

Eren Erken

List of Publications by Year in descending order

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58
papers

1,060
citations

394421

19
h-index

434195

31
g-index

58
all docs

58
docs citations

58
times ranked

1840
citing authors

#	ARTICLE	IF	CITATIONS
1	Identification of susceptibility loci for Takayasu arteritis through a large multi-ancestral genome-wide association study. <i>American Journal of Human Genetics</i> , 2021, 108, 84-99.	6.2	26
2	MO947KILLER CELL IMMUNOGLOBULIN LIKE RECEPTOR (KIR) GENE DISTRIBUTION AND ALLOGRAFT KIDNEY FUNCTION. <i>Nephrology Dialysis Transplantation</i> , 2021, 36, .	0.7	0
3	Killer cell immunoglobulin-like receptor (KIR) gene distribution and allograft kidney function. <i>International Journal of Clinical Practice</i> , 2021, 75, e14790.	1.7	1
4	Predictive factors for workday loss in Behçet's syndrome: A multicenter study. <i>International Journal of Rheumatic Diseases</i> , 2020, 23, 240-246.	1.9	7
5	Moderation analysis exploring associations between age and mucocutaneous activity in Behçet's syndrome: A multicenter study from Turkey. <i>Journal of Dermatology</i> , 2020, 47, 1403-1410.	1.2	4
6	Importance of 14-3-3eta, anti-CarP, and anti-Sa in the diagnosis of seronegative rheumatoid arthritis. <i>Turkish Journal of Medical Sciences</i> , 2019, 49, 1498-1502.	0.9	9
7	The distribution of MEFV mutations in Turkish FMF patients: multicenter study representing results of Anatolia. <i>Turkish Journal of Medical Sciences</i> , 2019, 49, 472-477.	0.9	23
8	Exon 2: Is it the good police in familial mediterranean fever?. <i>European Journal of Rheumatology</i> , 2019, 6, 33-36.	0.6	10
9	INVESTIGATION OF THE FREQUENCY OF FC GAMMA RECEPTOR IIIA V/158/F GENE POLYMORPHISM AND COMPARISON OF CLINICAL AND LABORATORY FINDINGS IN RHEUMATOID ARTHRITIS (RA). <i>European Oral Research</i> , 2019, 81, 139-144.	0.9	0
10	Oral ulcer activity assessment with the composite index according to different treatment modalities in Behçet's syndrome: a multicentre study. <i>Clinical and Experimental Rheumatology</i> , 2019, 37 Suppl 121, 98-104.	0.8	3
11	Neuropathic pain: is it an underestimated symptom in systemic sclerosis?. <i>Clinical Rheumatology</i> , 2018, 37, 1845-1851.	2.2	5
12	Comparison of early versus late onset familial Mediterranean fever. <i>International Journal of Rheumatic Diseases</i> , 2018, 21, 880-884.	1.9	25
13	Cardiac disease in familial Mediterranean fever. <i>Rheumatology International</i> , 2018, 38, 51-58.	3.0	20
14	Mannose-Binding Lectin2 gene polymorphism in PANDAS patients. <i>Noropsikiyatri Arsivi</i> , 2018, 56, 99-105.	0.3	3
15	A Rare Side Effect of Intravesical Bacillus Calmette-Guérin Therapy: Reactive Arthritis. <i>Åeroonkoloji BÄ¼lteni</i> , 2018, 17, 33-35.	0.1	1
16	Criteria sets for primary Sjogren's syndrome are not adequate for those presenting with extraglandular organ involvements as their dominant clinical features. <i>Rheumatology International</i> , 2017, 37, 675-684.	3.0	8
17	The frequency of etarnecept related sarcoidosis. , 2017, , .		1
18	Remarkable damage along with poor quality of life in Takayasu arteritis: cross-sectional results of a long-term followed-up multicentre cohort. <i>Clinical and Experimental Rheumatology</i> , 2017, 35 Suppl 103, 77-82.	0.8	7

