

# Farrah Rajabi

## List of Publications by Year in descending order

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Version: 2024-02-01

17  
papers

88  
citations

1684188  
5  
h-index

1474206  
9  
g-index

17  
all docs

17  
docs citations

17  
times ranked

168  
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel variant in the <i>TSPAN12</i> gene-presenting as unilateral myopia, pediatric cataract, and heterochromia in a patient with familial exudative vitreoretinopathy. <i>European Journal of Ophthalmology</i> , 2022, 32, NP6-NP9.	1.3	1
2	Semaphorin-Plexin Signaling: From Axonal Guidance to a New X-Linked Intellectual Disability Syndrome. <i>Pediatric Neurology</i> , 2022, 126, 65-73.	2.1	8
3	Pontocerebellar hypoplasia due to bi-allelic variants in MINPP1. <i>European Journal of Human Genetics</i> , 2021, 29, 411-421.	2.8	13
4	Genetic Study in a Cohort of Children With ROHHAD Syndrome. <i>Journal of the Endocrine Society</i> , 2021, 5, A503-A504.	0.2	2
5	Genotype-phenotype correlations and novel molecular insights into the DHX30-associated neurodevelopmental disorders. <i>Genome Medicine</i> , 2021, 13, 90.	8.2	16
6	Graves™ disease in a five-month-old boy with an unusual treatment course. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 401-406.	0.9	3
7	OR33-07 ARNT2: A Potential Novel Candidate Gene for Monogenic Obesity in Humans. <i>Journal of the Endocrine Society</i> , 2020, 4, .	0.2	0
8	Imprinted genes in clinical exome sequencing: Review of 538 cases and exploration of mouse-human conservation in the identification of novel human disease loci. <i>European Journal of Medical Genetics</i> , 2020, 63, 103903.	1.3	4
9	Phenylalanine hydroxylase genotype-phenotype associations in the United States: A single center study. <i>Molecular Genetics and Metabolism</i> , 2019, 128, 415-421.	1.1	7
10	Acute Pancreatitis in a Patient with Maple Syrup Urine Disease: A Management Paradox. <i>Journal of Pediatrics</i> , 2018, 198, 313-316.	1.8	0
11	Gain-of-function variants in the <i>ODC1</i> gene cause a syndromic neurodevelopmental disorder associated with macrocephaly, alopecia, dysmorphic features, and neuroimaging abnormalities. <i>American Journal of Medical Genetics, Part A</i> , 2018, 176, 2554-2560.	1.2	26
12	Liver Failure as the Presentation of Ornithine Transcarbamylase Deficiency in a 13-Month-Old Female. <i>JIMD Reports</i> , 2017, 40, 17-22.	1.5	4
13	Confounding factors in identification of disease-resilient individuals. <i>Nature Biotechnology</i> , 2016, 34, 1103-1104.	17.5	0
14	Hyperphenylalaninemia and the genomic revolution. <i>Molecular Genetics and Metabolism</i> , 2015, 114, 380-381.	1.1	1
15	Expansion and Implications of Newborn Screening. <i>Current Genetic Medicine Reports</i> , 2015, 3, 110-117.	1.9	1
16	New Innovations: Therapies for Genetic Conditions. <i>Current Genetic Medicine Reports</i> , 2014, 2, 113-123.	1.9	1
17	Bilateral consecutive choroidal neovascularization in Best vitelliform macular dystrophy. <i>Baylor University Medical Center Proceedings</i> , 0, , 1-3.	0.5	1