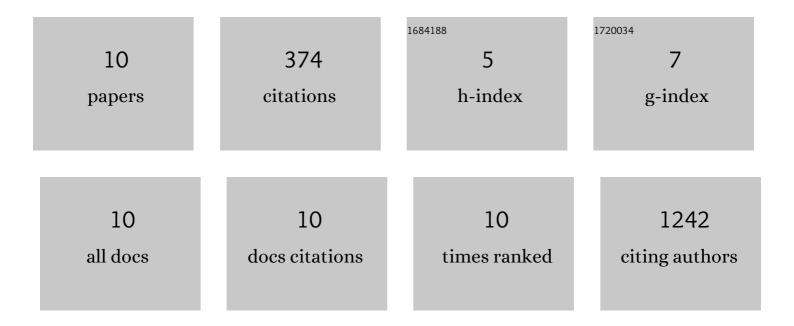
Resham Ejaz

List of Publications by Year in descending order

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RECHAM FIAZ

| # | Article | IF | CITATIONS |
|----|---|-----|-----------|
| 1 | Impact of Mobility Device Use on Quality of Life in Children With Friedreich Ataxia. Journal of Child Neurology, 2018, 33, 397-404. | 1.4 | 5 |
| 2 | Primary mediastinal paraganglioma associated with a familial variant in the succinate dehydrogenase B subunit gene. Journal of Surgical Oncology, 2018, 117, 160-162. | 1.7 | 6 |
| 3 | Riboflavin transporter deficiency mimicking mitochondrial myopathy caused by complex II deficiency. American Journal of Medical Genetics, Part A, 2018, 176, 399-403. | 1.2 | 18 |
| 4 | Stable transmission of an unbalanced chromosome 21 derived from chromoanasynthesis in a patient with a SYNGAP1 likely pathogenic variant. Molecular Cytogenetics, 2018, 11, 50. | 0.9 | 11 |
| 5 | Urgent Management of Bleeding in Immune Thrombocytopenia: Towards a Standardized Protocol in the Emergency Department. Blood, 2018, 132, 3517-3517. | 1.4 | Ο |
| 6 | De novo pathogenic variant in TUBB2A presenting with arthrogryposis multiplex congenita, brain abnormalities, and severe developmental delay. , 2017, 173, 2725-2730. | | 15 |
| 7 | Whole-genome sequencing expands diagnostic utility and improves clinical management in paediatric medicine. Npj Genomic Medicine, 2016, 1, . | 3.8 | 295 |
| 8 | Child Neurology: Diencephalic syndrome–like presentation of a cervicomedullary brainstem tumor. Neurology, 2016, 87, e248-e251. | 1.1 | 5 |
| 9 | Lateral meningocele (Lehman) syndrome: A child with a novel <i>NOTCH3</i> mutation. American Journal of Medical Genetics, Part A, 2016, 170, 1070-1075. | 1.2 | 19 |
| 10 | MG-119â€The evolving features of nicolaides-baraitser syndrome – a case report of a twenty-years follow-up. Journal of Medical Genetics, 2015, 52, A5.1-A5. | 3.2 | 0 |