

# Ori Scott

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4057233/publications.pdf>

Version: 2024-02-01

25  
papers

240  
citations

1039406

9  
h-index

996533

15  
g-index

25  
all docs

25  
docs citations

25  
times ranked

431  
citing authors

#	ARTICLE	IF	CITATIONS
1	NF- $\kappa$ B pathway and the Goldilocks principle: Lessons from human disorders of immunity and inflammation. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 1688-1701.	1.5	35
2	Anti-N-Methyl-D-Aspartate (NMDA) Receptor Encephalitis. <i>Journal of Child Neurology</i> , 2014, 29, 691-694.	0.7	33
3	Clinical clues for autoimmunity and neuroinflammation in patients with autistic regression. <i>Developmental Medicine and Child Neurology</i> , 2017, 59, 947-951.	1.1	30
4	Long-Term Outcome of Adenosine Deaminase-Deficient Patients—a Single-Center Experience. <i>Journal of Clinical Immunology</i> , 2017, 37, 582-591.	2.0	26
5	Targeted genome editing <i>in vivo</i> corrects a Dmd duplication restoring wild-type dystrophin expression. <i>EMBO Molecular Medicine</i> , 2021, 13, e13228.	3.3	18
6	STAT1 gain-of-function heterozygous cell models reveal diverse interferon-signature gene transcriptional responses. <i>Npj Genomic Medicine</i> , 2021, 6, 34.	1.7	13
7	Fetal Exposure to Alcohol, Developmental Brain Anomaly, and Vitamin A Deficiency: A Case Report. <i>Journal of Child Neurology</i> , 2011, 26, 231-234.	0.7	11
8	The Safety of Cruciferous Plants in Humans: A Systematic Review. <i>Journal of Biomedicine and Biotechnology</i> , 2012, 2012, 1-28.	3.0	11
9	Global Developmental Delay, Progressive Relapsing-Remitting Parkinsonism, and Spinal Syring in a Child With SOX6 Mutation. <i>Journal of Child Neurology</i> , 2014, 29, NP164-NP167.	0.7	11
10	Using the Test of Variables of Attention to Determine the Effectiveness of Modafinil in Children With Attention-Deficit Hyperactivity Disorder (ADHD): A Prospective Methylphenidate-Controlled Trial. <i>Journal of Child Neurology</i> , 2012, 27, 1547-1552.	0.7	10
11	Medical Student Perceptions of On-Call Modalities: A Focus Group Study. <i>Teaching and Learning in Medicine</i> , 2019, 31, 34-43.	1.3	7
12	DNA-Binding domain mutations confer severe outcome at an early age among STAT1 gain-of-function patients. <i>Pediatric Allergy and Immunology</i> , 2022, 33, .	1.1	7
13	Social Communication Features in Children following Moderate to Severe Acquired Brain Injury. <i>Journal of Child Neurology</i> , 2015, 30, 588-594.	0.7	6
14	Case Report: Eosinophilic Esophagitis in a Patient With a Novel STAT1 Gain-of-Function Pathogenic Variant. <i>Frontiers in Immunology</i> , 2022, 13, 801832.	2.2	5
15	A novel splice site variant in FOXN1 in a patient with abnormal newborn screening for severe combined immunodeficiency and congenital lymphopenia. <i>LymphoSign Journal</i> , 2021, 8, 1-4.	0.1	4
16	Basal Ganglia Injury With Extrapyramidal Presentation. <i>Journal of Child Neurology</i> , 2013, 28, 1489-1492.	0.7	3
17	Homozygous duplication identified by whole genome sequencing causes LRBA deficiency. <i>Npj Genomic Medicine</i> , 2021, 6, 96.	1.7	3
18	The Effect of Caffeine on the Neuropathological and Neurobehavioral Outcome in the Newborn Rat. <i>Journal of Caffeine and Adenosine Research</i> , 2018, 8, 143-152.	0.8	2

#	ARTICLE	IF	CITATIONS
19	Chronic mucocutaneous Candidiasis caused by a novel <i>STAT1</i> mutation: a report of 4 patients. <i>LymphoSign Journal</i> , 2021, 8, 64-67.	0.1	2
20	Three Cases of Cerebellar Hypoplasia and Vitamin A Deficiency. <i>Journal of Child Neurology</i> , 2011, 26, 1311-1315.	0.7	1
21	Decreased Levels of Nasal Nitric Oxide in Children With Midline Neuroanatomical Anomalies: A Possible Connection Between Ciliary Dysfunction and Isolated Nervous System Defects. <i>Pediatric Neurology</i> , 2015, 53, 324-329.	1.0	1
22	An atypical presentation of ataxia telangiectasia in a school-aged boy secondary to an intronic mutation. <i>LymphoSign Journal</i> , 2020, 7, 57-60.	0.1	1
23	A Toddler With Prolonged Fever and Intermittent Cough. <i>Global Pediatric Health</i> , 2019, 6, 2333794X1882299.	0.3	0
24	Hematopoietic stem cell transplantations for primary immune deficiencies associated with <i>NF1</i> mutations: A review of the literature. <i>LymphoSign Journal</i> , 0, , .	0.1	0
25	Novel mutation in <i>PIK3CD</i> affecting the Ras-binding domain. <i>LymphoSign Journal</i> , 2022, 9, 11-16.	0.1	0