

Mahesh Kamate

List of Publications by Year in descending order

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56
papers

287
citations

1163117

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1058476

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59
all docs

59
docs citations

59
times ranked

433
citing authors

#	ARTICLE	IF	CITATIONS
1	Disorders of Tetrahydrobiopterin Metabolism: Experience from South India. <i>Metabolic Brain Disease</i> , 2022, 37, 743-760.	2.9	4
2	Perinatal Infections: An Important Etiological Risk Factor for Mineralizing Angiopathy. <i>Indian Journal of Pediatrics</i> , 2021, 88, 58-60.	0.8	3
3	Mitochondrial DNA Depletion Syndrome: Mimicker for Hereditary Tyrosinemia. <i>Indian Journal of Pediatrics</i> , 2021, 88, 377-377.	0.8	2
4	Neuronal ceroid lipofuscinoses in children. <i>Annals of Indian Academy of Neurology</i> , 2021, 24, 192.	0.5	4
5	Acute Leucoencephalopathy with Restricted Diffusion in Children – A case series. <i>Neurology India</i> , 2021, 69, 466.	0.4	7
6	Role of Antiplatelet Therapy in Pediatric Stroke with Mineralizing Angiopathy. <i>Indian Journal of Pediatrics</i> , 2021, 88, 1152-1152.	0.8	2
7	Antenatal presentation and supratentorial brain abnormalities in a child with Poretti-Boltshauser syndrome. <i>Brain and Development</i> , 2021, , .	1.1	2
8	Early-onset phenotype of bi-allelic <i>GRN</i> mutations. <i>Brain</i> , 2021, 144, e22-e22.	7.6	5
9	Arterial ischemic stroke outcomes in children: Indian perspective. <i>Journal of Pediatric Neurosciences</i> , 2021, 16, 182.	0.3	0
10	Epilepsy Surgery in Children in Resource Poor Setting. <i>Indian Journal of Pediatrics</i> , 2020, 87, 1081-1081.	0.8	0
11	Theme: Neurology and Development. <i>Indian Pediatrics</i> , 2020, 57, 694-694.	0.4	0
12	Do All Children With Autoimmune Encephalitis Need Aggressive Immunotherapy?. <i>Indian Pediatrics</i> , 2020, 57, 1087-1088.	0.4	0
13	Clinico-Investigative Profile of Infantile Tremor Syndrome. <i>Indian Journal of Pediatrics</i> , 2020, 87, 169-170.	0.8	2
14	Clinical Profile and Short-term Outcome of Pediatric Status Epilepticus. <i>Indian Pediatrics</i> , 2020, 57, 207-208.	0.4	1
15	CAD Deficiency – Another Treatable Early Infantile Epileptic Encephalopathy. <i>Pediatric Neurology</i> , 2020, 110, 97-98.	2.1	3
16	Clinical Profile and Short-term Outcome of Pediatric Status Epilepticus. <i>Indian Pediatrics</i> , 2020, 57, 207-208.	0.4	0
17	Vitamin K Prophylaxis: Is There a Need for Policy Change?. <i>Indian Journal of Pediatrics</i> , 2019, 86, 975-976.	0.8	1
18	Neuronal ceroid lipofuscinosis type-11 in an adolescent. <i>Brain and Development</i> , 2019, 41, 542-545.	1.1	25

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19	Do All Children with Guillain Barré Syndrome Need Immunotherapy?. Indian Journal of Pediatrics, 2019, 86, 479-479.	0.8	0
20	Uncommon Treatable Genetic Epileptic Encephalopathies. Indian Pediatrics, 2019, 56, 427-428.	0.4	0
21	Effect of Risperidone on the Motor and Functional Disability in Children With Choreoathetoid Cerebral Palsy. Pediatric Neurology, 2018, 84, 46-48.	2.1	3
22	Prevalence of Hypocalcemia in Seizures in Infancy. Indian Journal of Pediatrics, 2018, 85, 307-308.	0.8	4
23	Bilateral Ptosis: A Rare Presentation of Hydrocephalus. Indian Journal of Pediatrics, 2018, 85, 398-399.	0.8	3
24	Subacute Sclerosing Panencephalitis: A Disease Not to be Forgotten. Indian Pediatrics, 2018, 55, 708-709.	0.4	2
25	Neonatal Rotaviral Encephalitis. Indian Journal of Pediatrics, 2017, 84, 865-866.	0.8	3
26	Guillaine-Barre syndrome with retained deep tendon reflexes. Indian Pediatrics, 2017, 54, 887-887.	0.4	0
27	The Effectiveness of a Computer Game-Based Rehabilitation Platform for Children With Cerebral Palsy: Protocol for a Randomized Clinical Trial. JMIR Research Protocols, 2017, 6, e93.	1.0	11
28	Autistic Regression: Should it Prompt Urgent EEG?. Indian Pediatrics, 2017, 54, 787.	0.4	0
29	Spongy White Matter: A Novel Neuroimaging Finding in Canavan Disease. Pediatric Neurology, 2016, 56, 92-93.	2.1	3
30	Familial hypomagnesemia with secondary hypocalcemia mimicking neurodegenerative disorder. Indian Pediatrics, 2015, 52, 521-522.	0.4	9
31	Macroamylasemia: A benign cause for high serum amylase. Indian Pediatrics, 2015, 52, 533-539.	0.4	2
32	Safety of lacosamide in children with refractory partial epilepsy. Saudi Pharmaceutical Journal, 2015, 23, 556-561.	2.7	8
33	CDKL5 Encephalopathy: A Rare Cause of Infantile Epileptic Encephalopathy. Indian Pediatrics, 2015, 52, 537.	0.4	0
34	Punched Holes in Globus Pallidi: A Novel Neuroimaging Finding in Wilson Disease. Pediatric Neurology, 2014, 50, 281-282.	2.1	2
35	Editorials. Indian Pediatrics, 2014, 51, 445-450.	0.4	0
36	L-2-Hydroxyglutaric Aciduria: Report of Two Indian Families. Indian Journal of Pediatrics, 2014, 81, 296-298.	0.8	2

