

Ashish Marwaha

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7771649/publications.pdf>

Version: 2024-02-01

13
papers

409
citations

1478505

6
h-index

1199594

12
g-index

13
all docs

13
docs citations

13
times ranked

792
citing authors

#	ARTICLE	IF	CITATIONS
1	Cutting Edge: Increased IL-17 ⁺ Secreting T Cells in Children with New-Onset Type 1 Diabetes. <i>Journal of Immunology</i> , 2010, 185, 3814-3818.	0.8	190
2	TH17 Cells in Autoimmunity and Immunodeficiency: Protective or Pathogenic?. <i>Frontiers in Immunology</i> , 2012, 3, 129.	4.8	102
3	Targeting the IL-17/IFN- γ axis as a potential new clinical therapy for type 1 diabetes. <i>Clinical Immunology</i> , 2014, 154, 84-89.	3.2	40
4	Pre-diagnostic genotyping identifies T1D subjects with impaired Treg IL-2 signaling and an elevated proportion of FOXP3+IL-17+ cells. <i>Genes and Immunity</i> , 2017, 18, 15-21.	4.1	23
5	<sc>The point-of-care</sc> use of a facial phenotyping tool in the genetics clinic: Enhancing diagnosis and education with machine learning. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 1151-1158.	1.2	18
6	Congenital hypothyroidism, cardiac defects, and pancreatic agenesis in an infant with <sc>GATA6</sc> mutation. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 1496-1499.	1.2	8
7	Two cases of carbonic anhydrase <sc>VA</sc> deficiencyâ€”An ultrarare metabolic decompensation syndrome presenting with hyperammonemia, lactic acidosis, ketonuria, and good clinical outcome. <i>JIMD Reports</i> , 2021, 57, 9-14.	1.5	7
8	Genotype-phenotype data from a case series of patients with mosaic neurofibromatosis type 1. <i>British Journal of Dermatology</i> , 2018, 179, 1216-1217.	1.5	6
9	Progressive decline of T and B cell numbers and function in a patient with CDC42 deficiency. <i>Immunologic Research</i> , 2021, 69, 53-58.	2.9	5
10	The utility of <sc>DNA</sc> methylation signatures in directing genome sequencing workflow: Kabuki syndrome and <sc>CDK13</sc>-related disorder. <i>American Journal of Medical Genetics, Part A</i> , 2022, 188, 1368-1375.	1.2	5
11	Epidermal growth factor receptor deficiency: Expanding the phenotype beyond infancy. <i>Journal of Dermatology</i> , 2020, 47, 898-902.	1.2	3
12	Phase II multicentre, double-blind, randomised trial of ustekinumab in adolescents with new-onset type 1 diabetes (USTEK1D): trial protocol. <i>BMJ Open</i> , 2021, 11, e049595.	1.9	2
13	Expansion of the neurodevelopmental phenotypic spectrum of <sc><i>CKAP2L</i></sc>-related Filippi syndrome to include an adolescent male with normal intellect. <i>American Journal of Medical Genetics, Part A</i> , 2022, 188, 1928-1929.	1.2	0