severe vision impairment; lesser eye; near total vision impairment	X	X			
369.14	Better eye sever vision impairment; lesser eye profound vision impairment	X	X			
369.2	Moderate or Severe vision Impairment, both eyes		X			
369.3	Unqualified visual loss both eyes		X			
369.6	Profound vision impairment, one eye	X				
374.30	Ptosis of the eyelid unspecified (interfering with vision)	X				
377.11	Primary optic atrophy	X	X			
377.75	Cortical blindness	X	X			
378	Strabismus	X				
379.31	Aphakia (absence of lens)	X				
385.3	Cholesteatoma of middle ear and mastoid	X				X
389	Hearing loss (defined as a threshold of 20 decibels or more in the in speech range frequencies (500-4000) in the better ear)	X				X
389.0	Conductive hearing loss	X				X
389.06	Conductive hearing loss, bilateral	X	X			
389.1	Sensorineural hearing loss	X				X
389.11	Sensory hearing loss (bilateral)	X	X			X
389.12	Neural hearing loss (bilateral)	X	X			X
389.18	Sensorineural hearing loss, bilateral	X	X			X
389.2	Mixed conductive and sensorineural hearing loss	X				X
389.22	Mixed hearing loss, bilateral	X	X			
389.8	Other specified forms of hearing loss (mild-moderate, bilateral)		X			X
389.9	Unspecified hearing loss (Deafness) (bilateral)	X	X			X

Child Health Programs Eligibility Conditions
2011-2012

NOTE: Diagnosis with an (X) are automatically eligible for the associated program.

		DIAGNOSES BY PROGRAM				
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
390-459: DISEASES OF THE CIRCULATORY SYSTEM						
393	Chronic rheumatic pericarditis	X				
394	Disease of mitral valve	X				
395	Disease of aortic valve	X				
396	Disease of mitral and aortic valves	X				
397.0	Disease of tricuspid valve	X				
397.1	Rheumatic disease of pulmonary valve	X				
397.9	Rheumatic disease of endocardium valve, unspecified	X				
398.0	Rheumatic myocarditis	X				
398.91	Rheumatic heart failure (congestive)	X				
414.10	Aneurysm of heart wall	X				
414.11	Aneurysm of coronary vessel	X				
415.0	Acute cor pulmonale	X				
416.0	Primary pulmonary hypertension	X				X
416.1	Kyphoscoliotic heart disease	X				
422	Acute myocarditis	X				
424.0	Mitral valve disorders	X				
424.1	Aortic valve disorders	X				
424.2	Tricuspid valve disorders	X				
424.3	Pulmonary valve disorders	X				
424.9	Endocarditis, valve unspecified	X				
425	Cardiomyopathy	X				
426	Conduction disorders	X				
427	Cardiac dysrhythmia	X				
428	Heart failure	X				
429.0	Myocarditis, unspecified	X				
429.3	Cardiomegaly	X				
446.1	Acute febrile mucocutaneous lymph node syndrome (includes **Kawasaki disease**)	X				X
460-519: DISEASES OF THE RESPIRATORY SYSTEM						
478.4	Polyp of vocal cord or larynx	X				
478.74	Stenosis of larynx	X				
493	Asthma	X				
496	Chronic airway obstruction not elsewhere classified	X				
520-579: DISEASES OF THE DIGESTIVE SYSTEM						
520	Anodontia	X				
524	Dentofacial anomalies, malocclusion (not to include routine dental care or cosmetic orthodonture)	X				
530.13	Eosinophilic Esophagitis	X				
555.9	Regional enteritis of unspecified site (includes **Crohn's Disease**)	X				
580-629: DISEASES OF THE GENITOURINARY SYSTEM						
593.7	Vesicoureteral reflux	X				
630-679: COMPLICATIONS OF PREGNANCY, CHILDBIRTH, AND THE PUERPERIUM						
	No listings					
680-709: DISEASES OF THE SKIN AND SUBCUTANEOUS TISSUE						
701.4	Keloid scar (only if painful, not for cosmetic purposes)	X				