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37	Efficacy and Tolerability of Lacosamide as an Adjunctive Therapy in Children With Refractory Partial Epilepsy. <i>Pediatric Neurology</i> , 2014, 51, 509-514.	2.1	24
38	Fructose-1,6-diphosphatase deficiency: a treatable neurometabolic disorder. <i>BMJ Case Reports</i> , 2014, 2014, bcr2013201553-bcr2013201553.	0.5	6
39	“Reversible Inattention and hyperactivity in a child on Lacosamide - In south Indian Subpopulation”™. <i>IOSR Journal of Pharmacy and Biological Sciences</i> , 2014, 9, 66-67.	0.1	6
40	INDT-ADHD as a diagnostic tool for ADHD in Indian children. <i>Indian Pediatrics</i> , 2014, 51, 449-50.	0.4	0
41	Glutaric aciduria type I: A treatable neurometabolic disorder. <i>Annals of Indian Academy of Neurology</i> , 2012, 15, 31.	0.5	16
42	Clinico-investigative profile of infantile and late-infantile neuronal ceroid lipofuscinoses. <i>Neurology India</i> , 2012, 60, 316.	0.4	11
43	Novel Neuroimaging Finding in Palmitoyl Protein Thioesterase-1-Related Neuronal Ceroid Lipofuscinosis. <i>Pediatric Neurology</i> , 2012, 46, 325-328.	2.1	4
44	Subacute Sclerosing Panencephalitis in a Toddler. <i>Indian Journal of Pediatrics</i> , 2012, 79, 1384-1385.	0.8	2
45	Predominant Corticospinal Tract Involvement in Early-Onset Krabbe Disease. <i>Pediatric Neurology</i> , 2011, 44, 155-156.	2.1	9
46	Normal Neuroimaging in Early-Onset Krabbe Disease. <i>Pediatric Neurology</i> , 2011, 44, 374-376.	2.1	3
47	“Eye-of-the-Tiger”™ Sign and Classic Pantothenate Kinase Associated Neurodegeneration. <i>Indian Journal of Pediatrics</i> , 2011, 78, 121-122.	0.8	1
48	Accidental Endosulfan Ingestion in a Toddler. <i>Indian Journal of Pediatrics</i> , 2011, 78, 884-885.	0.8	1
49	Andersen-Tawil syndrome “” Periodic paralysis with dysmorphism. <i>Indian Pediatrics</i> , 2011, 48, 64-65.	0.4	3
50	Benign partial seizures of adolescence. <i>Indian Pediatrics</i> , 2011, 48, 393-395.	0.4	0
51	MRI abnormalities of the anterior temporal lobe: a new indicator of congenital cytomegalovirus infection. <i>Indian Pediatrics</i> , 2011, 48, 325-8.	0.4	3
52	Profile of inborn errors of metabolism in a tertiary care centre PICU. <i>Indian Journal of Pediatrics</i> , 2010, 77, 57-60.	0.8	25
53	Cerebrotendinous xanthomatosis. <i>Indian Journal of Pediatrics</i> , 2010, 77, 697-698.	0.8	5
54	“Glutaric aciduria type I- an easily diagnosable and treatable metabolic disorder”™. <i>Indian Journal of Pediatrics</i> , 2009, 76, 562-563.	0.8	4

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55	Fulminant Cerebellitis: A Fatal, Clinically Isolated Syndrome. <i>Pediatric Neurology</i> , 2009, 41, 220-222.	2.1	41
56	Early myoclonic encephalopathy. <i>Indian Pediatrics</i> , 2009, 46, 804-6.	0.4	1