Stephanie Fehr

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5862576/publications.pdf

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9	639	1163117 8	1474206 9
papers	citations	h-index	g-index

9	9	9	764
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The CDKL5 disorder is an independent clinical entity associated with early-onset encephalopathy. European Journal of Human Genetics, 2013, 21, 266-273.	2.8	220
2	Trends in the Diagnosis of Rett Syndrome in Australia. Pediatric Research, 2011, 70, 313-319.	2.3	119
3	There is variability in the attainment of developmental milestones in the CDKL5 disorder. Journal of Neurodevelopmental Disorders, 2015, 7, 2.	3.1	74
4	Seizure variables and their relationship to genotype and functional abilities in the CDKL5 disorder. Neurology, 2016, 87, 2206-2213.	1.1	74
5	Functional abilities in children and adults with the CDKL5 disorder. American Journal of Medical Genetics, Part A, 2016, 170, 2860-2869.	1.2	65
6	Altered Attainment of Developmental Milestones Influences the Age of Diagnosis of Rett Syndrome. Journal of Child Neurology, 2011, 26, 980-987.	1.4	37
7	Atypical presentations and specific genotypes are associated with a delay in diagnosis in females with Rett syndrome. American Journal of Medical Genetics, Part A, 2010, 152A, 2535-2542.	1.2	31
8	Overview of Health Issues in School-aged Children with Down Syndrome. International Review of Research in Mental Retardation, 2010, 39, 67-106.	0.7	12
9	What does the nature of the MECP2 mutation tell us about parental origin and recurrence risk in Rett syndrome?. Clinical Genetics, 2012, 82, 526-533.	2.0	7