

Ahmet Gl

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

244
papers

11,116
citations

55
h-index

102
g-index

278
ext. papers

13,358
ext. citations

4.3
avg, IF

6.03
L-index

#	Paper	IF	Citations
244	Immunosuppressive therapy in lupus nephritis: the Euro-Lupus Nephritis Trial, a randomized trial of low-dose versus high-dose intravenous cyclophosphamide. <i>Arthritis and Rheumatism</i> , 2002 , 46, 2121-31		649
243	Familial Mediterranean fever (FMF) in Turkey: results of a nationwide multicenter study. <i>Medicine (United States)</i> , 2005 , 84, 1-11	1.8	495
242	EULAR recommendations for the management of Behçet disease. <i>Annals of the Rheumatic Diseases</i> , 2008 , 67, 1656-62	2.4	478
241	Genome-wide association study identifies variants in the MHC class I, IL10, and IL23R-IL12RB2 regions associated with Behçet's disease. <i>Nature Genetics</i> , 2010 , 42, 698-702	36.3	475
240	The 10-year follow-up data of the Euro-Lupus Nephritis Trial comparing low-dose and high-dose intravenous cyclophosphamide. <i>Annals of the Rheumatic Diseases</i> , 2010 , 69, 61-4	2.4	399
239	Genome-wide association analysis identifies new susceptibility loci for Behçet's disease and epistasis between HLA-B*51 and ERAP1. <i>Nature Genetics</i> , 2013 , 45, 202-7	36.3	375
238	Loss-of-function mutations in TNFAIP3 leading to A20 haploinsufficiency cause an early-onset autoinflammatory disease. <i>Nature Genetics</i> , 2016 , 48, 67-73	36.3	359
237	2018 update of the EULAR recommendations for the management of Behçet's syndrome. <i>Annals of the Rheumatic Diseases</i> , 2018 , 77, 808-818	2.4	277
236	Early response to immunosuppressive therapy predicts good renal outcome in lupus nephritis: lessons from long-term followup of patients in the Euro-Lupus Nephritis Trial. <i>Arthritis and Rheumatism</i> , 2004 , 50, 3934-40		255
235	Prevalence of Behçet's disease in Istanbul, Turkey. <i>International Journal of Dermatology</i> , 2003 , 42, 803-6	1.7	251
234	Efficacy of infliximab in the treatment of uveitis that is resistant to treatment with the combination of azathioprine, cyclosporine, and corticosteroids in Behçet's disease: an open-label trial. <i>Arthritis and Rheumatism</i> , 2005 , 52, 2478-84		235
233	Anti-TNF therapy in the management of Behçet's disease--review and basis for recommendations. <i>Rheumatology</i> , 2007 , 46, 736-41	3.9	231
232	Canakinumab for the Treatment of Autoinflammatory Recurrent Fever Syndromes. <i>New England Journal of Medicine</i> , 2018 , 378, 1908-1919	59.2	214
231	Behçet's disease as an autoinflammatory disorder. <i>Inflammation and Allergy: Drug Targets</i> , 2005 , 4, 81-3		181
230	Interleukin-1β-regulating antibody XOMA 052 (gevokizumab) in the treatment of acute exacerbations of resistant uveitis of Behçet's disease: an open-label pilot study. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 563-6	2.4	173
229	Pulmonary manifestations of Behçet's disease. <i>Thorax</i> , 2001 , 56, 572-8	7.3	162
228	Classification criteria for autoinflammatory recurrent fevers. <i>Annals of the Rheumatic Diseases</i> , 2019 , 78, 1025-1032	2.4	159

