

Jane Peake

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1918234/publications.pdf>

Version: 2024-02-01

19
papers

2,860
citations

623574

14
h-index

713332

21
g-index

24
all docs

24
docs citations

24
times ranked

4803
citing authors

#	ARTICLE	IF	CITATIONS
1	Hyper-IgE Syndrome due to an Elusive Novel Intronic Homozygous Variant in DOCK8. <i>Journal of Clinical Immunology</i> , 2022, 42, 119-129.	2.0	4
2	A loss-of-function <i>IFNAR1</i> allele in Polynesia underlies severe viral diseases in homozygotes. <i>Journal of Experimental Medicine</i> , 2022, 219, .	4.2	28
3	Effects of an Amino Acid-Based Formula Supplemented with Two Human Milk Oligosaccharides on Growth, Tolerability, Safety, and Gut Microbiome in Infants with Cow's Milk Protein Allergy. <i>Nutrients</i> , 2022, 14, 2297.	1.7	12
4	Somatic reversion of pathogenic DOCK8 variants alters lymphocyte differentiation and function to effectively cure DOCK8 deficiency. <i>Journal of Clinical Investigation</i> , 2021, 131, .	3.9	18
5	Human <i>STAT3</i> variants underlie autosomal dominant hyper-IgE syndrome by negative dominance. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	30
6	Making a diagnosis of periodic fever syndrome: Experience from a single tertiary centre. <i>Journal of Paediatrics and Child Health</i> , 2021, , .	0.4	0
7	Diagnosis and management of severe combined immunodeficiency in Australia and New Zealand. <i>Journal of Paediatrics and Child Health</i> , 2020, 56, 1508-1513.	0.4	4
8	Heterozygous <i>FOXN1</i> Variants Cause Low TRECs and Severe T Cell Lymphopenia, Revealing a Crucial Role of <i>FOXN1</i> in Supporting Early Thymopoiesis. <i>American Journal of Human Genetics</i> , 2019, 105, 549-561.	2.6	52
9	Hematopoietic stem cell transplant effectively rescues lymphocyte differentiation and function in DOCK8-deficient patients. <i>JCI Insight</i> , 2019, 4, .	2.3	23
10	Therapeutic plasma exchange normalizes insulin-mediated response in a child with type 1 diabetes and insulin autoimmune syndrome. <i>Pediatric Diabetes</i> , 2018, 19, 171-179.	1.2	8
11	DOCK8 Drives Src-Dependent NK Cell Effector Function. <i>Journal of Immunology</i> , 2017, 199, 2118-2127.	0.4	18
12	Dedicator of cytokinesis 8-deficient CD4 + T _H 2 cells are biased to a T _H 2 effector fate at the expense of T _H 1 and T _H 17 cells. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 933-949.	1.5	69
13	Exome Sequencing Reveals Primary Immunodeficiencies in Children with Community-Acquired <i>Pseudomonas aeruginosa</i> Sepsis. <i>Frontiers in Immunology</i> , 2016, 7, 357.	2.2	21
14	STAT3 is a critical cell-intrinsic regulator of human unconventional T cell numbers and function. <i>Journal of Experimental Medicine</i> , 2015, 212, 855-864.	4.2	70
15	Compound heterozygous mutations in the noncoding <i>RNU4ATAC</i> cause Roifman Syndrome by disrupting minor intron splicing. <i>Nature Communications</i> , 2015, 6, 8718.	5.8	104
16	Functional STAT3 deficiency compromises the generation of human T follicular helper cells. <i>Blood</i> , 2012, 119, 3997-4008.	0.6	267
17	IL-21 is the primary common γ chain-binding cytokine required for human B-cell differentiation in vivo. <i>Blood</i> , 2011, 118, 6824-6835.	0.6	132
18	B cell-intrinsic signaling through IL-21 receptor and STAT3 is required for establishing long-lived antibody responses in humans. <i>Journal of Experimental Medicine</i> , 2010, 207, 155-171.	4.2	346

#	ARTICLE	IF	CITATIONS
19	X-linked neonatal diabetes mellitus, enteropathy and endocrinopathy syndrome is the human equivalent of mouse scurfy. <i>Nature Genetics</i> , 2001, 27, 18-20.	9.4	1,648