## Francesca Pala

List of Publications by Year in descending order

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FRANCESCA PALA

#	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	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
3	Gut Microbiota–Host Interactions in Inborn Errors of Immunity. International Journal of Molecular Sciences, 2021, 22, 1416.	1.8	18
4	RAG deficiencies: Recent advances in disease pathogenesis and novel therapeutic approaches. European Journal of Immunology, 2021, 51, 1028-1038.	1.6	22
5	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
6	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
7	<i>SASH3</i> variants cause a novel form of X-linked combined immunodeficiency with immune dysregulation. Blood, 2021, 138, 1019-1033.	0.6	28
8	An Integrated Epigenomic and Transcriptomic Map of Mouse and Human αβ T Cell Development. Immunity, 2020, 53, 1182-1201.e8.	6.6	49
9	Phosphate Transporter Profiles in Murine and Human Thymi Identify Thymocytes at Distinct Stages of Differentiation. Frontiers in Immunology, 2020, 11, 1562.	2.2	3
10	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
11	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
12	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
13	A distinct cardiopharyngeal mesoderm genetic hierarchy establishes antero-posterior patterning of esophagus striated muscle. ELife, 2019, 8, .	2.8	20
14	Distinct metabolic states govern skeletal muscle stem cell fates during prenatal and postnatal myogenesis. Journal of Cell Science, 2018, 131, .	1.2	109
15	Isolation of Muscle Stem Cells from Mouse Skeletal Muscle. Methods in Molecular Biology, 2017, 1556, 23-39.	0.4	19
16	In Vivo Chronic Stimulation Unveils Autoreactive Potential of Wiskott–Aldrich Syndrome Protein-Deficient B Cells. Frontiers in Immunology, 2017, 8, 490.	2.2	10
17	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
18	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

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19	Lentiviral-mediated gene therapy restores B cell tolerance in Wiskott-Aldrich syndrome patients. Journal of Clinical Investigation, 2015, 125, 3941-3951.	3.9	43
20	Wiskott–Aldrich Syndrome protein deficiency perturbs the homeostasis of B-cell compartment in humans. Journal of Autoimmunity, 2014, 50, 42-50.	3.0	72
21	Autoimmunity in Wiskott–Aldrich Syndrome: An Unsolved Enigma. Frontiers in Immunology, 2012, 3, 209.	2.2	110