Francesca Pala

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9354068/publications.pdf

Version: 2024-02-01

21 888 15
papers citations h-index

27 27 27 1376
all docs docs citations times ranked citing authors

23

g-index

#	Article	IF	CITATIONS
1	Immunopathological signatures in multisystem inflammatory syndrome in children and pediatric COVID-19. Nature Medicine, 2022, 28, 1050-1062.	15.2	144
2	Autoimmunity in Wiskott–Aldrich Syndrome: An Unsolved Enigma. Frontiers in Immunology, 2012, 3, 209.	2.2	110
3	Distinct metabolic states govern skeletal muscle stem cell fates during prenatal and postnatal myogenesis. Journal of Cell Science, 2018, 131, .	1.2	109
4	Wiskott–Aldrich Syndrome protein deficiency perturbs the homeostasis of B-cell compartment in humans. Journal of Autoimmunity, 2014, 50, 42-50.	3.0	72
5	Artificial thymic organoids represent a reliable tool to study T-cell differentiation in patients with severe T-cell lymphopenia. Blood Advances, 2020, 4, 2611-2616.	2.5	65
6	An Integrated Epigenomic and Transcriptomic Map of Mouse and Human $\hat{l}\pm\hat{l}^2$ T Cell Development. Immunity, 2020, 53, 1182-1201.e8.	6.6	49
7	Lentiviral-mediated gene therapy restores B cell tolerance in Wiskott-Aldrich syndrome patients. Journal of Clinical Investigation, 2015, 125, 3941-3951.	3.9	43
8	B-cell reconstitution after lentiviral vector–mediated gene therapy in patients with Wiskott-Aldrich syndrome. Journal of Allergy and Clinical Immunology, 2015, 136, 692-702.e2.	1.5	41
9	Cysteine and hydrophobic residues in CDR3 serve as distinct T-cell self-reactivity indices. Journal of Allergy and Clinical Immunology, 2019, 144, 333-336.	1.5	31
10	Comparison of multiple transcriptomes exposes unified and divergent features of quiescent and activated skeletal muscle stem cells. Skeletal Muscle, 2017, 7, 28.	1.9	29
11	<i>SASH3</i> variants cause a novel form of X-linked combined immunodeficiency with immune dysregulation. Blood, 2021, 138, 1019-1033.	0.6	28
12	Gene Editing Rescues In vitro T Cell Development of RAG2-Deficient Induced Pluripotent Stem Cells in an Artificial Thymic Organoid System. Journal of Clinical Immunology, 2021, 41, 852-862.	2.0	27
13	RAG deficiencies: Recent advances in disease pathogenesis and novel therapeutic approaches. European Journal of Immunology, 2021, 51, 1028-1038.	1.6	22
14	F-BAR domain only protein 1 (FCHO1) deficiency is a novel cause of combined immune deficiency in human subjects. Journal of Allergy and Clinical Immunology, 2019, 143, 2317-2321.e12.	1.5	21
15	A distinct cardiopharyngeal mesoderm genetic hierarchy establishes antero-posterior patterning of esophagus striated muscle. ELife, 2019, 8, .	2.8	20
16	Isolation of Muscle Stem Cells from Mouse Skeletal Muscle. Methods in Molecular Biology, 2017, 1556, 23-39.	0.4	19
17	Gut Microbiota–Host Interactions in Inborn Errors of Immunity. International Journal of Molecular Sciences, 2021, 22, 1416.	1.8	18
18	In Vivo Chronic Stimulation Unveils Autoreactive Potential of Wiskott–Aldrich Syndrome Protein-Deficient B Cells. Frontiers in Immunology, 2017, 8, 490.	2.2	10

#	Article	IF	CITATIONS
19	POLD1 Deficiency Reveals a Role for POLD1 in DNA Repair and T and B Cell Development. Journal of Clinical Immunology, 2021, 41, 270-273.	2.0	10
20	Skewed TCR Alpha, but not Beta, Gene Rearrangements and Lymphoma Associated with a Pathogenic TRAC Variant. Journal of Clinical Immunology, 2021, 41, 1395-1399.	2.0	4
21	Phosphate Transporter Profiles in Murine and Human Thymi Identify Thymocytes at Distinct Stages of Differentiation. Frontiers in Immunology, 2020, 11, 1562.	2.2	3