

Hagit Baris-Feldman

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9938391/publications.pdf>

Version: 2024-02-01

26
papers

5,050
citations

686830

13
h-index

580395

25
g-index

27
all docs

27
docs citations

27
times ranked

8787
citing authors

#	ARTICLE	IF	CITATIONS
1	CD55-deficiency in Jews of Bukharan descent is caused by the Cromer blood type Dr(a ⁺) variant. <i>Human Genetics</i> , 2023, 142, 683-690.	1.8	3
2	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. <i>Science Immunology</i> , 2023, 8, .	5.6	35
3	Diagnostic criteria for constitutional mismatch repair deficiency (CMMRD): recommendations from the international consensus working group. <i>Journal of Medical Genetics</i> , 2022, 59, 318-327.	1.5	57
4	A recurrent pathogenic BRCA2 exon 5 ^{â€} 11 duplication in the Christian Arab population in Israel. <i>Familial Cancer</i> , 2022, 21, 289-294.	0.9	2
5	A global effort to dissect the human genetic basis of resistance to SARS-CoV-2 infection. <i>Nature Immunology</i> , 2022, 23, 159-164.	7.0	41
6	Human genetic and immunological determinants of critical COVID-19 pneumonia. <i>Nature</i> , 2022, 603, 587-598.	13.7	216
7	Bi-allelic variants in neuronal cell adhesion molecule cause a neurodevelopmental disorder characterized by developmental delay, hypotonia, neuropathy/spasticity. <i>American Journal of Human Genetics</i> , 2022, 109, 518-532.	2.6	8
8	Non-immune hydrops fetalis caused by <i>PIEZO1</i> compound heterozygous deletions detected only by exome sequencing. <i>Prenatal Diagnosis</i> , 2022, , .	1.1	3
9	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2200413119.	3.3	110
10	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. <i>Journal of Experimental Medicine</i> , 2022, 219, .	4.2	59
11	Heterozygous loss of <i>WBP11</i> function causes multiple congenital defects in humans and mice. <i>Human Molecular Genetics</i> , 2021, 29, 3662-3678.	1.4	14
12	Eculizumab-Responsive Adult Onset Protein Losing Enteropathy, Caused by Germline CD55-Deficiency and Complicated by Aggressive Angiosarcoma. <i>Journal of Clinical Immunology</i> , 2021, 41, 477-481.	2.0	6
13	Pathogenic variants in <i>SMARCA5</i> , a chromatin remodeler, cause a range of syndromic neurodevelopmental features. <i>Science Advances</i> , 2021, 7, .	4.7	17
14	A novel truncating variant in the <i>FGD1</i> gene associated with Aarskog-Scott syndrome in a family previously diagnosed with Tel Hashomer camptodactyly. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 3161-3166.	0.7	0
15	Non-immune Hemolysis in Gaucher Disease and Review of the Literature. <i>Rambam Maimonides Medical Journal</i> , 2021, 12, e0025.	0.4	4
16	Clinical outcomes after 4.5%years of eliglustat therapy for <i>Gaucher</i> disease type 1: Phase 3 <i>ENGAGE</i> trial final results. <i>American Journal of Hematology</i> , 2021, 96, 1156-1165.	2.0	22
17	Experts TM views on COVID-19 vaccination and the impact of the pandemic on patients with Gaucher disease. <i>British Journal of Haematology</i> , 2021, 195, e135-e137.	1.2	3
18	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. <i>Science Immunology</i> , 2021, 6, .	5.6	357

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19	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. <i>Science Immunology</i> , 2021, 6, .	5.6	267
20	PI(3,4)P2-mediated cytokinetic abscission prevents early senescence and cataract formation. <i>Science</i> , 2021, 374, eabk0410.	6.0	37
21	Spectrum of genes for inherited hearing loss in the Israeli Jewish population, including the novel human deafness gene <i>ATOH1</i> . <i>Clinical Genetics</i> , 2020, 98, 353-364.	1.0	15
22	The National Autism Database of Israel: a Resource for Studying Autism Risk Factors, Biomarkers, Outcome Measures, and Treatment Efficacy. <i>Journal of Molecular Neuroscience</i> , 2020, 70, 1303-1312.	1.1	22
23	Inborn errors of type I IFN immunity in patients with life-threatening COVID-19. <i>Science</i> , 2020, 370, .	6.0	1,749
24	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. <i>Science</i> , 2020, 370, .	6.0	1,983
25	Prenatal clubfoot increases the risk for clinically significant chromosomal microarray results – Analysis of 269 singleton pregnancies. <i>Early Human Development</i> , 2020, 145, 105047.	0.8	10
26	Diagnostic yield of multigene panel testing in an Israeli cohort: enrichment of low-penetrance variants. <i>Breast Cancer Research and Treatment</i> , 2020, 181, 445-453.	1.1	7