Shivaram Shastri

List of Publications by Year
in descending order

12	273	1163117 8	1281871 11
papers	citations	h-index	g-index

13	13	13	409
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Genetic polymorphisms associated with obesity and non-alcoholic fatty liver disease in Asian Indian adolescents. Journal of Pediatric Endocrinology and Metabolism, 2019, 32, 749-758.	0.9	13
2	Aquagenic Wrinkling of Skin: A Screening Test for Cystic Fibrosis. Indian Pediatrics, 2019, 56, 109-113.	0.4	9
3	Aquagenic Wrinkling of Skin: A Screening Test for Cystic Fibrosis. Indian Pediatrics, 2019, 56, 109-113.	0.4	0
4	Identification of a novel homozygous mutation in transmembrane channel like 1 () gene, one of the second-tier hearing loss genes after in India. Indian Journal of Medical Research, 2017, 145, 492-497.	1.0	3
5	ADRB2 polymorphism and salbutamol responsiveness in Northern Indian children with mild to moderate exacerbation of asthma. Indian Pediatrics, 2016, 53, 211-215.	0.4	8
6	Zinc Supplementation for One Year Among Children with Cystic Fibrosis Does Not Decrease Pulmonary Infection. Respiratory Care, 2016, 61, 78-84.	1.6	26
7	Abstract 2040: Influence of MDR1 and CYP3A5 genetic polymorphisms on trough levels and therapeutic response of imatinib in patients with chronic myeloid leukemia. , 2016, , .		0
8	Prevalence of UGT1A6 polymorphisms in children with epilepsy on valproate monotherapy. Neurology India, 2015, 63, 35.	0.4	16
9	Glutaric Acidemia Type 1-Clinico-Molecular Profile and Novel Mutations in GCDH Gene in Indian Patients. JIMD Reports, 2014, 21, 45-55.	1.5	25
10	Norrie Disease: First Mutation Report and Prenatal Diagnosis in an Indian Family. Indian Journal of Pediatrics, 2012, 79, 1529-1531.	0.8	5
11	Idiopathic chronic pancreatitis in India: phenotypic characterisation and strong genetic susceptibility due to SPINK1 and CFTR gene mutations. Gut, 2010, 59, 800-807.	12.1	100
12	Screening of families with autosomal recessive non-syndromic hearing impairment (ARNSHI) for mutations in GJB2 gene: Indian scenario. American Journal of Medical Genetics Part A, 2003, 120A, 180-184.	2.4	55