Nutan Kamath

List of Publications by Year
in descending order

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27 papers	134 citations	1478280 6 h-index	1372474 10 g-index
27 all docs	27 docs citations	27 times ranked	185 citing authors

#	Article	IF	Citations
1	Pseudoachondroplasia: Phenotype and genotype in 11 Indian patients. American Journal of Medical Genetics, Part A, 2022, 188, 751-759.	0.7	1
2	Clinical and Genetic Spectrum of Inborn Errors of Immunity in a Tertiary Care Center in Southern India. Indian Journal of Pediatrics, 2022, 89, 233-242.	0.3	4
3	Improving Underweight Mothers' Essential Newborn Care During Early Infancy: A Single-Blinded, Parallel-Randomized, Controlled Trial. Journal of Pediatrics, 2022, 244, 72-78.e2.	0.9	1
4	Perceptions of teachers towards COVID appropriate behaviors for school children in coastal South India. Current Psychology, 2022, 41, 8112-8122.	1.7	4
5	Can Flip-Chart Assisted Maternal Education Improve Essential New Born Care Knowledge and Skills? A Randomized Controlled Trial. Maternal and Child Health Journal, 2022, , 1.	0.7	0
6	A Delphi survey based construction and validation of test for oropharyngeal dysphagia in Indian neonates. International Journal of Pediatric Otorhinolaryngology, 2021, 140, 110306.	0.4	1
7	Clinical and genetic spectrum of 104 Indian families with central nervous system white matter abnormalities. Clinical Genetics, 2021, 100, 542-550.	1.0	12
8	Hypovitaminosis D and Parathyroid Hormone Response in Critically Ill Children with Sepsis: A Case-control Study. Indian Journal of Critical Care Medicine, 2021, 25, 923-927.	0.3	2
9	Development and utility of tools to identify preventable perinatal deaths: results from a community-based interventional study in two districts of Karnataka State, India. Indian Journal of Community Medicine, 2021, 46, 631.	0.2	0
10	Phenotypic diversity and genetic complexity of PAX3 â€related Waardenburg syndrome. American Journal of Medical Genetics, Part A, 2020, 182, 2951-2958.	0.7	8
11	Resource limited centres can deliver treatment for children with acute lymphoblastic leukaemia with risk-stratified minimal residual disease based UKALL 2003 protocol with no modification and a good outcome. Expert Review of Hematology, 2020, 13, 1143-1151.	1.0	6
12	A case report of acute disseminated encephalomyelitis following severe dengue in a child. Germs, 2020, 10, 115-119.	0.5	1
13	Antenatal Ovarian Torsion Presenting With Features of Intestinal Obstruction in a Neonate. Journal of Nepal Paediatric Society, 2020, 40, 265-269.	0.1	0
14	Documentation and Reporting of Perinatal Deaths in Two Districts of Karnataka, India: A Situational Analysis. Indian Pediatrics, 2020, 57, 1006-1009.	0.2	2
15	Modified Sick Neonatal Score (MSNS): A Novel Neonatal Disease Severity Scoring System for Resource-Limited Settings. Critical Care Research and Practice, 2019, 2019, 1-6.	0.4	12
16	Thyroid Function in Chronically Transfused Children with Beta Thalassemia Major: A Cross-Sectional Hospital Based Study. International Journal of Pediatrics (United Kingdom), 2018, 2018, 1-5.	0.2	13
17	Electrocardiographic Analysis of Repolarization Changes in South Indian Children with Kawasaki Disease after the Acute Phase of Illness. International Journal of Pediatrics (United Kingdom), 2018, 2018, 1-5.	0.2	3
18	The emergence of Kawasaki disease in India and China. Global Cardiology Science & Practice, 2018, 2017, e201721.	0.3	20

#	Article	IF	Citations
19	Influenza B virus triggering macrophage activation syndrome in an infant. Indian Journal of Critical Care Medicine, 2017, 21, 802-803.	0.3	2
20	Audit of Organic Acidurias from a Single Centre: Clinical and Metabolic Profile at Presentation with Long Term Outcome. Journal of Clinical and Diagnostic Research JCDR, 2017, 11, SC11-SC14.	0.8	7
21	Homozygous deletion of exons 2 and 3 of NPC2 associated with Niemannâ€“Pick disease type C. American Journal of Medical Genetics, Part A, 2016, 170, 2486-2489.	0.7	5
22	Metabolic Syndrome in Childhood: Rare Case of Alstrom Syndrome with Blindness. Indian Journal of Clinical Biochemistry, 2016, 31, 480-482.	0.9	3
23	Recurrent and novel GLB1 mutations in India. Gene, 2015, 567, 173-181.	1.0	22
24	Necrotizing fasciitis in children: Experience in a teaching hospital. Journal of Pediatric Infectious Diseases, 2015, 02, 225-229.	0.1	1
25	Bronze Diabetes. Journal of Clinical and Diagnostic Research JCDR, 2015, 9, BD01-2.	0.8	0
26	DEVELOPMENTAL DELAY IN CHILDHOOD CATARACT: A CAVEAT MARINESCO-SJÃ–GREN SYNDROME. Journal of Health and Allied Sciences NU, 2014, 04, 121-123.	0.1	0
27	Biochemical Basis of Heterogeneity in Acute Presentations of Propionic Acidemia. Indian Journal of Clinical Biochemistry, 2013, 28, 95-97.	0.9	4