



CMS CLINICAL ELIGIBILITY ATTESTATION

Patient Name: _____ DOB: _____ Medicaid and/or KidCare ID: _____

Parent/Legal Guardian Name: _____ Phone number: _____

Initial all that Apply:

Initials	ICD 10	Descriptor
		<u>B Codes- Viral Infections with Skin and Mucous Membrane Lesions Certain infectious and parasitic diseases</u>
	B20	Human immunodeficiency virus (HIV) disease
		<u>C and D Codes- Neoplasmstic Lesions</u>
	C00-D48	All malignancies (except those in remission greater than five years), (list specific ICD10 code and descriptor) _____
		<u>D Codes ≥ 50- Diseases of the blood and blood-forming organs-and certain disorders involving the immune mechanism</u>
	D57.00- D57.1	Hb-SS disease, (list specific ICD10 code and descriptor) _____
	D66	Hereditary factor VIII deficiency
	D81.0- D81.4	Severe combined immunodeficiency [SCID], (list specific ICD10 code and descriptor) _____
	D82.1	Digeorge’s syndrome
		<u>E Codes- Endocrine, nutritional and metabolic disorders</u>
	<u>E03.1</u>	<u>Congenital Hypothyroidism</u>
	<u>E10.10-E10.9</u>	<u>Type 1 diabetes mellitus,</u> (list specific ICD10 code and descriptor) _____
	<u>E11.00- E11.9</u>	<u>Type 2 diabetes mellitus,</u> (list specific ICD10 code and descriptor) _____
	<u>E25.0</u>	<u>Congenital adrenal hyperplasia</u>
	E70.0	Classical phenylketonuria
	E70.21	Tyrosinemia
	E71.0	Maple Syrup Urine Disease
	E71.110	Isovaleric acidemia
	E71.111	3-methylglutaconic aciduria
	E71.120	Methylmalonic acidemia
	E71.121	Propionic acidemia
	<u>E71.310</u>	<u>LCAD/LCHAD/VLCAD Fatty acid oxidation disorders and ketogenesis disorders</u>
	<u>E71.311</u>	<u>MCAD Fatty acid oxidation disorders and ketogenesis disorders</u>
	<u>E71.312</u>	<u>SCAD Fatty acid oxidation disorders and ketogenesis disorders</u>
	<u>E71.313</u>	<u>Organic Acidurias (glutaric aciduria, Canavan)</u>
	E72.11	H homocystinuria
	<u>E72.21</u>	<u>Argininemia Urea Cycle Disorder</u>
	E72.22	Arginosuccinic aciduria
	E72.23	Citrulinemia

<u>E72.4</u>	<u>Ornithine Transcarbamylase Deficiency Urea Cycle Disorder</u>
<u>E72.51</u>	<u>Non-ketotic Hyperglycinemia</u>
<u>E74.09</u>	<u>Glycogen storage disease</u>
E74.21	Galactosemia
E75.23	Krabbe disease
E75.240- E75.249	Niemann-Pick disease, (list specific ICD10 code and descriptor) _____
E75.25	Metachromatic leukodystrophy
E76.01	Hurler's syndrome
E76.1	Mucopolysaccharidosis, type II
E76.22	Sanfilippo mucopolysaccharidoses
E84.0	Cystic fibrosis, (list specific ICD10 code and descriptor) _____
	<u>F Codes- Mental, behavioral and neurodevelopmental disorders</u>
F20.0- F20.9	Schizophrenia (list specific ICD10 code and descriptor) _____
F25.9	Schizoaffective disorder
F30.13	Manic episode, severe, without psychotic symptoms
F30.2	Manic episode, severe with psychotic symptoms
F31.13	Bipolar disorder, current episode manic without psychotic features, severe
F31.2	Bipolar disorder, current episode manic severe with psychotic features
F31.4	Bipolar disorder, current episode depressed, severe, without psychotic features
F31.5	Bipolar disorder, current episode depressed, severe, with psychotic features
F31.63	Bipolar disorder, current episode mixed, severe, without psychotic features
F31.64	Bipolar disorder, current episode mixed, severe, with psychotic features
F31.81	Bipolar disorder II
F33.2	Major depressive disorder, recurrent severe without psychotic features
F33.3	Major depressive disorder, recurrent, severe with psychotic symptoms
	<u>G Codes- Diseases of the nervous system</u>
G31.81	Alpers disease
G31.82	Leigh's disease
G40.001- G40.B19	Epilepsy, (list specific ICD10 code and descriptor) _____
G71.11	Myotonic muscular dystrophy
G80.0	Spastic quadriplegia cerebral palsy
G80.1	Spastic diplegic cerebral palsy
G80.2	Spastic hemiplegic cerebral palsy
G82.50- G82.54	Quadriplegia, (list specific ICD10 code and descriptor) _____
G91.0	Communicating hydrocephalus
G91.1	Obstructive hydrocephalus
G93.1	Anoxic brain damage, not elsewhere classified
	<u>H Codes- H00 to H59 Diseases of the eye and adnexa and H60 to H95 ear and mastoid</u>
H35.051- H35.059	Retinal neovascularization, (list specific ICD10 code and descriptor) _____
H35.171- H35.179	Retrolental fibroplasia, (list specific ICD10 code and descriptor) _____
H54.0	Blindness, binocular