227	Induced-Pluripotent-Stem-Cell-Derived Primitive Macrophages Provide a Platform for Modeling Tissue-Resident Macrophage Differentiation and Function. <i>Immunity</i> , 2017 , 47, 183-198.e6	32.3	153
226	Two-year results from an open-label, multicentre, phase III study evaluating the safety and efficacy of canakinumab in patients with cryopyrin-associated periodic syndrome across different severity phenotypes. <i>Annals of the Rheumatic Diseases</i> , 2011 , 70, 2095-102	2.4	146
225	Familial aggregation of Behçet's disease in Turkey. <i>Annals of the Rheumatic Diseases</i> , 2000 , 59, 622-5	2.4	134
224	Tuberculosis and other opportunistic infections in tofacitinib-treated patients with rheumatoid arthritis. <i>Annals of the Rheumatic Diseases</i> , 2016 , 75, 1133-8	2.4	134
223	Management of Behçet disease: a systematic literature review for the European League Against Rheumatism evidence-based recommendations for the management of Behçet disease. <i>Annals of the Rheumatic Diseases</i> , 2009 , 68, 1528-34	2.4	131
222	'MHC-I-opathy'-unified concept for spondyloarthritis and Behçet disease. <i>Nature Reviews Rheumatology</i> , 2015 , 11, 731-40	8.1	124
221	Diagnostic criteria for cryopyrin-associated periodic syndrome (CAPS). <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 942-947	2.4	122
220	COVID-19 is a Real Headache!. <i>Headache</i> , 2020 , 60, 1415-1421	4.2	115
219	A20 haploinsufficiency (HA20): clinical phenotypes and disease course of patients with a newly recognised NF-κB-mediated autoinflammatory disease. <i>Annals of the Rheumatic Diseases</i> , 2018 , 77, 728-734	7.4	111
218	Targeted resequencing implicates the familial Mediterranean fever gene MEFV and the toll-like receptor 4 gene TLR4 in Behçet disease. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013 , 110, 8134-9	11.5	105
217	CT findings of pulmonary artery aneurysms during treatment for Behçet's disease. <i>American Journal of Roentgenology</i> , 1999 , 172, 729-33	5.4	105
216	HLA-DRB1*11 and variants of the MHC class II locus are strong risk factors for systemic juvenile idiopathic arthritis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015 , 112, 15970-5	11.5	103
215	Immunology and functional genomics of Behçet's disease. <i>Cellular and Molecular Life Sciences</i> , 2003 , 60, 1903-22	10.3	100
214	Behçet disease-associated MHC class I residues implicate antigen binding and regulation of cell-mediated cytotoxicity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014 , 111, 8867-72	11.5	95
213	Immunohistology of skin pathergy reaction in Behçet's disease. <i>British Journal of Dermatology</i> , 1995 , 132, 901-7	4	93
212	Dense genotyping of immune-related loci implicates host responses to microbial exposure in Behçet's disease susceptibility. <i>Nature Genetics</i> , 2017 , 49, 438-443	36.3	89
211	Genetic architecture distinguishes systemic juvenile idiopathic arthritis from other forms of juvenile idiopathic arthritis: clinical and therapeutic implications. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 906-913	2.4	89
210	Whole-genome screening for susceptibility genes in multicase families with Behçet's disease. <i>Arthritis and Rheumatism</i> , 2005 , 52, 1836-42		89

