

Baran Erman

List of Publications by Year in descending order

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Version: 2024-02-01

22
papers

464
citations

933264

10
h-index

794469

19
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22
all docs

22
docs citations

22
times ranked

1176
citing authors

#	ARTICLE	IF	CITATIONS
1	STK4 (MST1) deficiency in two siblings with autoimmune cytopenias: A novel mutation. <i>Clinical Immunology</i> , 2015, 161, 316-323.	1.4	73
2	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	0.6	64
3	Role of natural killer cells in lung cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2018, 144, 997-1003.	1.2	52
4	Necessity of Keratinized Tissues for Dental Implants: A Clinical, Immunological, and Radiographic Study. <i>Clinical Implant Dentistry and Related Research</i> , 2015, 17, 1-12.	1.6	50
5	Combined immunodeficiency with CD4 lymphopenia and sclerosing cholangitis caused by a novel loss-of-function mutation affecting IL21R. <i>Haematologica</i> , 2015, 100, e216-e219.	1.7	46
6	Statins and IL-1 β , IL-10, and MPO Levels in Gingival Crevicular Fluid: Preliminary Results. <i>Inflammation</i> , 2016, 39, 1547-1557.	1.7	31
7	Investigation of Genetic Defects in Severe Combined Immunodeficiency Patients from Turkey by Targeted Sequencing. <i>Scandinavian Journal of Immunology</i> , 2017, 85, 227-234.	1.3	30
8	Course of IL-2-inducible T-cell kinase deficiency in a family: lymphomatoid granulomatosis, lymphoma and allogeneic bone marrow transplantation in one sibling; and death in the other. <i>Bone Marrow Transplantation</i> , 2017, 52, 126-129.	1.3	21
9	Two siblings with PRKDC defect who presented with cutaneous granulomas and review of the literature. <i>Clinical Immunology</i> , 2018, 197, 1-5.	1.4	18
10	IgE and IgG4 binding to lentil epitopes in children with red and green lentil allergy. <i>Pediatric Allergy and Immunology</i> , 2020, 31, 158-166.	1.1	15
11	Expanding the Clinical and Immunological Phenotypes and Natural History of MALT1 Deficiency. <i>Journal of Clinical Immunology</i> , 2022, 42, 634-652.	2.0	12
12	C1q deficiency: identification of a novel missense mutation and treatment with fresh frozen plasma. <i>Clinical Rheumatology</i> , 2012, 31, 1123-1126.	1.0	10
13	Invasive <i>Saprochaete capitata</i> Infection in a Patient with Autosomal Recessive CARD9 Deficiency and a Review of the Literature. <i>Journal of Clinical Immunology</i> , 2020, 40, 466-474.	2.0	10
14	Diagnosis of Interstitial Lung Disease Caused by Possible Hypersensitivity Pneumonitis in a Child: Think CGD. <i>Journal of Clinical Immunology</i> , 2017, 37, 269-272.	2.0	9
15	A Novel Homozygous Mutation With Different Clinical Presentations in 2 IRAK-4 Deficient Siblings: First Case With Recurrent Salmonellosis and Non-Hodgkin Lymphoma. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2018, 28, 271-273.	0.6	6
16	Successful outcome with second hematopoietic stem cell transplantation in a patient with IL-10R deficiency. <i>Bone Marrow Transplantation</i> , 2016, 51, 615-616.	1.3	5
17	Genetic Screening of the Patients with Primary Immunodeficiency by Whole-Exome Sequencing. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2020, 33, 19-24.	0.3	5
18	Biallelic Form of a Known CD3E Mutation in a Patient with Severe Combined Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2020, 40, 539-542.	2.0	5

#	ARTICLE	IF	CITATIONS
19	Defective pneumococcal antibody response in patients with recurrent respiratory tract infections. Turkish Journal of Pediatrics, 2017, 59, 555-560.	0.3	1
20	Talasemi MajÅ¶r HastalarÄ±nda KÅ¶k Hücre Transplantasyonu SonrasÄ± Immun Yeniden YapÄ±lanmanÄ±n DeÄŸerlendirilmesi. Turkish Journal of Immunology, 2020, 8, 113-119.	0.1	1
21	Investigation of causative genetic defects in patients with primary immunodeficiency by next generation sequencing. Annals of Medical Research, 2021, 28, 1038.	0.0	0
22	Apoptosis in Autoimmune Lymphoproliferative Syndrome Suspected Patients with Clinical and Laboratory Findings. Turkish Journal of Immunology, 2013, 1, 5-12.	0.1	0