Rowmika Ravi

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1434562/publications.pdf

Version: 2024-02-01

1937685 1372567 10 105 4 10 citations h-index g-index papers 10 10 10 201 docs citations times ranked citing authors all docs

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Saliva microbiome in primary Sjögren's syndrome reveals distinct set of diseaseâ€associated microbes. Oral Diseases, 2020, 26, 295-301. | 3.0 | 39 |
| 2 | Pharmacogenetic landscape of <i>DPYD</i> variants in south Asian populations by integration of genome-scale data. Pharmacogenomics, 2018, 19, 227-241. | 1.3 | 25 |
| 3 | SAGE: a comprehensive resource of genetic variants integrating South Asian whole genomes and exomes. Database: the Journal of Biological Databases and Curation, 2018, 2018, 1-10. | 3.0 | 20 |
| 4 | <scp>E</scp> gyptian tale from <scp>I</scp> ndia: application of wholeâ€exome sequencing in diagnosis of atypical familial <scp>M</scp> editerranean fever. International Journal of Rheumatic Diseases, 2017, 20, 1770-1775. | 1.9 | 7 |
| 5 | Understanding the complexity of epimorphic regeneration in zebrafish caudal fin tissue: A transcriptomic and proteomic approach. Genomics, 2022, 114, 110300. | 2.9 | 5 |
| 6 | Case Report: Whole exome sequencing reveals a novel frameshift deletion mutation p.G2254fs in COL7A1 associated with autosomal recessive dystrophic epidermolysis bullosa. F1000Research, 2016, 5, 900. | 1.6 | 3 |
| 7 | Case Report: Whole exome sequencing reveals a novel frameshift deletion mutation p.G2254fs in COL7A1 associated with autosomal recessive dystrophic epidermolysis bullosa. F1000Research, 2016, 5, 900. | 1.6 | 2 |
| 8 | Case Report:ÂWhole exome sequencing identifies variation c.2308G>A p.E770K in RAG1 associated with B- T- NK+ severe combined immunodeficiency. F1000Research, 2016, 5, 2532. | 1.6 | 2 |
| 9 | Case Report: Whole exome sequencing identifies variation c.2308G>A p.E770K in RAG1 associated with B-T-NK+ severe combined immunodeficiency. F1000Research, 2016, 5, 2532. | 1.6 | 1 |
| 10 | Metabolite Signature in the Carriers of Pathogenic Genetic Variants for Cardiomyopathy: A Population-Based METSIM Study. Metabolites, 2022, 12, 437. | 2.9 | 1 |