209	Coagulation factor V gene mutation increases the risk of venous thrombosis in behçt's disease. <i>Rheumatology</i> , 1996 , 35, 1178-80	3.9	83
208	Pathogenesis of Behçt's disease: autoinflammatory features and beyond. <i>Seminars in Immunopathology</i> , 2015 , 37, 413-8	12	79
207	Quality of life changes with canakinumab therapy in adults with colchicine resistant FMF. <i>Pediatric Rheumatology</i> , 2015 , 13,	3.5	78
206	Evidence for linkage of the HLA-B locus in Behçt's disease, obtained using the transmission disequilibrium test. <i>Arthritis and Rheumatism</i> , 2001 , 44, 239-40		78
205	Tuberculosis in Turkish patients with systemic lupus erythematosus: increased frequency of extrapulmonary localization. <i>Lupus</i> , 2004 , 13, 274-8	2.6	74
204	Association of specific interleukin 1 gene cluster polymorphisms with increased susceptibility for Behçet's disease. <i>British Journal of Rheumatology</i> , 2003 , 42, 860-4		74
203	Adult-onset Still's disease. <i>International Journal of Clinical Practice</i> , 2009 , 63, 1050-5	2.9	72
202	Efficacy and safety of canakinumab in adolescents and adults with colchicine-resistant familial Mediterranean fever. <i>Arthritis Research and Therapy</i> , 2015 , 17, 243	5.7	71
201	Comparative study of the skin pathergy test with blunt and sharp needles in Behçt's disease: confirmed specificity but decreased sensitivity with sharp needles. <i>Annals of the Rheumatic Diseases</i> , 1993 , 52, 823-5	2.4	71
200	Use of laser flare-cell photometry to quantify intraocular inflammation in patients with Behçt uveitis. <i>Graefes Archive for Clinical and Experimental Ophthalmology</i> , 2008 , 246, 1169-77	3.8	66
199	Colchicine resistance and intolerance in familial mediterranean fever: Definition, causes, and alternative treatments. <i>Seminars in Arthritis and Rheumatism</i> , 2017 , 47, 115-120	5.3	65
198	Consensus proposal for taxonomy and definition of the autoinflammatory diseases (AIDs): a Delphi study. <i>Annals of the Rheumatic Diseases</i> , 2018 , 77, 1558-1565	2.4	61
197	Cluster analysis of autoantibodies in 852 patients with systemic lupus erythematosus from a single center. <i>Journal of Rheumatology</i> , 2014 , 41, 1304-10	4.1	60
196	HLA-B*51 and Behçt Disease. <i>Ocular Immunology and Inflammation</i> , 2012 , 20, 37-43	2.8	60
195	Characteristics Predicting Tuberculosis Risk under Tumor Necrosis Factor- α Inhibitors: Report from a Large Multicenter Cohort with High Background Prevalence. <i>Journal of Rheumatology</i> , 2016 , 43, 524-9	4.1	58
194	Management of major organ involvement of Behçt's syndrome: a systematic review for update of the EULAR recommendations. <i>Rheumatology</i> , 2018 , 57, 2200-2212	3.9	57
193	Efficacy of anakinra treatment in a patient with colchicine-resistant familial Mediterranean fever. <i>Rheumatology International</i> , 2012 , 32, 3277-9	3.6	57
192	Linkage mapping of a novel susceptibility locus for Behçt's disease to chromosome 6p22-23. <i>Arthritis and Rheumatism</i> , 2001 , 44, 2693-6		56

191	International Retrospective Chart Review of Treatment Patterns in Severe Familial Mediterranean Fever, Tumor Necrosis Factor Receptor-Associated Periodic Syndrome, and Mevalonate Kinase Deficiency/Hyperimmunoglobulinemia D Syndrome. <i>Arthritis Care and Research</i> , 2017 , 69, 578-586	4.7	55
190	Genetics of Behçet's disease: lessons learned from genomewide association studies. <i>Current Opinion in Rheumatology</i> , 2014 , 26, 56-63	5.3	55
189	A weak association of HLA-B*2702 with Behçet's disease. <i>Genes and Immunity</i> , 2002 , 3, 368-72	4.4	55
188	Risk factors for avascular bone necrosis in patients with systemic lupus erythematosus. <i>Rheumatology International</i> , 2012 , 32, 177-82	3.6	54
187	International multi-centre study of pregnancy outcomes with interleukin-1 inhibitors. <i>Rheumatology</i> , 2017 , 56, 2102-2108	3.9	54
186	Development and initial validation of international severity scoring system for familial Mediterranean fever (ISSF). <i>Annals of the Rheumatic Diseases</i> , 2016 , 75, 1051-6	2.4	52
185	Use of Gevokizumab in Patients with Behçet's Disease Uveitis: An International, Randomized, Double-Masked, Placebo-Controlled Study and Open-Label Extension Study. <i>Ocular Immunology and Inflammation</i> , 2018 , 26, 1023-1033	2.8	50
184	Lack of association of HLA-B*51 with a severe disease course in Behçet's disease. <i>British Journal of Rheumatology</i> , 2001 , 40, 668-72		49
183	FMF50: a score for assessing outcome in familial Mediterranean fever. <i>Annals of the Rheumatic Diseases</i> , 2014 , 73, 897-901	2.4	48
182	Serologic response to Epstein-Barr virus antigens in patients with systemic lupus erythematosus: a controlled study. <i>Rheumatology International</i> , 2012 , 32, 79-83	3.6	48
181	Perinatal outcomes in severe preeclampsia-eclampsia with and without HELLP syndrome. <i>Gynecologic and Obstetric Investigation</i> , 2005 , 59, 113-8	2.5	47
180	Peripheral blood T cell expansions in patients with Behçet's disease. <i>Clinical and Experimental Immunology</i> , 1997 , 107, 520-7	6.2	47
179	Inflammatory/demyelinating central nervous system involvement in familial Mediterranean fever (FMF): coincidence or association?. <i>Journal of Neurology</i> , 2006 , 253, 928-34	5.5	46
178	Low frequency of HLA-B27 in ankylosing spondylitis patients from Turkey. <i>Joint Bone Spine</i> , 2008 , 75, 299-302	2.9	45
177	Association of familial Mediterranean fever-related MEFV variations with ankylosing spondylitis. <i>Arthritis and Rheumatism</i> , 2010 , 62, 3232-6		42
176	Association of Major Histocompatibility Complex Class I Chain-Related Gene A and HLA-B Alleles with Behçet's Disease in Turkey. <i>Japanese Journal of Ophthalmology</i> , 2007 , 51, 431-6	2.6	42
175	Procoagulant mutations and venous thrombosis in Behçet's disease. <i>Rheumatology</i> , 1999 , 38, 1298-9	3.9	42
174	A single endoplasmic reticulum aminopeptidase-1 protein allotype is a strong risk factor for Behçet's disease in HLA-B*51 carriers. <i>Annals of the Rheumatic Diseases</i> , 2016 , 75, 2208-2211	2.4	42

