

Esra Hazar Sayar

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7342507/publications.pdf>

Version: 2024-02-01

18
papers

542
citations

933447

10
h-index

996975

15
g-index

18
all docs

18
docs citations

18
times ranked

1649
citing authors

#	ARTICLE	IF	CITATIONS
1	Long-Term Experience of Subcutaneous Immunoglobulin Therapy in Pediatric Primary Immunodeficient Patients with Low and Normal Body Weight. <i>Journal of Clinical Immunology</i> , 2022, 42, 64-71.	3.8	2
2	Primary antibody deficiencies in Turkey: molecular and clinical aspects. <i>Immunologic Research</i> , 2021, , 1.	2.9	2
3	Borderless collaboration is needed for COVID-19â€™A disease that knows no borders. <i>Infection Control and Hospital Epidemiology</i> , 2020, 41, 1245-1246.	1.8	64
4	Mutational landscape of severe combined immunodeficiency patients from Turkey. <i>International Journal of Immunogenetics</i> , 2020, 47, 529-538.	1.8	14
5	The frequency of vitamin B12, iron, folic acid deficiency in the neonatal period and infancy and relationship with maternal levels. <i>Turk Pediatri Arsivi</i> , 2020, 55, 139-148.	0.9	7
6	International Efforts to Save Healthcare Personnel during COVID-19. <i>Acta Biomedica</i> , 2020, 91, e2020044.	0.3	12
7	Aeroallergen Sensitivity of Atopic Children in Alanya Region. <i>Selcuk Tip Dergisi</i> , 2020, 3, 226-231.	0.1	1
8	Inherited IFNAR1 deficiency in otherwise healthy patients with adverse reaction to measles and yellow fever live vaccines. <i>Journal of Experimental Medicine</i> , 2019, 216, 2057-2070.	8.5	127
9	Measles, mumps, rubella vaccination experience in infants with egg allergy. <i>Acta Medica Alanya</i> , 2019, 3, 283-286.	0.2	1
10	Type I IFNâ€™related NETosis in ataxia telangiectasia and Artemis deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 246-257.	2.9	47
11	Tuberculosis and impaired IL-23â€™dependent IFN-Î³ immunity in humans homozygous for a common <i>TYK2</i> missense variant. <i>Science Immunology</i> , 2018, 3, .	11.9	148
12	Hematopoietic stem cell transplantation from unrelated donors in children with <sc>DOCK</sc>8 deficiency. <i>Pediatric Transplantation</i> , 2017, 21, e13015.	1.0	12
13	Nijmegen Breakage Sendromlu Bir Olgunun Uzun SÃ¼reli izlemi. <i>Asim, Allerji, Immunoloji</i> , 2017, , .	0.0	0
14	The Ineffectiveness of Tacrolimus in an Infant With a Mutation in the IL-10 Receptor. <i>Journal of Clinical Gastroenterology</i> , 2016, 50, 352-353.	2.2	0
15	How effective are the 6 European Society of Immunodeficiency warning signs for primary immunodeficiency disease?. <i>Annals of Allergy, Asthma and Immunology</i> , 2016, 116, 151-155.e1.	1.0	16
16	Ocular Findings in Children With 22q11.2 Deletion Syndrome. <i>Journal of Pediatric Ophthalmology and Strabismus</i> , 2016, 53, 218-222.	0.7	13
17	<i>DCLRE1C</i> (ARTEMIS) mutations causing phenotypes ranging from atypical severe combined immunodeficiency to mere antibody deficiency. <i>Human Molecular Genetics</i> , 2015, 24, 7361-7372.	2.9	72
18	Langerhans cell histiocytosis in <sc>IPEX</sc> syndrome: Possible role for natural regulatory T cells?. <i>Pediatric Allergy and Immunology</i> , 2014, 25, 601-603.	2.6	4