



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			
E75.22,Gaucher's Disease		<11			
E77.1,Fucosidosis, Schindler Disease Type 1			<11		
F70,Mild Intellectual Disability	<11	<11	52	<11	16
F71,Moderate Intellectual Disability	<11	<11	45	<11	<11
F72,Severe Intellectual Disability	<11	23	35	13	<11
F73,Profound Intellectual Disability	35	257	59	94	93
F79,Unspecified Intellectual Disability		<11	<11		<11
F84.0,Autistic Disorder (Childhood Autism, Infantile Psychosis, Kanner's Syndrome)	12	69	41	30	28
F84.5,Other Specified Pervasive Developmental Disorders(Alcohol Embryo and Fetopathy, Asperger Syndrome, Atypical Childhood Psychosis; Borderline Psychosis of Childhood)		<11	<11		<11
F84.9,Unspecified Pervasive Developmental Disorder, Pervasive Developmental Disorder- NOS	<11	32	48	18	17
F89,Unspecified Delay in Development (Developmental Disorder NOS)	<11	<11	<11		<11
F99,Psychiatric Disorder or Problem	18	67	121	<11	87
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	17	72	49	25	38
G40.401,Generalized Non-Convulsive Epilepsy	<11	<11	<11	<11	<11
G40.409,Grand Mal Status		<11	<11	<11	<11
G40.901,Epilepsy, unspecified with status epilepticus			<11		



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G40.909,Epilepsy, unspecified without status epilepticus	11	47	14	27	<11
G60.0,Charcot Marie Tooth Disease		<11			
G80.0,Spastic Cerebral Palsy, Quadriplegic	<11	25		<11	<11
G80.1,Cerebral Palsy, Paraplegic, Congenital	<11	<11	<11	11	<11
G80.2,Cerebral Palsy, Hemiplegic, Congenital/Spastic Hemiplegia		<11	<11	<11	<11
G80.9,Cerebral Palsy, unspecified, Infantile Cerebral Palsy, unspecified	17	20	14	16	28
G82.20,Paraplegia (Paralysis of Both Lower Limbs)	<11	<11		<11	
G82.50,Quadriplegia and Quadriparesis, Unspecified		<11	<11	<11	
G93.40,Encephalopathy, not elsewhere classified	<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
Q85.89,Other phakomatoses, not elsewhere classified				<11	
Q86.0,Fetal Alcohol Syndrome		<11		<11	<11
Q87.11,Prader-Willi Syndrome			<11		
Q87.19, Cockayne Syndrome, Cornelia de Lange Syndrome and Other congenital malformation syndromes predominantly associated with short stature				<11	
Q87.2,Rubinstien-Taybi Syndrome, Vater Association		<11	<11		



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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
Q99.89, Other Specified Chromosome Abnormalities		<11	<11		
S04.019A,Unspecified (Traumatic Blindness NOS)	<11	<11		<11	39
S09.90XA,Head Injury, unspecified					<11