173	Pulmonary hypertension in systemic lupus erythematosus: relationship with antiphospholipid antibodies and severe disease outcome. <i>Rheumatology International</i> , 2011 , 31, 183-9	3.6	41
172	Pro-inflammatory cellular immune response in Behçet's disease. <i>Rheumatology International</i> , 2007 , 27, 1113-8	3.6	40
171	Organizing pneumonia associated with pulmonary artery aneurysms in Behçet's disease. <i>Rheumatology</i> , 1999 , 38, 1285-9	3.9	39
170	Ancient familial Mediterranean fever mutations in human pyrin and resistance to <i>Yersinia pestis</i> . <i>Nature Immunology</i> , 2020 , 21, 857-867	19.1	38
169	Management of skin, mucosa and joint involvement of Behçet's syndrome: A systematic review for update of the EULAR recommendations for the management of Behçet's syndrome. <i>Seminars in Arthritis and Rheumatism</i> , 2019 , 48, 752-762	5.3	38
168	Perinatal outcomes of twin pregnancies discordant for major fetal anomalies. <i>Fetal Diagnosis and Therapy</i> , 2005 , 20, 244-8	2.4	38
167	Polymorphisms in the endothelial nitric oxide synthase gene are associated with Behçet's disease. <i>Rheumatology</i> , 2005 , 44, 614-7	3.9	38
166	Safety and Efficacy of Gevokizumab in Patients with Behçet's Disease Uveitis: Results of an Exploratory Phase 2 Study. <i>Ocular Immunology and Inflammation</i> , 2017 , 25, 62-70	2.8	36
165	Anti-CCP and antikeratin antibodies in rheumatoid arthritis, primary Sjögren's syndrome, and Wegener's granulomatosis. <i>Clinical Rheumatology</i> , 2005 , 24, 673-6	3.9	33
164	Metabolic syndrome is not only a risk factor for cardiovascular diseases in systemic lupus erythematosus but is also associated with cumulative organ damage: a cross-sectional analysis of 311 patients. <i>Lupus</i> , 2016 , 25, 177-84	2.6	32
163	A comparison of clinical findings of familial Mediterranean fever patients with and without amyloidosis. <i>Rheumatology International</i> , 2005 , 25, 442-6	3.6	32
162	Standard and novel therapeutic approaches to Behçet's disease. <i>Drugs</i> , 2007 , 67, 2013-22	12.1	31
161	Familial Mediterranean fever and seronegative arthritis. <i>Current Rheumatology Reports</i> , 2011 , 13, 388-94	4.9	30
160	Impaired endothelium-dependent flow-mediated dilation in Behçet's disease: more prominent endothelial dysfunction in patients with vascular involvement. <i>International Journal of Clinical Practice</i> , 2005 , 59, 777-81	2.9	30
159	Treatment of recurrent perforating intestinal ulcers with thalidomide in Behçet's disease. <i>Annals of Pharmacotherapy</i> , 2004 , 38, 808-11	2.9	28
158	Comparison of Disease Characteristics, Organ Damage, and Survival in Patients with Juvenile-onset and Adult-onset Systemic Lupus Erythematosus in a Combined Cohort from 2 Tertiary Centers in Turkey. <i>Journal of Rheumatology</i> , 2017 , 44, 619-625	4.1	27
157	Association of the MEFV gene variations with inflammatory bowel disease in Turkey. <i>Journal of Clinical Gastroenterology</i> , 2013 , 47, e23-7	3	27
156	Update on Outcome Measure Development for Large Vessel Vasculitis: Report from OMERACT 12. <i>Journal of Rheumatology</i> , 2015 , 42, 2465-9	4.1	26