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19	Is cystatin C an evaluative marker for right heart functions in systemic sclerosis?. International Journal of Cardiology, 2016, 221, 478-483.	1.7	3
20	Response rate of initial conventional treatments, disease course, and related factors of patients with adult-onset Still's disease: Data from a large multicenter cohort. Journal of Autoimmunity, 2016, 69, 59-63.	6.5	62
21	SPO27KILLER CELL IMMUNOGLOBULIN LIKE RECEPTOR GENOTYPE DISTRIBUTION IN FAMILIAL MEDITERRANEAN FEVER: NO ASSOCIATION WITH DISEASE PHENOTYPE AND RENAL AMYLOIDOSIS. Nephrology Dialysis Transplantation, 2015, 30, iii389-iii389.	0.7	0
22	A novel recessive 15-hydroxyprostaglandin dehydrogenase mutation in a family with primary hypertrophic osteoarthropathy. Modern Rheumatology, 2015, 25, 315-321.	1.8	21
23	Identification of Susceptibility Loci in <i>IL6</i> , <i>RPS9</i> / <i>LILRB3</i> , and an Intergenic Locus on Chromosome 21q22 in Takayasu Arteritis in a Genome-Wide Association Study. Arthritis and Rheumatology, 2015, 67, 1361-1368.	5.6	79
24	Incidence of Cyclophosphamide-induced Urotoxicity and Protective Effect of Mesna in Rheumatic Diseases. Journal of Rheumatology, 2015, 42, 1661-1666.	2.0	43
25	Behçet Disease With Vascular Involvement. Medicine (United States), 2015, 94, e494.	1.0	114
26	Sacroiliac joint involvement in systemic sclerosis. International Journal of Rheumatic Diseases, 2015, 18, 84-90.	1.9	9
27	Killer Cell Immunoglobulin-Like Receptor (KIR) Genotype Distribution in Familial Mediterranean Fever (FMF) Patients. Medical Science Monitor, 2015, 21, 3547-3554.	1.1	5
28	Erythema elevatum diutinum coexisting with ankylosing spondylitis. European Journal of Rheumatology, 2015, 2, 73-75.	0.6	2
29	Current antiviral practice and course of Hepatitis B virus infection in inflammatory arthritis: a multicentric observational study (A + HBV study). European Journal of Rheumatology, 2015, 2, 149-154.	0.6	5
30	Amyloidosis and its related factors in Turkish patients with familial Mediterranean fever: a multicentre study. Rheumatology, 2014, 53, 741-745.	1.9	96
31	FMF50: a score for assessing outcome in familial Mediterranean fever. Annals of the Rheumatic Diseases, 2014, 73, 897-901.	0.9	57
32	Mevalonate kinase gene mutations and their clinical correlations in Behçet's disease. International Journal of Rheumatic Diseases, 2014, 17, 435-443.	1.9	7
33	Role of KIR genes and genotypes in susceptibility to or protection against hepatitis B virus infection in a Turkish cohort. Medical Science Monitor, 2014, 20, 28-34.	1.1	15
34	Analysis of Cytotoxic T Lymphocyte Antigen-4 (CTLA-4) Promoter -318C/T and +49A/G Gene Polymorphisms in Turkish Patients with Familial Mediterranean Fever. Cell Biochemistry and Biophysics, 2013, 65, 181-186.	1.8	1
35	A novel recessive 15-hydroxyprostaglandin dehydrogenase mutation in a family with primary hypertrophic osteoarthropathy. Modern Rheumatology, 2013, , 1.	1.8	8
36	Impaired quality of life, disability and mental health in Takayasu's arteritis. Rheumatology, 2013, 52, 1898-1904.	1.9	53

