Xiaodong Zhao

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8164848/publications.pdf

Version: 2024-02-01

18	276	1307594 7	996975 15
papers	citations	h-index	g-index

22	22	22	434
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Reference values for peripheral blood lymphocyte subsets of healthy children in China. Journal of Allergy and Clinical Immunology, 2018, 142, 970-973.e8.	2.9	93
2	T-cell receptor diversity is selectively skewed in T-cell populations of patients with Wiskott-Aldrich syndrome. Journal of Allergy and Clinical Immunology, 2015, 135, 209-216.e8.	2.9	39
3	A nanoscale Co3O4â€“WO3 pâ€“n junction sensor with enhanced acetone responsivity. Journal of Materials Science: Materials in Electronics, 2015, 26, 8217-8223.	2.2	30
4	Molecular and Phenotypic Characterization of Nine Patients with STAT1 GOF Mutations in China. Journal of Clinical Immunology, 2020, 40, 82-95.	3.8	30
5	Novel biallelic TRNT1 mutations lead to atypical SIFD and multiple immune defects. Genes and Diseases, 2020, 7, 128-137.	3.4	16
6	Novel DOCK8 gene mutations lead to absence of protein expression in patients with hyper-IgE syndrome. Immunologic Research, 2016, 64, 260-271.	2.9	11
7	Autosomal dominant hyper IgE syndrome from a single centre in Chongqing, China (2009â€2018). Scandinavian Journal of Immunology, 2020, 91, e12885.	2.7	9
8	Activated Phosphoinositide 3-Kinase Î´ Syndrome: a Large Pediatric Cohort from a Single Center in China. Journal of Clinical Immunology, 2022, 42, 837-850.	3.8	9
9	Abatacept is effective in Chinese patients with LRBA and CTLA4 deficiency. Genes and Diseases, 2021, 8, 662-668.	3.4	8
10	Phenotypic characterization of patients with activated PI3KÎ´ syndrome 1 presenting with features of systemic lupus erythematosus. Genes and Diseases, 2021, 8, 907-917.	3.4	7
11	E1021K Homozygous Mutation in PIK3CD Leads to Activated PI3K-Delta Syndrome 1. Journal of Clinical Immunology, 2020, 40, 378-387.	3.8	7
12	Combined Immunodeficiency Caused by a Novel De Novo Gain-of-Function RAC2 Mutation. Journal of Clinical Immunology, 2022, 42, 1280-1292.	3.8	7
13	Novel CD81 Mutations in a Chinese Patient Led to IgA Nephropathy and Impaired BCR Signaling. Journal of Clinical Immunology, 2022, 42, 1672-1684.	3.8	4
14	Progranulin regulates the development and function of NKT2 cells through EZH2 and PLZF. Cell Death and Differentiation, 2022, 29, 1901-1912.	11.2	3
15	Design of building component library based on IFC and PLIB standard. , 2010, , .		2
16	Primary Immune Deficiencies â€“ A rapidly emerging area of basic and clinical research. Genes and Diseases, 2020, 7, 1-2.	3.4	1
17	Research on integrated digital city model and its application. , 2010, , .		0
18	Generation of human induced pluripotent stem cell line from peripheral blood mononuclear cells from an activated phosphoinositide 3-kinase Î´ syndrome patient. Stem Cell Research, 2022, 62, 102822.	0.7	0