

Peter Humphreys

List of Publications by Year in descending order

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Version: 2024-02-01

15
papers

495
citations

933447

10
h-index

1058476

14
g-index

15
all docs

15
docs citations

15
times ranked

1169
citing authors

#	ARTICLE	IF	CITATIONS
1	Epidemiology and Outcomes of Arterial Ischemic Stroke in Children: The Canadian Pediatric Ischemic Stroke Registry. <i>Pediatric Neurology</i> , 2017, 69, 58-70.	2.1	213
2	Homozygous nonsense mutation in <i>SYNJ1</i> associated with intractable epilepsy and tau pathology. <i>Neurobiology of Aging</i> , 2015, 36, 1222.e1-1222.e5.	3.1	50
3	Congenital Visual Impairment and Progressive Microcephaly Due to Lysyl-Transfer Ribonucleic Acid (RNA) Synthetase (<i>KARS</i>) Mutations. <i>Journal of Child Neurology</i> , 2015, 30, 1037-1043.	1.4	47
4	Clinical Guidelines for Management of Bone Health in Rett Syndrome Based on Expert Consensus and Available Evidence. <i>PLoS ONE</i> , 2016, 11, e0146824.	2.5	45
5	The Incidence and Evolution of Parkinsonian Rigidity in Rett Syndrome: A Pilot Study. <i>Canadian Journal of Neurological Sciences</i> , 2016, 43, 567-573.	0.5	28
6	Factors Associated With Epilepsy in Children With Periventricular Leukomalacia. <i>Journal of Child Neurology</i> , 2007, 22, 598-605.	1.4	26
7	Brain malformations in a patient with deletion 2p16.1: A refinement of the phenotype to <i>BCL11A</i> . <i>European Journal of Medical Genetics</i> , 2015, 58, 351-354.	1.3	24
8	Association of Early-Onset Spasticity and Risk for Cognitive Impairment With Mutations at Amino Acid 499 in <i>SPAST</i> . <i>Journal of Child Neurology</i> , 2018, 33, 329-332.	1.4	20
9	Focal Cerebral Mantle Disruption in Fetal Hydrocephalus. <i>Pediatric Neurology</i> , 2007, 36, 236-243.	2.1	14
10	Clinical delineation of <i>GTPBP2</i> -associated neuroectodermal syndrome: Report of two new families and review of the literature. <i>Clinical Genetics</i> , 2019, 95, 601-606.	2.0	11
11	Septo-optic-pituitary dysplasia. <i>Handbook of Clinical Neurology</i> / Edited By P J Vinken and G W Bruyn, 2007, 87, 39-52.	1.8	5
12	Atypical Rett Syndrome and Intractable Epilepsy With Novel <i>GRIN2B</i> Mutation. <i>Child Neurology Open</i> , 2018, 5, 2329048X1878794.	1.1	5
13	Analysis of cerebrocortical neuronal migration in three-dimensional fetal mouse cerebral explants: comparison with in vivo. <i>International Journal of Developmental Neuroscience</i> , 2000, 18, 573-584.	1.6	4
14	The Magic Mountain—A Time Capsule of Tuberculosis Treatment in the Early Twentieth Century. <i>Canadian Bulletin of Medical History = Bulletin Canadien D'histoire De La Médecine</i> , 1989, 6, 147-163.	0.1	3
15	What are the reasons for improved survival following spine fusion for severe scoliosis in Rett syndrome?. <i>Developmental Medicine and Child Neurology</i> , 2016, 58, 538-539.	2.1	0