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
709.2	Scar conditions and fibrosis of skin- Follow-up	X				
710-739: DISEASES OF THE MUSCULOSKELETAL SYSTEM AND CONNECTIVE TISSUE						
710.0	Systemic lupus erythematosus	X				
714.0	Rheumatoid arthritis	X				
714.3	Juvenile chronic polyarthritis	X				
727.81	Contracture of tendons	X				
728.3	Other specific muscle disorders (includes **Arthrogryposis**)	X				
730.1	Osteomyelitis - chronic	X				
732.1	Juvenile osteochondrosis of hip and pelvis (includes **Legg-Calve-Perthes**)	X				
732.2	Nontraumatic slipped upper femoral epiphysis	X				
732.4	Juvenile osteochondrosis of lower extremity, excluding lower foot (includes **Osgood-Schlatter's disease**)	X				
733.4	Aseptic necrosis of bone	X				
733.41	Aseptic necrosis of head of humerus	X				
733.42	Aseptic necrosis of head and neck of femur	X				
733.43	Aseptic necrosis of medial femoral condyle	X				
733.8	Malunion and nonunion of fracture	X				
736.2	Other acquired deformities of finger	X				
736.3	Acquired deformities of hip	X				
736.31	Coxa valga (acquired)	X				
736.32	Coxa vara (acquired)	X				
736.4	Genu valgum or varum (acquired)	X				
736.5	Genu recurvatum (acquired)	X				
736.6	Other acquired deformities of knee	X				
736.7	Other acquired deformities of ankle and foot	X				
736.8	Acquired deformities of other parts of limbs	X				
736.81	Unequal leg length (acquired)	X				
736.9	Acquired deformity of limb, site unspecified	X				
737	Curvature of spine	X				
737.1	Kyphosis (acquired)	X				
737.2	Lordosis (acquired)	X				
737.30	Scoliosis (and kyphoscoliosis) idioipathic	X				
737.4	Curvature of spine associated with other conditions	X				
740-759: CONGENITAL ANOMALIES						
740.0	Anencephalus	X	X			
740.1	Craniorachischisis	X	X			
740.2	Iniencephaly	X	X			
741	Spina Bifida	X	X	X		X
741.0	Spina Bifida with hydrocephalus (includes Arnold-Chiari Syndrome, Chiari Malformation)	X	X	X		X
741.01	Spina bifida cervical region, with hydrocephalus	X	X	X		
741.02	Spina bifida dorsal (thoracic) region with hydrocephalus	X	X	X		
741.03	Spina bifida lumbar region with hydrocephalus	X	X	X		
741.91	Spina bifida cervical region without hydrocephalus	X	X	X		
741.92	Spina bifida dorsal (thoracic) region without hydrocephalus	X	X	X		
741.93	Spina bifida lumbar region without hydrocephalus	X	X	X		
741.9	Spina Bifida without mention of hydrocephalus (includes **Myelomeningocele without hydrocephalus**)	X	X	X		X

Child Health Programs Eligibility Conditions
2011-2012

NOTE: Diagnosis with an (X) are automatically eligible for the associated program.

	CATEGORIES, CODES, DIAGNOSES:	DIAGNOSES BY PROGRAM				
		CMS	BCW	FIRST CARE	NEWBORN SCREENING	HEARING
742.0	Encephalocele	X	X	X		
742.1	Microcephalus	X	X	X		X
742.2	Congenital reduction deformities of brain (includes **Agenesis of corpus callosum**, **Holoprosencephaly**, **Lissencephaly**, **Agyria**, **Microgyria**, **Polymicrogyria**, **Septo-optic dysplasia, Aicardi syndrome**)	X	X			X
742.3	Congenital hydrocephalus (includes **Dandy Walker Syndrome, Walker-Warburg Syndrome**, **Hydranencephaly**)	X	X	X		X
742.4	Other specified congenital anomalies of brain (includes **Macroencephaly**, ****Megalencephaly**, ****Porencephaly**, ****Schizencephaly**)	X	**X (only those with **)**			
742.51	Diastematomyelia	X	X			
742.53	Hydromyelia	X	X			
742.59	Other specified congenital anomalies of spinal cord	X	X			
742.8	Other specified congenital anomalies of nervous system (includes **Riley-Day syndrome**, **Familial dysautonomia**)	X	X			
742.9	Unspecified congenital anomaly of brain spinal cord and nervous system (includes **Cebocephaly**, **Colpocephaly**, **Ethmocephaly**, **Isotretinoin or Accutane Embryopathy/IE**, **Schinzel-Giedion syndrome/SGS**)	X	X			
743	Congenital anomalies of eye	X				
743.0	Anophthalmos	X	X			
743.1	Microphthalmos (bilateral)	X	X			
743.57	Specified congenital anomalies of optic disc (includes **Coloboma of optic disc [congenital], bilateral**)	X	X			
743.62	Congenital deformities of eyelids (includes **Ablepharon Macrostomia syndrome**)	X	X			
744.0	Congenital anomalies of ear causing impairment of hearing	X				X
745.0	Common truncus	X				X
745.1	Transposition of great vessels	X		X		X
745.2	Tetralogy of Fallot	X		X		X
745.3	Common ventricle	X				X
745.4	Ventricular septal defect	X				X
745.5	Atrial septal defect - ostium secundum type	X				X
745.6	Endocardial cushion defects	X				X
745.69	Other endocardial cushion defects (includes **Atrioventricular canal**)	X				X
746	Other congenital anomalies of heart	X				X
746.0	Anomalies of pulmonary valve congenital	X				X
746.1	Tricuspid atresia and stenosis congenital	X	X			X
746.2	Ebstein's anomaly	X	X			X
746.3	Congenital stenosis of aortic valve	X				X
746.5	Congenital mitral stenosis	X				X
746.6	Congenital mitral insufficiency	X				X
746.7	Hypoplastic left heart syndrome	X	X	X		X
746.86	Congenital heart block	X				X
746.87	Malposition of heart and cardiac apex	X				X
746.9	Unspecified congenital anomaly of heart (**for **BCW only**: **Congenital Heart Disease co-occurring with either 759.0 [Polysplenia or Asplenia] or 759.3 [Situs inversus]**)	X	**X****			X
747.0	Patent ductus arteriosus	X				X
747.1	Coarctation of aorta	X		X		X
747.3	Congenital anomalies of pulmonary artery	X	X			X

