Jean-Baptiste Le Pichon

List of Publications by Year
in descending order

22	839	933447 10	888059 17
papers	citations	h-index	g-index

23	23	23	2187
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genomic answers for children: Dynamic analyses of >1000 pediatric rare disease genomes. Genetics in Medicine, 2022, 24, 1336-1348.	2.4	37
2	Combined use of the ketogenic diet and vagus nerve stimulation in pediatric drugâ€resistant epilepsy. Epilepsia Open, 2021, 6, 112-119.	2.4	4
3	The efficacy and tolerability of auto-stimulation-VNS in children with Lennox-Gastaut syndrome. Seizure: the Journal of the British Epilepsy Association, 2021, 86, 168-174.	2.0	12
4	Diagnostic yield of genetic testing in 324 infants with hypotonia. Clinical Genetics, 2021, 100, 752-757.	2.0	4
5	Pediatric Head Trauma: A Review and Update. , 2021, , 28-39.		0
6	Seven-Year Experience From the National Institute of Neurological Disorders and Strokeâ€“Supported Network for Excellence in Neuroscience Clinical Trials. JAMA Neurology, 2020, 77, 755.	9.0	6
7	Transition to Adult Care in Youth with Epilepsy: One Center's Experience with a Transition Program and Its Integration within Telemedicine. Journal of Pediatric Epilepsy, 2020, 09, 119-124.	0.2	1
8	The 2018 Pediatric Neurology Trainee Publication Award. Pediatric Neurology, 2019, 101, 1.	2.1	0
9	Safety and tolerability of adjunctive lacosamide in a pediatric population with focal seizures â€“ An open-label trial. Seizure: the Journal of the British Epilepsy Association, 2019, 71, 166-173.	2.0	21
10	Initiating the ketogenic diet in infants with treatment refractory epilepsy while maintaining a breast milk diet. Seizure: the Journal of the British Epilepsy Association, 2019, 69, 41-43.	2.0	16
11	The 2017 Pediatric Neurology Training Publication Award. Pediatric Neurology, 2018, 86, 4.	2.1	0
12	Brown-Vialetto-Van Laere Syndrome as a Mimic of Neuroimmune Disorders: 3 Cases From the Clinic and Review of the Literature. Journal of Child Neurology, 2017, 32, 528-532.	1.4	15
13	Editorial: The 2016 Pediatric Neurology Trainee Publication Award. Pediatric Neurology, 2017, 75, 3.	2.1	0
14	The Neurological Sequelae of Neonatal Hyperbilirubinemia: Definitions, Diagnosis and Treatment of the Kernicterus Spectrum Disorders (KSDs). Current Pediatric Reviews, 2017, 13, 199-209.	0.8	96
15	A Hypothesis for Using Pathway Genetic Load Analysis for Understanding Complex Outcomes in Bilirubin Encephalopathy. Frontiers in Neuroscience, 2016, 10, 376.	2.8	14
16	Biallelic Mutations in TBCD , Encoding the Tubulin Folding Cofactor D, Perturb Microtubule Dynamics and Cause Early-Onset Encephalopathy. American Journal of Human Genetics, 2016, 99, 962-973.	6.2	66
17	Hyperactivity in the Gunn rat model of neonatal jaundice: age-related attenuation and emergence of gait deficits. Pediatric Research, 2015, 77, 434-439.	2.3	9
18	The Future of Next-Generation Sequencing in Neurology. JAMA Neurology, 2015, 72, 971.	9.0	6

#	Article	IF	Citations
19	Effectiveness of exome and genome sequencing guided by acuity of illness for diagnosis of neurodevelopmental disorders. Science Translational Medicine, 2014, 6, 265ra168.	12.4	440
20	Genome-wide gene expression in a patient with 15q13.3 homozygous microdeletion syndrome. European Journal of Human Genetics, 2013, 21, 1093-1099.	2.8	23
21	A 15q13.3 homozygous microdeletion associated with a severe neurodevelopmental disorder suggests putative functions of the TRPM1, CHRNA7, and other homozygously deleted genes. American Journal of Medical Genetics, Part A, 2010, 152A, 1300-1304.	1.2	55
22	Acute bilirubin encephalopathy and its progression to kernicterus: current perspectives. Research and Reports in Neonatology, 0, Volume 8, 33-44.	0.2	14