ICD Code/Description	EI Eligible/Diagnosed Condition With a High Probability of Developmental Delay?
E44.1 - Mild protein-calorie malnutrition	No
F81.89 - Other developmental disorders of scholastic skills	No
G11.3 - Cerebellar ataxia with defective DNA repair	No
G12.9 - Spinal muscular atrophy, unspecified	No
G31.82 - Leigh's disease	No
G40.201 - Local-rel symptc epi w cmplx prt seiz, not ntrct, w stat epi	No
G70.89 - Other specified myoneural disorders	No
G81.10 - Spastic hemiplegia affecting unspecified side	No
G93.40 - Encephalopathy, unspecified	No
H35.109 - Retinopathy of prematurity, unspecified, unspecified eye	No
H52.13 - Myopia, bilateral	No
H54.12 - Blindness, left eye, low vision right eye	No
H55.09 - Other forms of nystagmus	No
H91.93 - Unspecified hearing loss, bilateral	No
I42.9 - Cardiomyopathy, unspecified	No
I69.351 - Hemiplegia and hemiparesis following cerebral infarction affecting right dominant side	No
J38.5 - Laryngeal spasm	No
M08.3 - Juvenile rheumatoid polyarthritis (seronegative)	No
M08.90 - Juvenile arthritis, unspecified, unspecified site	No
M24.562 - Contracture, left knee	No
M24.9 - Joint derangement, unspecified	No
M62.49 - Contracture of muscle, multiple sites	No
P07.24 - Extreme immaturity of newborn, gestational age 25 completed weeks	No
P91.63 - Severe hypoxic ischemic encephalopathy [HIE]	No
P94.9 - Disorder of muscle tone of newborn, unspecified	No
P96.89 - Other specified conditions originating in the perinatal period	No
Q07.9 - Congenital malformation of nervous system, unspecified	No
Q12.0 - Congenital cataract	No
Q15.0 - Congenital glaucoma	No
Q15.8 - Other specified congenital malformations of eye	No
Q21.9 - Congenital malformation of cardiac septum, unspecified	No
Q24.9 - Congenital malformation of heart, unspecified	No

ICD Code/Description	EI Eligible/Diagnosed Condition With a High Probability of Developmental Delay?
Q26.1 - Persistent left superior vena cava	No
Q31.9 - Congenital malformation of larynx, unspecified	No
Q62.0 - Congenital hydronephrosis	No
Q66.2 - Congenital metatarsus (primus) varus	No
Q66.50 - Congenital pes planus, unspecified foot	No
Q71.00 - Congenital complete absence of unspecified upper limb	No
Q74.0 - Other congenital malformations of upper limb(s), including shoulder girdle	No
Q74.9 - Unspecified congenital malformation of limb(s)	No
Q82.8 - Other specified congenital malformations of skin	No
Q87.0 - Congenital malformation syndromes predominantly affecting facial appearance	No
Q93.4 - Deletion of short arm of chromosome 5	No
Q96.9 - Turner's syndrome, unspecified	No
S14.3XXS - Injury of brachial plexus, sequela	No
Z72.4 - Inappropriate diet and eating habits	No
Z82.79 - Family history of other congenital malformations, deformations and chromosomal abnormalities	No