	Diseases of the ear and mastoid process
H90.3	Sensorineural hearing loss, bilateral
H91.9	Hearing Loss
	I Codes- Diseases of the circulatory system
I34.0- I34.2	Nonrheumatic mitral (valve), (list specific ICD10 code and descriptor) _____
I35.0- I35.2	Nonrheumatic aortic (valve), (list specific ICD10 code and descriptor) _____
I36.1- I36.2	Nonrheumatic tricuspid (valve), (list specific ICD10 code and descriptor) _____
I50.1-I50.9	All heart failure, (list specific ICD10 code and descriptor) _____
	J Codes- Diseases of the respiratory system
J45.50- J45.52	Severe persistent asthma, (list specific ICD10 code and descriptor) _____
	K Codes- Diseases of the digestive system
K72.10	Chronic hepatic failure without coma
K72.11	Chronic hepatic failure with coma
<u>K91.2</u>	<u>Short gut syndrome, aka Postsurgical malabsorption</u>
	M Codes- Diseases of the musculoskeletal system and connective tissue
M08.00- M08.48	Juvenile rheumatoid arthritis, (list specific ICD10 code and descriptor) _____
M32.0-M32.9	Systemic lupus erythematosus
M91.11	Juvenile osteochondrosis of head of femur [Legg-Calve-Perthes], right leg
M91.12	Juvenile osteochondrosis of head of femur [Legg-Calve-Perthes], left leg
	N Codes- Diseases of the genitourinary system
N18.4	Chronic kidney disease stage IV (severe)
N18.5	Chronic kidney disease stage V
N18.6	End stage renal disease
	P Codes- Certain conditions originating in the perinatal period
P27.0	Wilson-Mikity syndrome
P27.1	Bronchopulmonary dysplasia originating in the perinatal period
P52.21	Intraventricular (nontraumatic) hemorrhage, grade 3, of newborn
P52.22	Intraventricular (nontraumatic) hemorrhage, grade 4, of newborn
P78.81	Congenital cirrhosis (of liver)
P96.0	Congenital renal failure
	Q Codes- Congenital malformations, deformations, and chromosomal abnormalities
Q00.0	Anencephaly
Q00.1	Craniorachischisis
Q00.2	Iniiencephaly
Q01.0	Frontal encephalocele
Q01.1	Nasofrontal encephalocele
Q01.2	Occipital encephalocele
<u>Q02</u>	<u>Microcephaly</u>
Q03.0	Malformations of aqueduct of Sylvius
Q03.1	Atresia of foramina of Magendie and Luschka
Q04.2	Holoprosencephaly
Q04.4	Septo-optic dysplasia of brain

Q04.5	Megalencephaly
Q05.0- Q05.3 and Q05.5- Q05.8	Spina bifida, (list specific ICD10 code and descriptor) _____
Q06.0	Amyelia
Q06.1	Hypoplasia and dysplasia of spinal cord
Q07.00- Q07.03	Arnold-Chiari syndrome, (list specific ICD10 code and descriptor) _____
<u>Q20.0 to</u> <u>Q26.9</u>	<u>Congenital Malformations of the Circulatory System except Peripheral Vascular System</u> <u>Specify Specific ICD-10 Code</u> _____
<u>Q20.0</u>	<u>Common Arterial Trunk</u>
<u>Q20.1</u>	<u>Double outlet right ventricle</u>
<u>Q20.3</u>	<u>Discordant ventriculoarterial connection</u>
<u>Q20.4</u>	<u>Double inlet ventricle</u>
<u>Q21.3</u>	<u>Tetralogy of Fallot</u>
<u>Q22.0</u>	<u>Pulmonary valve atresia</u>
<u>Q22.5</u>	<u>Ebstein's anomaly</u>
<u>Q22.6</u>	<u>Hypoplastic right heart syndrome</u>
<u>Q23.4</u>	<u>Hypoplastic left heart syndrome</u>
<u>Q24.6</u>	<u>Congenital heart block</u>
<u>Q25.1</u>	<u>Coarctation of aorta</u>
<u>Q25.2</u>	<u>Atresia of aorta</u>
<u>Q26.2</u>	<u>Total anomalous pulmonary venous connection</u>
<u>Q33.3</u>	<u>Agenesis of lung</u>
<u>Q33.6</u>	<u>Congenital hypoplasia and dysplasia of lung</u>
<u>Q35.1</u>	<u>Cleft hard palate</u>
<u>Q35.3</u>	<u>Cleft soft palate</u>
<u>Q35.5</u>	<u>Cleft hard palate with cleft soft palate</u>
<u>Q35.9</u>	<u>Cleft palate, unspecified</u>
<u>Q36.0</u>	<u>Cleft lip, bilateral</u>
<u>Q36.1</u>	<u>Cleft lip, median</u>
<u>Q36.9</u>	<u>Cleft lip, unilateral</u>
<u>Q37.0</u>	<u>Cleft hard palate with bilateral cleft lip</u>
<u>Q37.1</u>	<u>Cleft hard palate with unilateral cleft lip</u>
<u>Q37.2</u>	<u>Cleft soft palate with bilateral cleft lip</u>
<u>Q37.3</u>	<u>Cleft soft palate with unilateral cleft lip</u>
<u>Q37.4</u>	<u>Cleft hard and soft palate with bilateral cleft lip</u>
<u>Q37.5</u>	<u>Cleft hard and soft palate with unilateral cleft lip</u>
<u>Q37.8</u>	<u>Unspecified cleft palate with bilateral cleft lip</u>
<u>Q37.9</u>	<u>Unspecified cleft palate with unilateral cleft lip</u>
<u>Q39.1</u>	<u>Atresia of esophagus with tracheo-esophageal fistula</u>
<u>Q43.1</u>	<u>Hirschsprung's disease</u>
<u>Q44.3</u>	<u>Congenital stenosis and stricture of bile ducts</u>
<u>Q60.1</u>	<u>Renal agenesis, bilateral</u>
<u>Q75.0</u>	<u>Craniosynostosis</u>
<u>Q75.1</u>	<u>Craniofacial dysostosis (Crouzan Syndrome)</u>
<u>Q75.4</u>	<u>Mandibulofacial dysostosis (Treacher Collins)</u>
<u>Q75.5</u>	<u>Oculomandibular dysostosis (Hallermann-Streif syndrome)</u>