155	Diagnostic performance of amyloid A protein quantification in fat tissue of patients with clinical AA amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2007 , 14, 133-40	2.7	26
154	Development of a Core Set of Outcome Measures for Large-vessel Vasculitis: Report from OMERACT 2016. <i>Journal of Rheumatology</i> , 2017 , 44, 1933-1937	4.1	25
153	Anti-neuronal and stress-induced-phosphoprotein 1 antibodies in neuro-Behçet's disease. <i>Journal of Neuroimmunology</i> , 2011 , 239, 91-7	3.5	25
152	Predictors of damage and survival in patients with Wegener's granulomatosis: analysis of 50 patients. <i>Journal of Rheumatology</i> , 2010 , 37, 374-8	4.1	25
151	Follow-up results of 702 patients receiving tumor necrosis factor- α antagonists and evaluation of risk of tuberculosis. <i>Rheumatology International</i> , 2010 , 30, 1459-63	3.6	25
150	Expression of KIR and C-type lectin receptors in Behçet's disease. <i>British Journal of Rheumatology</i> , 2004 , 43, 423-7		25
149	Primary antiphospholipid syndrome associated with mesenteric inflammatory veno-occlusive disease. <i>Clinical Rheumatology</i> , 1996 , 15, 207-10	3.9	25
148	Common genetic susceptibility loci link PFAPA syndrome, Behçet's disease, and recurrent aphthous stomatitis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020 , 117, 14405-14411	11.5	24
147	The regulation of MEFV expression and its role in health and familial Mediterranean fever. <i>Genes and Immunity</i> , 2011 , 12, 497-503	4.4	23
146	Predicting important residues and interaction pathways in proteins using Gaussian Network Model: binding and stability of HLA proteins. <i>PLoS Computational Biology</i> , 2010 , 6, e1000845	5	23
145	IL1RN Variation Influences Both Disease Susceptibility and Response to Recombinant Human Interleukin-1 Receptor Antagonist Therapy in Systemic Juvenile Idiopathic Arthritis. <i>Arthritis and Rheumatology</i> , 2018 , 70, 1319-1330	9.5	22
144	Approach to the patients with inadequate response to colchicine in familial Mediterranean fever. <i>Best Practice and Research in Clinical Rheumatology</i> , 2016 , 30, 296-303	5.3	22
143	HSP 60 expression in recurrent oral ulcerations of Behçet's disease. <i>Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics</i> , 2010 , 110, 196-200		22
142	Safety of high-dose intravenous immunoglobulin in systemic autoimmune diseases. <i>Clinical Rheumatology</i> , 2007 , 26, 1913-5	3.9	21
141	Impaired brachial endothelial function in patients with primary anti-phospholipid syndrome. <i>International Journal of Clinical Practice</i> , 2004 , 58, 1003-7	2.9	21
140	Re-evaluation of heterogeneity in HLA-B*510101 associated with Behçet's disease. <i>Tissue Antigens</i> , 2008 , 72, 347-53		20
139	Association between reduced levels of MEFV messenger RNA in peripheral blood leukocytes and acute inflammation. <i>Arthritis and Rheumatism</i> , 2007 , 56, 345-50		20
138	Experience with monthly, high-dose, intravenous immunoglobulin therapy in patients with different connective tissue diseases. <i>Rheumatology International</i> , 2005 , 25, 211-4	3.6	20