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
747.4	Congenital anomalies of great veins	X				X
747.41	Total anomalous pulmonary venous connection	X	X	X		X
748.0	Choanal atresia	X				
748.3	Other congenital anomalies of larynx, trachea, bronchus (includes **Tracheomalacia, congenital**)	X				
748.61	Congenital bronchiectasis	X				
749	Cleft palate and cleft lip	X		X		X
749.10	Cleft lip, unspecified	X		X		X
750.0	Tongue tie	X				
750.3	Congenital tracheoesophageal fistula, esophageal atresia and stenosis	X		X		
751.0	Meckel's diverticulum	X				
751.1	Congenital atresia and stenosis of small intestine	X				
751.2	Congenital atresia and stenosis of large intestine rectum and anal canal (includes **Bowel atresia, Imperforate anus**)	X				
751.3	Hirschsprungs disease and other congenital function disorders of colon	X				
751.5	Other congenital anomalies of intestine (includes **Duplication of rectum**)	X				
752	Congenital anomalies of genital organs					
752.5	Undescended and retractile testicle(s)	X				
752.6	Hypospadias and epispadias and other penile anomalies	X				
752.7	Indeterminate sex and pseudohermaphroditism	X				
753	Congenital anomalies of urinary system					
753.1	Cystic kidney disease	X				
753.20	Congenital obstructive defect of renal pelvis and ureter	X				
753.3	Other specified anomalies of kidney (includes **Horseshoe kidney**)	X				
753.4	Other specified anoamlies of ureter (includes **Duplication of ureters**)	X				
753.5	Exstrophy of urinary bladder	X				
753.6	Congenital atresia and stenosis of urethra and bladder neck	X				
753.8	Other specified anomalies of bladder and urethra	X				
754.0	Congenital musculoskeletal deformities of skull, face, and jaw	X				
754.1	Congenital musculoskeletal deformities of sternocleidomastoid muscle (includes Torticollis)	X				
754.2	Congenital musculoskeletal deformities of spine	X				
754.3	Congenital dislocation of hip	X				
754.50	Congenital talipes varus	X				
754.51	Congenital talipes equinovarus (includes Club foot)	X				
754.6	Congenital valgus deformities of feet	X				
754.81	Pectus excavatum (funnel chest)	X				
754.82	Pectus carinatum (pigeon chest)	X				
754.89	Other specified nonteratogenic anomalies (includes **Arthrogryposis Muliplex Congenita/AMC, Club hand**)	X	X			
755.0	Polydactyly	X				
755.1	Syndactyly	X				
755.2	Reduction deformities of upper limb congenital	X				
755.3	Congenital reduction deformities of lower limb	X				
755.55	Acrocephalosyndactyly, Type I (**Apert Syndrome, Pfeiffer's Syndrome**)	X	X			X
755.61	Coxa valga, congenital	X				
755.62	Coxa vara, congenital	X				
755.64	Congenital deformity of knee (joint)	X				