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37	Periodic Fever and Hyperimmunoglobulin D Syndrome in a Boy with Pediatric Autoimmune Neuropsychiatric Disorders Associated with Group A β -Hemolytic Streptococcus. <i>Journal of Child and Adolescent Psychopharmacology</i> , 2013, 23, 302-304.	1.3	10
38	Pigmented Purpuric Dermatitis Associated with Primary Antiphospholipid Syndrome. <i>Internal Medicine</i> , 2013, 52, 1255-1257.	0.7	3
39	Evaluating hand in systemic sclerosis. <i>Rheumatology International</i> , 2012, 32, 3581-3586.	3.0	27
40	Different clinical presentation of the hyperimmunoglobulin D syndrome (HIDS) (four cases from) <i>Tj ETQq0 0 0 rgBT/Overlock 10 Tf 50 6</i>	2.2	13
41	Reply: Sacralization is not associated with elongated cervical costal process and cervical rib. <i>Clinical Anatomy</i> , 2011, 24, 918-919.	2.7	1
42	The significance of E266K polymorphism in the NOD1 gene on Helicobacter Pylori infection: an effective force on pathogenesis?. <i>Clinical and Experimental Medicine</i> , 2010, 10, 107-112.	3.6	29
43	Investigation of C5a receptor gene 450 C/T polymorphism in Turkish patients with familial Mediterranean fever. <i>Molecular Biology Reports</i> , 2010, 37, 273-276.	2.3	7
44	Early suppression of familial Mediterranean fever attacks by single medium dose methyl-prednisolone infusion. <i>Joint Bone Spine</i> , 2008, 75, 370-372.	1.6	29
45	CT-based Morphometric Data of L3-L5 Vertebrae: Anatomic and Surgical Approach. <i>Neurosurgery Quarterly</i> , 2007, 17, 92-97.	0.1	6
46	VEGF 936 C/T gene polymorphism in renal transplant recipients: Association of the T allele with good graft outcome. <i>Human Immunology</i> , 2007, 68, 599-602.	2.4	20
47	Evaluation of the turkish version of the bath ankylosing Spondylitis Patient Global Score (BAS-G). <i>Clinical Rheumatology</i> , 2006, 25, 136-139.	2.2	5
48	Plasma interleukin-10 and interleukin-12 levels in patients with familial Mediterranean fever. <i>Rheumatology International</i> , 2006, 26, 862-864.	3.0	23
49	Serum RANTES, MIP-1 β , and MCP-1 levels in Behçet's disease. <i>Rheumatology International</i> , 2005, 25, 487-488.	3.0	14
50	Evaluation of the Turkish version of the Dougados functional index in ankylosing spondylitis. <i>Rheumatology International</i> , 2005, 25, 368-372.	3.0	2
51	The Turkish version of the Bath Ankylosing Spondylitis Functional Index: reliability and validity. <i>Clinical Rheumatology</i> , 2005, 24, 123-128.	2.2	40
52	Comparison of Amplification Refractory Mutation System and Polymerase Chain Reaction-Restriction Fragment Length Polymorphism Techniques Used for the Investigation of MEFV Gene Exon 10 Point Mutations in Familial Mediterranean Fever Patients Living in Aşkurova Region (Turkey). <i>Genetic Testing and Molecular Biomarkers</i> , 2005, 9, 220-225.	1.7	14
53	Successful Medical Management of Intracardiac Thrombus in a Young Man with Behçet's Disease. <i>Journal of Clinical Rheumatology</i> , 2000, 6, 303-304.	0.9	0
54	Sequence analysis of amyloid protein AA from a Turkish patient with familial Mediterranean fever: Documentation of its SAA1 β derivation. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 1996, 3, 173-176.	3.0	8

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55	The association between HLA B5 and ocular involvement in Behçet's disease in southern Turkey. Acta Ophthalmologica, 1992, 70, 786-789.	1.1	37
56	First Turkish Family with FAP has Homozygous Met 30 TTR. , 1991, , 595-598.		0
57	Homozygosity for the met30 transthyretin gene in a Turkish kindred with familial amyloidotic polyneuropathy. Human Genetics, 1990, 86, 89-90.	3.8	36
58	MANNANOSE BINDING LECTIN (MBL) GENE POLYMORPHISMS AND THEIR RELATIONS WITH CLINICAL FEATURES IN PATIENTS WITH FAMILIAL MEDITERRANEAN FEVER (FMF). İstanbul Tıp Fakültesi Dergisi, 0, , 125-130.	0.0	3