137	Developing a Core Set of Outcome Measures for Behçet Disease: Report from OMERACT 2016. <i>Journal of Rheumatology</i> , 2017 , 44, 1750-1753	4.1	18
136	Anti-interferon alpha antibodies and autoantibodies in patients with Behçet's disease uveitis treated with recombinant human interferon alpha-2a. <i>Graefes Archive for Clinical and Experimental Ophthalmology</i> , 2015 , 253, 457-65	3.8	18
135	Expression of regulatory receptors on T cells and their cytokine production in Behçet's disease. <i>Arthritis Research and Therapy</i> , 2013 , 15, R15	5.7	18
134	Effect of Interferon alfa-2a Treatment on Adaptive and Innate Immune Systems in Patients With Behçet Disease Uveitis 2019 , 60, 52-63		17
133	Treatment of familial Mediterranean fever: colchicine and beyond. <i>Israel Medical Association Journal</i> , 2014 , 16, 281-4	0.9	17
132	Small bowel mucosal damage in familial Mediterranean fever: results of capsule endoscopy screening. <i>Scandinavian Journal of Gastroenterology</i> , 2014 , 49, 1414-8	2.4	16
131	Identification of possible pathogenic pathways in Behçet's disease using genome-wide association study data from two different populations. <i>European Journal of Human Genetics</i> , 2015 , 23, 678-87	5.3	15
130	Comparison of perinatal and maternal outcomes of severe preeclampsia, eclampsia, and HELLP syndrome. <i>Journal of the Turkish German Gynecology Association</i> , 2011 , 12, 90-6	1.1	15
129	Distribution of common CARD15 variants in patients with sporadic Crohn's disease: cases from Turkey. <i>Digestive Diseases and Sciences</i> , 2006 , 51, 706-10	4	15
128	Current Status, Goals, and Research Agenda for Outcome Measures Development in Behçet Syndrome: Report from OMERACT 2014. <i>Journal of Rheumatology</i> , 2015 , 42, 2436-41	4.1	14
127	Mitochondrial carrier homolog 1 (Mtch1) antibodies in neuro-Behçet's disease. <i>Journal of Neuroimmunology</i> , 2013 , 263, 139-44	3.5	14
126	Low-dose natural human interferon-alpha lozenges in the treatment of Behçet's syndrome. <i>Rheumatology</i> , 2009 , 48, 1388-91	3.9	14
125	No association of the TLR2 gene Arg753Gln polymorphism with rheumatic heart disease and Behçet's disease. <i>Clinical Rheumatology</i> , 2009 , 28, 1385-8	3.9	14
124	Antibodies reactive with HIV-1 antigens in systemic lupus erythematosus. <i>Lupus</i> , 1996 , 5, 120-2	2.6	14
123	Eosinophilia and hyperimmunoglobulinemia E as the presenting manifestations of Wegener's granulomatosis. <i>Clinical Rheumatology</i> , 2003 , 22, 333-5	3.9	14
122	Increased IL-23 receptor, TNF- α and IL-6 expression in individuals with the IL23R-IL12RB2 locus polymorphism. <i>Immunology Letters</i> , 2014 , 160, 96-98	4.1	13
121	Re-evaluation of 129 patients with systemic necrotizing vasculitides by using classification algorithm according to consensus methodology. <i>Clinical Rheumatology</i> , 2012 , 31, 325-8	3.9	13
120	Behçet's disease: immunological relevance with arthritis of ankylosing spondylitis. <i>Rheumatology International</i> , 2013 , 33, 733-41	3.6	13