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
756.0	Congenital anomalies of skull and face bones (includes **Crouzon's**, **Pierre-Robin**, **Treacher Collins**, **Craniofacial dysplasia**, **Craniostenosis**, **Craniosynostosis**, **Shprintzen-Goldberg Craniosynostosis Syndrome**)	X				X
756.1	Congenital anomalies of spine	X				
756.12	Spondylolisthesis	X				
756.14	Hemivertebra	X				
756.16	Klippel-Feil Syndrome	X	X			X
756.3	Other conenital anomalies of ribs and sternum	X				
756.4	Chondrodystrophy (includes **Achondroplasia, Chondrodysplasia**)	X				X
756.5	Congenital osteodystrophies (includes **Bannayan-Riley-Ruvalcaba Syndrome**)	X	X			
756.51	Osteogenesis imperfecta	X		X		X
756.6	Congenital anomalies of diaphragm (includes **Congenital diaphragmatic hernia**)	X				
756.73	Gastroschisis	X				
756.83	Ehlers Danlos Syndrome	X				
756.9	Other and unspecified congenital anomalies of musculoskeletal system (includes **Camptomelic dysplasia**)	X	X			
757.1	Ichthyosis congenita	X				
757.33	Incontinentia pigmenti (also known as **Hypomelanosis of Ito**)	X	X			
757.39	Other specified congenital anomalies of skin	X				
758	Chromosomal anomalies	X				X
758.0	Trisomy 21 (**Down Syndrome**)	X	X	X		X
758.0	Trisomy 22/**Cat Eye Syndrome** (**Coloboma of Iris-Anal Atresia Syndrome**)	X	X			
758.1	Trisomy 13 (**Patau's Syndrome**)	X	X			X
758.2	Trisomy 16, Trisomy 18 (**Edwards' Syndrome**)	X	X			X
758.3	Trisomy 11 mosaicism	X	X			
758.31	Cri-du-Chat Syndrome (**Deletion 5p syndrome**)	X	X			X
758.32	Velo-Cardio-Facial Syndrome; Shprintzen Syndrome (**Deletion 22q11 syndrome**)	X	X			X
758.33	Other Autosomal deletion syndromes del. 3p, 4p, 4q, 7, 9p, 9q, 11q, 13q, 18p, 18q, 22 (includes **Miller-Deiker syndrome**, **Smith-Magenis syndrome**)	X	X			X
758.39	Other autosomal deletions (Includes **Aniridia-Wilms Tumor Association/WAGR syndrome - del. 11p13**)	X	X			
758.4	Balanced autosomal translocation in normal individual	X				
758.4	Trisomy 16 mosaicism	X	X			
758.5	Other conditions due to autosomal anomalies (includes **Tetrasomy 12p/Pallister-Killain syndrome**, **Trisomy 8**, **Trisomy 9,10,15, & 19 mosaicism**)	X	X			X
758.6	Gonadal dysgenesis (includes **XO syndrome**, **Turner syndrome**)	X	X			X
758.7	Klinefelter's Syndrome	X	X			X
758.8	Triploidy and diploid/triploid mixooploid syn (69,xxy; 46,xx/69xxy)	X	X			
758.8	Other chromosmal anomalies dup. 3q, 4p,9q, 10q, 15q	X	X			
758.81	Other conditions due to sex chromosome anomalies (includes **XYY, XXX, XXXX XXXY, XXXXX XXXXY syndromes**)	X	X			
758.89	Other conditions due to chromosome anomalies	X	X			
758.9	Trisomy 7 mosaicism	X				
759.5	Tuberous Sclerosis (TS)	X	X			
759.6	Other congenital hamartoses not elsewhere classified (includes **Sturge-Weber Syndrome**)	X	X			

