S Y Patel

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5366449/publications.pdf

Version: 2024-02-01

32 papers	1,582 citations	393982 19 h-index	414034 32 g-index
33 all docs	33 docs citations	33 times ranked	3451 citing authors

#	Article	IF	Citations
1	Resolving the polygenic aetiology of a late onset combined immune deficiency caused by NFKB1 haploinsufficiency and modified by PIK3R1 and TNFRSF13B variants. Clinical Immunology, 2022, 234, 108910.	1.4	3
2	SARS-CoV-2 Vaccine Responses in Individuals with Antibody Deficiency: Findings from the COV-AD Study. Journal of Clinical Immunology, 2022, 42, 923-934.	2.0	37
3	Screening for Immunodeficiencies in Children With Invasive Pneumococcal Disease: Six-year Experience From a UK Childrenâ€™s Hospital. Pediatric Infectious Disease Journal, 2022, 41, 575-578.	1.1	3
4	Decreased ATM Function Causes Delayed DNA Repair and Apoptosis in Common Variable Immunodeficiency Disorders. Journal of Clinical Immunology, 2021, 41, 1315-1330.	2.0	6
5	A Novel, Heterozygous Three Base-Pair Deletion in CARD11 Results in B Cell Expansion with NF-ÎºB and T Cell Anergy Disease. Journal of Clinical Immunology, 2020, 40, 406-411.	2.0	10
6	Histology of Interstitial Lung Disease in Common Variable Immune Deficiency. Frontiers in Immunology, 2020, 11, 605187.	2.2	17
7	Impaired lymphocyte function and differentiation in CTPS1-deficient patients result from a hypomorphic homozygous mutation. JCI Insight, 2020, 5, .	2.3	29
8	Defining B-cell defects and correlation with complications in patients with common variable immune deficiency. Journal of Allergy and Clinical Immunology, 2019, 144, 654-655.	1.5	1
9	Interstitial lung disease in patients with common variable immunodeficiency disorders: several different pathologies?. Clinical and Experimental Immunology, 2019, 198, 212-223.	1.1	21
10	Sequencing of human genomes with nanopore technology. Nature Communications, 2019, 10, 1869.	5.8	140
11	Classical and Non-classical Presentations of Complement Factor I Deficiency: Two Contrasting Cases Diagnosed via Genetic and Genomic Methods. Frontiers in Immunology, 2019, 10, 1150.	2.2	21
12	Imaging of Bronchial Pathology in Antibody Deficiency: Data from the European Chest CT Group. Journal of Clinical Immunology, 2019, 39, 45-54.	2.0	32
13	Identification of CVID Patients With Defects in Immune Repertoire Formation or Specification. Frontiers in Immunology, 2018, 9, 2545.	2.2	38
14	The United Kingdom Primary Immune Deficiency (UKPID) registry 2012 to 2017. Clinical and Experimental Immunology, 2018, 192, 284-291.	1.1	57
15	Is It Safe to Switch From Intravenous Immunoglobulin to Subcutaneous Immunoglobulin in Patients With Common Variable Immunodeficiency and Autoimmune Thrombocytopenia?. Frontiers in Immunology, 2018, 9, 1656.	2.2	12
16	Clinical and laboratory features of seventy-eight UK patients with Goodâ€™s syndrome (thymoma and) Tj ETQq0 0 0 rgBT /Overlock 10	1.1	45
17	The role of genomics in common variable immunodeficiency disorders. Clinical and Experimental Immunology, 2017, 188, 326-332.	1.1	75
18	British Lung Foundation/United Kingdom Primary Immunodeficiency Network Consensus Statement on the Definition, Diagnosis, and Management of Granulomatous-Lymphocytic Interstitial Lung Disease in Common Variable Immunodeficiency Disorders. Journal of Allergy and Clinical Immunology: in Practice, 2017, 5, 938-945.	2.0	138

#	Article	IF	Citations
19	The primary immunodeficiency disorders. Medicine, 2017, 45, 597-604.	0.2	5
20	Prognosis of Good syndrome: mortality and morbidity of thymoma associated immunodeficiency in perspective. Clinical Immunology, 2016, 171, 12-17.	1.4	55
21	Chronic mucocutaneous candidiasis: characterization of a family with STAT-1 gain-of-function and development of an ex-vivo assay for Th17 deficiency of diagnostic utility. Clinical and Experimental Immunology, 2016, 184, 216-227.	1.1	25
22	Variable phenotype and discrete alterations of immune phenotypes in CTP synthase 1 deficiency: Report of 2 siblings. Journal of Allergy and Clinical Immunology, 2016, 138, 1722-1725.e6.	1.5	18
23	Clinical and laboratory correlates of lung disease and cancer in adults with idiopathic hypogammaglobulinaemia. Clinical and Experimental Immunology, 2016, 184, 73-82.	1.1	24
24	Hypomorphic function and somatic reversion of DOCK8 cause combined immunodeficiency without hyper-IgE. Clinical Immunology, 2016, 163, 17-21.	1.4	18
25	Identification of a Novel Mutation in MAGT1 and Progressive Multifocal Leucoencephalopathy in a 58-Year-Old Man with XMEN Disease. Journal of Clinical Immunology, 2015, 35, 112-118.	2.0	52
26	Key stages of bone marrow B-cell maturation are defective in patients with common variable immunodeficiency disorders. Journal of Allergy and Clinical Immunology, 2015, 136, 487-490.e2.	1.5	20
27	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. Nature Genetics, 2015, 47, 717-726.	9.4	310
28	Application of whole genome and RNA sequencing to investigate the genomic landscape of common variable immunodeficiency disorders. Clinical Immunology, 2015, 160, 301-314.	1.4	100
29	A mutation in X-linked inhibitor of apoptosis (G466X) leads to memory inflation of Epsteinâ€“Barr virus-specific T cells. Clinical and Experimental Immunology, 2014, 178, 470-482.	1.1	15
30	The United Kingdom Primary Immune Deficiency (UKPID) Registry: report of the first 4 years' activity 2008â€“2012. Clinical and Experimental Immunology, 2013, 175, 68-78.	1.1	85
31	Confirmation and improvement of criteria for clinical phenotyping in common variable immunodeficiency disorders in replicate cohorts. Journal of Allergy and Clinical Immunology, 2012, 130, 1197-1198.e9.	1.5	129
32	Lymphoid Proliferations of Indeterminate Malignant Potential arising in Adults with Common Variable Immunodeficiency Disorders: Unusual Case Studies and Immunohistological Review in the Light of Possible Causative Events. Journal of Clinical Immunology, 2011, 31, 784-791.	2.0	40