



Number of individuals receiving DDD-funded services per developmental center, per diagnosis code. Report total may be greater than the total of individuals because some have more than one diagnosis. Due to Federal Limited Data Set requirements, values less than 11 are suppressed.

Center / ICD	GREENBROOK DC	HUNTERDON DC	NEW LISBON DC	VINELAND DC	WOODBINE DC
D84.1,Biotinidase Deficiency	<11				
E70.0,Untreated Phenylketonuria		<11			
E71.121,Propionic Acidemia				<11	
E75.22,Gaucher's Disease		<11			
E77.1,Fucosidosis, Schindler Disease Type 1			<11		
F70,Mild Intellectual Disability	<11	<11	65	<11	14
F71,Moderate Intellectual Disability	<11	11	53	11	<11
F72,Severe Intellectual Disability	<11	29	40	13	<11
F73,Profound Intellectual Disability	39	291	63	106	104
F79,Unspecified Intellectual Disability		<11	<11		11
F84.0,Autistic Disorder (Childhood Autism, Infantile Psychosis, Kanner's Syndrome)	13	79	42	31	29
F84.2,Rett's Syndrome			<11		
F84.5,Other Specified Pervasive Developmental Disorders(Alcohol Embryo and Fetopathy, Asperger Syndrome, Atypical Childhood Psychosis; Borderline Psychosis of Childhood)		<11	<11	<11	<11
F84.9,Unspecified Pervasive Developmental Disorder, Pervasive Developmental Disorder- NOS	<11	35	54	19	16
F89,Unspecified Delay in Development (Developmental Disorder NOS)	<11	<11	<11		11
F99,Psychiatric Disorder or Problem	21	76	142	<11	94
G04.90,Unspecified Cause of Encephalitis		<11	<11	<11	<11
G40.201,Partial Epilepsy, with Impairment of Consciousness (Psychomotor Epilepsy)	<11	<11	<11	<11	<11
G40.309,Generalized Convulsive Epilepsy, Intractable Seizure Disorder	18	82	58	27	43
G40.401,Generalized Non-Convulsive Epilepsy	<11	<11	<11	<11	<11



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G40.409,Grand Mal Status	<11	<11	<11	<11	<11
G40.901,Epilepsy, unspecified with status epilepticus			<11		
G40.909,Epilepsy, unspecified without status epilepticus	12	55	18	29	<11
G60.0,Charcot Marie Tooth Disease		<11			
G80.0,Spastic Cerebral Palsy, Quadriplegic	<11	29	<11	<11	<11
G80.1,Cerebral Palsy, Paraplegic, Congenital	<11	<11	<11	12	<11
G80.2,Cerebral Palsy, Hemiplegic, Congenital/Spastic Hemiplegia		<11	<11	<11	<11
G80.9,Cerebral Palsy, unspecified, Infantile Cerebral Palsy, unspecified	19	24	14	21	29
G82.20,Paraplegia (Paralysis of Both Lower Limbs)	<11	<11		<11	
G82.50,Quadriplegia and Quadriparesis, Unspecified		11	<11	<11	<11
G93.40,Encephalopathy, not elsewhere classified	<11	<11	<11	<11	<11
H54.10,Moderate or Severe Impairment, Better Eye, Profound Impairment Lesser Eye		<11	<11		
H90.8,Mixed Conductive and Sensorineural Hearing Loss		<11			<11
P91.63,Severe Hypoxic Ischemic CNS Injury		<11			<11
Q02,Microcephaly		<11	<11	<11	<11
Q04.3,Agenesis of Septum Pellucidum, Agenesis of the Corpus Callosum, Argyria, Pachygyria, Microgyria, Lissencephaly					<11
Q05.9,Spina Bifida , Unspecified		<11			
Q07.9,Unspecified Anomaly of Brain, Spinal Cord, and Nervous System		<11	<11	<11	<11
Q85.1,Tuberous Sclerosis	<11	<11	<11	<11	<11
Q85.8,Sturge-Weber Syndrome				<11	
Q86.0,Fetal Alcohol Syndrome		<11		<11	<11
Q87.11,Prader-Willi Syndrome			<11		



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Q87.19,Other congenital malformation syndromes predominantly associated with short stature				<11	
Q87.2,Rubinstien-Taybi Syndrome, Vater Association		<11	<11		
Q89.8,Kabuki Syndrome, Coffin-Lowry Syndrome, Cockayne Syndrome, Cornelia de Lange Syndrome, CHARGE Association				<11	
Q90.9,Down Syndrome	<11	<11	<11	<11	<11
Q91.3,Trisomy 18, Unspecified		<11			
Q93.51,Angelman Syndrome		<11		<11	
Q98.4,Klinefelter's Syndrome					<11
Q99.2,Fragile X Syndrome					<11
Q99.8,Other Chromosomal Abnormalities, not elsewhere classified		<11			
S04.019A,Unspecified (Traumatic Blindness NOS)	<11	<11		<11	44
S09.90XA,Head Injury, unspecified					<11