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
759.7	Multiple congenital anomalies so described	X				X
759.81	Prader-Willi Syndrome	X	X			
759.82	Marfan Syndrome	X	X			
759.83	Fragile X Syndrome	X	X			X
759.89	Other specified congenital anomalies (includes the following syndromes: **Angelman**, **Brachman-de Lange**, **CHARGE Association**, C**ockayne's**, **Cornelia de Lange's**, **Joubert**, **Mohr's -types I and II**, **Noonan's**, **Rubinstein-Taybi's**, **Smith-Lemli-Opitz, Williams**)	X	X			X
759.89	Congenital malformation syndrome affecting multiple systems, not elsewhere classified - **Beckwith, Vater, Goldenhar, Knist**, etc.	X				X
760-779: CERTAIN CONDITIONS ORIGINATING IN THE PERINATAL PERIOD						
760.71	Alcohol affecting fetus or newborn via placenta or breast milk (includes **Fetal Alcohol Syndrome**)		X			X
760.74	Anti-infectives affecting fetus or newborn via placenta or breast milk (includes **Methylmercury poisoning, severe**)		X			X
760.77	Anticonvulsants affecting fetus or newborn via placenta or breastmilk (includes **Phenytoin/Dilantin**, **Valproic acid, Phenobarital**, **Carbamazepine**)		X			
760.78	Antimetabolic agents affecting fetus or newborn via placenta or breast milk (includes **Aminopterin**)		X			
760.79	Other noxious influences affecting fetus or newborn via placenta or breast milk (includes **Accutane**, **Dicumarol**, **Thalidomide**)		X			
765.00	Disorders relating to extreme immaturity of infant unspecified weight			X		
765.01	Disorders relating to extreme immaturity of infant <500 gms			X		
765.02	Disorders relating to extreme immaturity of infant 500-749 gms			X		
765.03	Disorders relating to extreme immaturity of infant 750-999 gms			X		
765.04	Disorders relating to extreme immaturity of infant 1000-1249 gms			X		
765.05	Disorders relating to extreme immaturity of infant 1250-1499 gms			X		
765.06	Disorders relating to extreme immaturity of infant 1500-1749 gms			X		
765.07	Disorders relating to extreme immaturity of infant 1750-1999 gms			X		
765.08	Disorders relating to extreme immaturity of infant 2000-2499 gms			X		
765.10	Disorders relating to other preterm infants unspecified weight			X		
765.11	Disorders relating to other preterm infants <500 gms			X		
765.12	Disorders relating to other preterm infants 500-749 gms			X		
765.13	Disorders relating to other preterm infants 750-999 gms			X		
765.14	Disorders relating to other preterm infants 1000-1249 gms			X		
765.15	Disorders relating to other preterm infants 1250-1499 gms			X		
765.16	Disorders relating to other preterm infants 1500-1749 gms			X		
765.17	Disorders relating to other preterm infants 1750-1999 gms			X		
765.18	Disorders relating to other preterm infants 2000-2499 gms			X		
767.0	Subdural and cerebral hemorrhage due to birth trauma (Injury during perinatal period)			X		
768.5	Severe birth asphyxia (with neurlogic involvement - APGAR < or equal to 3 at 5 minutes)	X		X		
768.70	Hypoxic-ischemic encephalopathy (HIE) unspecified	X	X	X		
769	Respiratory distress syndrome in newborn			X		

Child Health Programs Eligibility Conditions
2011-2012

		DIAGNOSES BY PROGRAM				
NOTE: Diagnosis with an (X) are automatically eligible for the associated program.						
CATEGORIES, CODES, DIAGNOSES:		**CMS**	**BCW**	**FIRST CARE**	**NEWBORN SCREENING**	**HEARING**
770.7	Chronic respiratory disease arising in the perinatal period (includes **Bronchopulmonary dysplasia**)	X		X		
770.9	Unpecified respiratory condition of fetus and newborn (significant respiratory distress - vent. > 48 hrs.)			X		
771.0	Congenital rubella		X			X
771.1	Congenital cytomegalovirus (CMV) infection		X			X
771.2	Other congenital infections specific to the perinatal period (includes **Herpes Simplex - congenital**, **Toxoplasmosis**)		X	X		X
772.13	Intraventricular Hemorrhage (IVH) Grade III		X	X		X
772.14	Intraventricular Hemorrhage (IVH) Grade IV		X	X		X
774.6	Unspecified fetal and neonatal jaundice (Hyperbilirubinemia requiring exchange transfusion)					
775.2	Neonatal myasthenia gravis		X			
777.53	Stage III necrotizing enterocolitis (NEC) in newborn			X		
779.0	Convulsions in newborn (seizures)			X		
779.7	Periventricular (or Preventricular) Leukomalacia (PVL)		X	X		X
780-799: SYMPTOMS, SIGNS, AND ILL-DEFINED CONDITIONS						
783.41	Failure to Thrive			X		
796.4	Other abnormal clinical findings **FOR DIAGNOSTIC PERIOD ONLY - NOT TO EXCEED 6 (SIX) MONTHS**	X				
800-999: INJURY AND POISONING						
887	Traumatic amputation of arm and hand (partial)(complete)	X				
896	Traumatic amputation of foot (complete)(partial)	X				
897	Traumatic amputation of leg(s) (partial)(complete)	X				
959.01	Other and unspecified injury to head NEC		X			X
984	Toxic effect of lead and its compounds (including fumes)	X				X
995.55	Shaken baby syndrome					X
996.83	Complicatons of transplanted heart		X			X