Q76.1	Klippel-Feil syndrome
Q78.0	Osteogenesis imperfecta
Q81.2	<u>Epidermolysis bullosa dystrophica</u>
Q91.0- Q91.2	Trisomy 18, nonmosaicism, (list specific ICD10 code and descriptor) _____
Q91.4-Q91.6	Trisomy 13, (list specific ICD10 code and descriptor) _____
	R Codes- Symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified
R40.3	Persistent vegetative state
R75	<u>Inconclusive laboratory evidence of human immunodeficiency virus (HIV)</u>
	T Codes- Injury, poisoning and certain other consequences of external causes
T74.1__	Physical abuse, confirmed, (list specific ICD10 code and descriptor) _____
T74.3__	Psychological abuse, confirmed, (list specific ICD10 code and descriptor) _____
T74.4XXA- T74.4XXS	Shaken infant syndrome, (list specific ICD10 code and descriptor) _____
	Z Codes- Factors influencing health status and contact with health services
Z21	Asymptomatic human immunodeficiency virus [HIV] infection status
Z76.82	Awaiting organ transplant status
Z94.0	Kidney transplant status
Z94.1	Heart transplant status
Z94.2	Lung transplant status
Z94.3	Heart and lungs transplant status
Z94.4	Liver transplant status
Z94.81	Bone marrow transplant status

*** If you are unable to identify the correct ICD 10 CM diagnosis for your patient that represents a condition that is both chronic and serious with functional limitation, at your option as the patient's physician you may request a Medical Panel Review by providing the following additional information for review after obtaining and including consent from the responsible parent or guardian:

1. Specify the ICD 10 Diagnosis with the Alphabetic Code and up to 5 digits _____.

Add Additional Diagnoses, if appropriate _____

2. Include the most recent specialty consultant summary or hospital discharge summary or a summary of the consideration and description of current condition not to exceed three pages together with the following attestations and Florida licensed physician signature

I attest that the child identified above is actively under my care for the management of the chronic and serious condition(s) identified on this form by my initials.

I attest that my patient is under 21 years of age with a chronic physical, developmental, behavioral, or emotional condition, and also requires health care and related services of a type or amount beyond that which is generally required by children.

I attest my patient has functional limits related to his or her ability to do the things most children of the same age can do.

Draft April 12, 2016

Patient Name: _____ DOB: _____

I understand this attestation will be used to establish my patient's clinical eligibility for the CMS Managed Care Plan based on the qualifying diagnoses.

Signed: _____

Print Name: _____

Physician Medicaid #: _____

Physician Medical License #: _____

Date: _____

Office/Practice/Facility Name: _____

Office/Practice/Facility Address: _____

Office/Practice/Facility Phone Number: _____

Office/Practice/Facility Facsimile Number: _____

INSTRUCTIONS: AFTER COMPLETION OF FORM, RETURN TO FLORIDA DEPARTMENT OF HEALTH, OFFICE OF THE CMS MANAGED CARE PLAN BY WAY OF ENCRYPTED EMAIL (CMS.ClinicalEligibilityScreening@flhealth.gov); FACSIMILE (850.488-3813); OR MAIL (Attention: Clinical Eligibility, 4052 Bald Cypress Way, Bin A-06, Tallahassee, Florida, 32399).