119	Evaluation of KIR3DL1/KIR3DS1 polymorphism in Behçet's disease. <i>Genes and Immunity</i> , 2016 , 17, 396-399	4.4	12
118	The use of two different Health Assessment Questionnaires in Turkish rheumatoid arthritis population and assessment of the associations with disability. <i>Clinical Rheumatology</i> , 1999 , 18, 33-7	3.9	12
117	An International Delphi Survey for the Definition of New Classification Criteria for Familial Mediterranean Fever, Mevalonate Kinase Deficiency, TNF Receptor-associated Periodic Fever Syndromes, and Cryopyrin-associated Periodic Syndrome. <i>Journal of Rheumatology</i> , 2019 , 46, 429-436	4.1	12
116	Dynamics of Inflammatory Response in Autoinflammatory Disorders: Autonomous and Hyperinflammatory States. <i>Frontiers in Immunology</i> , 2018 , 9, 2422	8.4	12
115	Recovery from hypoxia and hypercapnic hypoxia: impacts on the transcription of key antioxidants in the shrimp <i>Litopenaeus vannamei</i> . <i>Comparative Biochemistry and Physiology - B Biochemistry and Molecular Biology</i> , 2014 , 170, 43-9	2.3	11
114	Amyloid arthropathy mimicking seronegative rheumatoid arthritis in multiple myeloma: case reports and review of the literature. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2009 , 16, 226-31	2.7	11
113	Generation of integration-free induced pluripotent stem cells from a patient with Familial Mediterranean Fever (FMF). <i>Stem Cell Research</i> , 2015 , 15, 694-6	1.6	10
112	TNF-alpha antagonist therapy modify the tuberculin skin test response. <i>Rheumatology International</i> , 2011 , 31, 1147-51	3.6	10
111	Amyopathic dermatomyositis associated with cervical cancer. <i>Clinical Rheumatology</i> , 2001 , 20, 438-40	3.9	10
110	Shared epitope homozygosity' is strongly associated with rheumatoid arthritis in Turkey. Istanbul Rheumatology Study Group. <i>British Journal of Rheumatology</i> , 1998 , 37, 1126-8		10
109	Natural killer cells dominate a Th-1 polarized response in Behçet's disease patients with uveitis. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, S24-9	2.2	10
108	Unified Modeling of Familial Mediterranean Fever and Cryopyrin Associated Periodic Syndromes. <i>Computational and Mathematical Methods in Medicine</i> , 2015 , 2015, 893507	2.8	9
107	An Algorithm for the Diagnosis of Behçet Disease Uveitis in Adults. <i>Ocular Immunology and Inflammation</i> , 2020 , 1-10	2.8	9
106	HLA-B51 negative monozygotic twins discordant for Behçet's disease. <i>Rheumatology</i> , 1997 , 36, 922-3	3.9	8
105	Colchicine-induced myopathy in a teenager with familial Mediterranean fever. <i>Annals of Pharmacotherapy</i> , 2003 , 37, 1821-4	2.9	8
104	Direct and indirect costs associated with ankylosing spondylitis and related disease activity scores in Turkey. <i>Rheumatology International</i> , 2015 , 35, 1473-8	3.6	7
103	The OMERACT Core Set of Domains for Outcome Measures in Behçet Syndrome. <i>Arthritis Care and Research</i> , 2020 ,	4.7	7
102	The effect of tocilizumab, anakinra and prednisolone on antibody response to SARS-CoV-2 in patients with COVID-19: A prospective cohort study with multivariate analysis of factors affecting the antibody response. <i>International Journal of Infectious Diseases</i> , 2021 , 105, 756-762	10.5	7

101	Phenotypic variability including Behçet's disease-like manifestations in DADA2 patients due to a homozygous c.973-2A>G splice site mutation. <i>Clinical and Experimental Rheumatology</i> , 2019 , 37 Suppl 121, 142-146	2.2	7
100	Microscopic colitis in patients with Takayasu's arteritis: a potential association between the two disease entities. <i>Clinical Rheumatology</i> , 2016 , 35, 2495-9	3.9	6
99	MEFV gene variations in patients with systemic lupus erythematosus. <i>Modern Rheumatology</i> , 2014 , 24, 93-6	3.3	6
98	Recurrent anterior uveitis in Henoch Schonlein's vasculitis. <i>Rheumatology International</i> , 2010 , 30, 1377-9	3.6	6
97	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	2.2	6
96	The efficacy of anti- IL-1 treatment in three patients with coexisting familial Mediterranean fever and multiple sclerosis. <i>Multiple Sclerosis and Related Disorders</i> , 2020 , 45, 102332	4	5
95	Comment on the article by Durmus et al. "Clinical significance of MEFV mutations in ankylosing spondylitis". <i>Joint Bone Spine</i> , 2010 , 77, 281	2.9	5
94	Systemic necrotizing vasculitides in Turkey: a comparative analysis of 40 consecutive patients. <i>Rheumatology International</i> , 2005 , 26, 16-20	3.6	5
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