

# Shivaram Shastri

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8010751/publications.pdf>

Version: 2024-02-01

12  
papers

273  
citations

1163117

8  
h-index

1281871

11  
g-index

13  
all docs

13  
docs citations

13  
times ranked

409  
citing authors

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