JoÃ£o Farela Neves

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9450838/publications.pdf

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30	603	933264 10	642610 23
papers	citations	h-index	g-index

32	32	32	1163
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Homozygous V377I mutation causing mevalonate kinase. BMJ Case Reports, 2022, 15, e249135.	0.2	0
2	Case Report: Wide Spectrum of Manifestations of Ligase IV Deficiency: Report of 3 Cases. Frontiers in Immunology, 2022, 13, 869728.	2.2	1
3	Haploidentical Î±/Î² T-cell and B-cell depleted stem cell transplantation in severe mevalonate kinase deficiency. Rheumatology, 2021, 60, 4850-4854.	0.9	6
4	Kingella kingae. Pediatric Infectious Disease Journal, 2021, 40, e247-e249.	1.1	1
5	Case Report: Primary Immunodeficiencies, Massive EBV+ T-Cell Lympoproliferation Leading to the Diagnosis of ICF2 Syndrome. Frontiers in Immunology, 2021, 12, 654167.	2.2	3
6	A Novel TRAF3IP2 Mutation Causing Chronic Mucocutaneous Candidiasis. Journal of Clinical Immunology, 2021, 41, 1376-1379.	2.0	11
7	Variants in STXBP3 are Associated with Very Early Onset Inflammatory Bowel Disease, Bilateral Sensorineural Hearing Loss and Immune Dysregulation. Journal of Crohn's and Colitis, 2021, 15, 1908-1919.	0.6	7
8	SARS-CoV-2â€“related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. Journal of Experimental Medicine, 2021, 218, .	4.2	100
9	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. Journal of Allergy and Clinical Immunology, 2021, 148, 1332-1341.e5.	1.5	75
10	Congenital nephrotic syndrome in IL7RÎ±-SCID: A rare feature of maternofetal graft-versus-host disease. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 4151-4153.e1.	2.0	2
11	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. Journal of Allergy and Clinical Immunology, 2020, 146, 901-911.	1.5	78
12	Fatal Meningitis in Patient with X-Linked Chronic Granulomatous Disease Caused by Virulent Granulibacter bethesdensis. Emerging Infectious Diseases, 2019, 25, 976-979.	2.0	2
13	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. World Allergy Organization Journal, 2019, 12, 100018.	1.6	83
14	Invasive Meningococcal Disease Unraveling a Novel Mutation in the C5 Gene in a Portuguese Family. Pediatric Infectious Disease Journal, 2019, 38, 416-418.	1.1	2
15	Fatal Central Nervous System Lymphocytic Vasculitis after Treatment for Burkitt Lymphoma in a Patient with a SH2D1A Mutation. Pediatric Infectious Disease Journal, 2019, 38, e29-e31.	1.1	4
16	Plerixafor for the Treatment of WHIM Syndrome. New England Journal of Medicine, 2019, 380, 163-170.	13.9	74
17	Fatal Meningitis in Patient with X-Linked Chronic Granulomatous Disease Caused by Virulent Granulibacter bethesdensis. Emerging Infectious Diseases, 2019, 25, 976-979.	2.0	0
18	25 MUTATIONS IN STXBP3 CONTRIBUTE TO VERY EARLY ONSET OF IBD, IMMUNODEFICIENCY AND HEARING LOSS. Inflammatory Bowel Diseases, 2018, 24, S28-S28.	0.9	0

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19	25 MUTATIONS IN STXBP3 CONTRIBUTE TO VERY EARLY ONSET OF IBD, IMMUNODEFICIENCY AND HEARING LOSS. Gastroenterology, 2018, 154, S40-S41.	0.6	0
20	Missense mutation of TTC7A mimicking tricho-hepato-enteric (SD/THE) syndrome in a patient with very-early onset inflammatory bowel disease. European Journal of Medical Genetics, 2018, 61, 185-188.	0.7	18
21	Severe Stevensâ€Johnson syndrome/toxic epidermal necrolysis overlap syndromeâ€”beyond skin involvement. Pediatric Dermatology, 2018, 35, e17-e19.	0.5	6
22	Novel PLCG2 Mutation in a Patient With APLAID and Cutis Laxa. Frontiers in Immunology, 2018, 9, 2863.	2.2	64
23	Sa2008 - Mutations in Stxbp3 Contribute to Very Early Onset of IBD Immunodeficieny and Hearing Loss. Gastroenterology, 2018, 154, S-445.	0.6	0
24	Methylotroph Infections and Chronic Granulomatous Disease. Emerging Infectious Diseases, 2016, 22, 404-409.	2.0	17
25	Primary immunodeficiency associated with chromosomal aberration â€“ an ESID survey. Orphanet Journal of Rare Diseases, 2016, 11, 110.	1.2	23
26	Hemophagocytic lymphohistiocytosis secondary to Falciparum malaria in a 5Âyear-old boy. Annals of Hematology, 2015, 94, 161-163.	0.8	3
27	Monoclonal IgG Kappa gammopathy previous to hematopoietic stem cell transplantation in an infant with severe combined immunodeficiency. Clinical Immunology, 2012, 145, 133-135.	1.4	2
28	Successful handling of autoimmunity in X-linked thrombocytopenia (XLT) using mycophenolate mofetil. Pediatric Blood and Cancer, 2012, 59, 961-961.	0.8	0
29	Successful Handling of Disseminated BCG Disease in a Child with Severe Combined Immunodeficiency. Case Reports in Medicine, 2011, 2011, 1-5.	0.3	17
30	ERITRODERMIA: PRIMEIRA MANIFESTAÃ‡ÃƒO DE DÃ‰FICE IMUNITÃRIO CONGÃ‰NITO. Journal of the Portuguese Society of Dermatology and Venereology, 2011, 69, 97-101.	0.0	0