Hunmin Kim

List of Publications by Year
in descending order

45	541	687220 13	752573 20
papers	citations	h-index	g-index

48	48	48	1042
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Surgical outcome and predictive factors of epilepsy surgery in pediatric isolated focal cortical dysplasia. Epilepsy Research, 2018, 139, 54-59.	0.8	55
2	Lacosamide as an adjunctive therapy in pediatric patients with refractory focal epilepsy. Brain and Development, 2014, 36, 510-515.	0.6	39
3	Clinical and EEG risk factors for subsequent epilepsy in patients with complex febrile seizures. Epilepsy Research, 2013, 105, 158-163.	0.8	27
4	Serum Î±-synuclein and IL-1Î² are increased and correlated with measures of disease severity in children with epilepsy: potential prognostic biomarkers?. BMC Neurology, 2020, 20, 85.	0.8	27
5	Diagnostic Yield of Epilepsy Panel Testing in Patients With Seizure Onset Within the First Year of Life. Frontiers in Neurology, 2019, 10, 988.	1.1	26
6	Screening Autoimmune Anti-neuronal Antibodies in Pediatric Patients with Suspected Autoimmune Encephalitis. Journal of Epilepsy Research, 2014, 4, 55-61.	0.1	24
7	Leigh Syndrome in Childhood: Neurologic Progression and Functional Outcome. Journal of Clinical		

#	Article	IF	Citations
19	Spike persistence and normalization in benign epilepsy with centrotemporal spikes â€“ Implications for management. Brain and Development, 2018, 40, 693-698.	0.6	11
20	Hypokalemic periodic paralysis; two different genes responsible for similar clinical manifestations. Korean Journal of Pediatrics, 2011, 54, 473.	1.9	11
21	Averaged EEG spike dipole analysis may predict atypical outcome in Benign Childhood Epilepsy with Centrotemporal Spikes (BCECTS). Brain and Development, 2016, 38, 903-908.	0.6	10
22	Familial cases of progressive myoclonic epilepsy caused by maternal somatic mosaicism of a recurrent KCNC1 p.Arg320His mutation. Brain and Development, 2018, 40, 429-432.	0.6	10
23	Antiepileptic Drug Withdrawal after Surgery in Children with Focal Cortical Dysplasia: Seizure		

#	Article	IF	Citations
37	The Role of Focal Epilepsy Features in Defining <i>SCN1A</i> Mutation-positive Dravet Syndrome as Generalized and Focal Epilepsy. Journal of Epilepsy Research, 2021, 11, 127-135.	0.1	3
38	Dissecting the phenotypic and genetic spectrum of early childhood-onset generalized epilepsies. Seizure: the Journal of the British Epilepsy Association, 2019, 71, 222-228.	0.9	2
39	Clinical and Genetic Spectrum ofATP1A3-Related Disorders in a Korean Pediatric Population.		