Anne Maria Rochtus

List of Publications by Year
in descending order

14	384	1040056 9	1199594 12
papers	citations	h-index	g-index

15	15	15	909
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Antiseizure medications and thyroid hormone homeostasis: Literature review and practical recommendations. Epilepsia, 2022, 63, 259-270.	5.1	9
2	Antiepileptic Drugs and Thyroid Hormone Homeostasis: Literature Review and Practical Guideline. Journal of the Endocrine Society, 2021, 5, A824-A824.	0.2	0
3	<scp>BRAT</scp>1 encephalopathy: a recessive cause of epilepsy of infancy with migrating focal seizures. Developmental Medicine and Child Neurology, 2020, 62, 1096-1099.	2.1	18
4	SAT-240 Hypothalamic Lipoma and Growth Hormone Deficiency. Journal of the Endocrine Society, 2020, 4, .	0.2	0
5	Antiepileptic Drug Teratogenicity and De Novo Genetic Variation Load. Annals of Neurology, 2020, 87, 897-906.	5.3	9
6	Genetic diagnoses in epilepsy: The impact of dynamic exome analysis in a pediatric cohort. Epilepsia, 2020, 61, 249-258.	5.1	85
7	Expanding the clinical spectrum of Fowler syndrome: Three siblings with survival into adulthood and systematic review of the literature. Clinical Genetics, 2020, 98, 423-432.	2.0	4
8	The role of sodium channels in sudden unexpected death in pediatrics. Molecular Genetics & Genomic Medicine, 2020, 8, e1309.	1.2	14
9	Hypothalamic lipoma and growth hormone deficiency. International Journal of Pediatric Endocrinology (Springer), 2020, 2020, 4.	1.6	5
10	The Genetic Landscape of Epilepsy of Infancy with Migrating Focal Seizures. Annals of Neurology, 2019, 86, 821-831.	5.3	96
11	Spectrum of neurodevelopmental disease associated with the GNAO1 guanosine triphosphateâ€“binding region. Epilepsia, 2019, 60, 406-418.	5.1	53
12	Mutations in NRXN1 and NRXN2 in a patient with early-onset epileptic encephalopathy and respiratory depression. Journal of Physical Education and Sports Management, 2019, 5, a003442.	1.2	20
13	Methylome analysis for spina bifida shows SOX18 hypomethylation as a risk factor with evidence for a complex (epi)genetic interplay to affect neural tube development. Clinical Epigenetics, 2016, 8, 108.	4.1	18
14	Genome-wide DNA methylation analysis of pseudohypoparathyroidism patients with GNAS imprinting defects. Clinical Epigenetics, 2016, 8, 10.	4.1	53