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Developmental and epileptic encephalopathies: recognition and approaches to care

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22	De novo SCN8A and inherited rare CACNA1H variants associated with severe developmental and epileptic encephalopathy. <i>Molecular Brain</i> , 2021 , 14, 126	4.5	1
21	The epilepsy-autism spectrum disorder phenotype in the era of molecular genetics and precision therapy. <i>Epilepsia</i> , 2021 ,	6.4	5
20	Multidisciplinary Care of Patients with Inherited Metabolic Diseases and Epilepsy: Current Perspectives <i>Journal of Multidisciplinary Healthcare</i> , 2022 , 15, 553-566	2.8	1
19	Genetic analysis using targeted exome sequencing of 53 Vietnamese children with developmental and epileptic encephalopathies <i>American Journal of Medical Genetics, Part A</i> , 2022 ,	2.5	
18	Difficulties of Prenatal Genetic Counseling for a Subsequent Child in a Family With Multiple Genetic Variations <i>Frontiers in Genetics</i> , 2021 , 12, 612100	4.5	
17	Detection of Deregulated miRNAs in Childhood Epileptic Encephalopathies <i>Journal of Molecular Neuroscience</i> , 2022 , 1	3.3	1
16	Precision medicine for developmental and epileptic encephalopathies in Africa Btrategies for a resource-limited setting.		
15	Monogenic developmental and epileptic encephalopathies of infancy and childhood, a population cohort from Norway. 10,		1
14	Emerging evidence of genotypephenotype associations of developmental and epileptic encephalopathy due to KCNC2 mutation: Identification of novel R405G. 15,		
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4	Cannabinoid Efficacy for Developmental Epileptic Encephalopathy (DEE) Intractable Seizure Control. 2023 , 76-102	O
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