Ahmet Gül

List of Publications by Year in descending order

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274 papers 14,919 citations

20817 60 h-index 117 g-index

278 all docs

278 docs citations

278 times ranked 11694 citing authors

#	Article	IF	CITATIONS
1	Core Set of Domains for Outcome Measures in Behçet's Syndrome. Arthritis Care and Research, 2022, 74, 691-699.	3.4	21
2	Tocilizumab treatment in severe COVID-19: a multicenter retrospective study with matched controls. Rheumatology International, 2022, 42, 457-467.	3.0	4
3	IL-17A and IFN-γ are Up-regulated in CD4 and γδT Cells in Active Behcet's Disease Patients. Immunology Letters, 2022, 242, 37-45.	2.5	8
4	Psychometric validation of Barkley's Adult Sluggish Cognitive Tempo (SCT) Ratings Scale -Turkish version and distinguishing SCT from attention deficit-hyperactivity disorder (ADHD) among Turkish adults. Research in Developmental Disabilities, 2022, 121, 104155.	2.2	6
5	Description of damage in different clusters of patients with antiphospholipid syndrome. Lupus, 2022, 31, 433-442.	1.6	5
6	A Case of Takayasu Arteritis with Thrombotic Microangiopathy Secondary to Malignant Hypertension Due to Bilateral Renal Artery Stenosis. Open Access Rheumatology: Research and Reviews, 2022, Volume 14, 39-42.	1.6	2
7	Frequency and severity of COVIDâ€19 in patients with various rheumatic diseases treated regularly with colchicine or hydroxychloroquine. Journal of Medical Virology, 2022, 94, 3431-3437.	5.0	4
8	The relationship between serum A proliferation-inducing ligand and B-cell activating factor levels with disease activity and organ involvement in systemic lupus erythematosus. Lupus, 2022, 31, 555-564.	1.6	1
9	Effect of different cytokines in combination with IL-15 on the expression of activating receptors in NK cells of patients with Behçet's disease. Immunologic Research, 2022, 70, 654-666.	2.9	4
10	An Algorithm for the Diagnosis of Beh $\tilde{\text{A}}$ set Disease Uveitis in Adults. Ocular Immunology and Inflammation, 2021, 29, 1154-1163.	1.8	26
11	Validation of the adjusted global antiphospholipid syndrome score in a single centre cohort of APS patients from Turkey. Journal of Thrombosis and Thrombolysis, 2021, 51, 466-474.	2.1	6
12	Defining colchicine resistance/intolerance in patients with familial Mediterranean fever: a modified-Delphi consensus approach. Rheumatology, 2021, 60, 3799-3808.	1.9	29
13	Long-term follow-up of 89 patients with giant cell arteritis: a retrospective observational study on disease characteristics, flares and organ damage. Rheumatology International, 2021, 41, 439-448.	3.0	2
14	mRNA Expression Levels of NKp30, NKp46, NKG2D, Perforin and Granzyme in the Behçet's Disease. Turkish Journal of Immunology, 2021, , .	0.1	0
15	A shared motif of hla-dpb1 affecting the susceptibility to pr3-anca positive granulomatosis with polyangiitis: comparative analysis of a Turkish cohort with matched healthy controls. Rheumatology International, 2021 , 41 , 1667 - 1672 .	3.0	2
16	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. International Journal of Infectious Diseases, 2021, 105, 756-762.	3. 3	16
17	POS0142â€MINIMAL DISEASE ACTIVITY IN PATIENTS WITH PSORIATIC ARTHRITIS AND ASSOCIATED FACTORS: REAL LIFE DATA FROM A SINGLE CENTER. Annals of the Rheumatic Diseases, 2021, 80, 282.3-283.	0.9	0
18	AB0766â€SUCCESSFUL TEATMENT OF ANKYLOSING SPONDILITIS ASSOCIATED AA AMYLOIDOSIS WITH SECUKINUMAB: A CASE SERIES WITH THREE PATIENTS. Annals of the Rheumatic Diseases, 2021, 80, 1409.3-1410.	0.9	1

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19	ABO448â€SYSTEMIC SCLEROSIS ASSOCIATED PULMONARY ARTERIAL HYPERTENSION: PREDOMINANCE OF PULMONARY FIBROSIS AS A RISK FACTOR FOR MORTALITY IN A SINGLE CENTER COHORT. Annals of the Rheumatic Diseases, 2021, 80, 1251.2-1252.	0.9	0
20	POS1339â€MORE FREQUENT AND EARLIER HIP INVOLVEMENT IN SPONDYLOARTHRITIS ASSOCIATED WITH FAMILIAL MEDITERRANEAN FEVER. Annals of the Rheumatic Diseases, 2021, 80, 951.3-952.	0.9	0
21	AB0653â€COURSE OF COVID-19 INFECTION IN A SERIES OF PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSL Annals of the Rheumatic Diseases, 2021, 80, 1359.1-1359.	IS. 0.9	0
22	POS0706â€PERFORMANCES OF DIFFERENT CLASSIFICATION CRITERIA FOR SYSTEMIC LUPUS ERYTHEMATOSUS IN A SINGLE CENTER COHORT FROM TURKEY. Annals of the Rheumatic Diseases, 2021, 80, 602.1-603.	S _{0.9}	0
23	POS0766â€CLUSTER ANALYSIS AND COMPARISON OF CUMULATIVE DAMAGE BY DIAPS IN A SINGLE CENTER COHORT OF APS PATIENTS. Annals of the Rheumatic Diseases, 2021, 80, 636.2-637.	0.9	0
24	POS1259â€FAVOURABLE SHORT-TERM COURSE OF COVID-19 IN PATIENTS WITH FAMILIAL MEDITERRANEAN FEVER USING BIOLOGIC AGENTS. Annals of the Rheumatic Diseases, 2021, 80, 913.2-914.	0.9	0
25	POS1445â€RETINOL BINDING PROTEIN 4 AS AN ACUTE PHASE REACTANT AND BIOMARKER IN PATIENTS WITH FAMILIAL MEDITERRANEAN FEVER AND AMYLOIDOSIS COMPARED TO INFECTIONS. Annals of the Rheumatic Diseases, 2021, 80, 1006.2-1007.	0.9	O
26	AB0331â€PULMONARY INVOLVEMENT IN A SINGLE CENTER COHORT OF PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSUS. Annals of the Rheumatic Diseases, 2021, 80, 1191.1-1191.	0.9	0
27	POSO710â€ANALYSIS OF 5-YEAR HOSPITALIZATION DATA OF PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOS DAMAGE IS A RISK FACTOR FOR FREQUENT AND LONGER STAYS. Annals of the Rheumatic Diseases, 2021, 80, 604.2-605.	US: 0.9	O
28	OP0313â€PRELIMINARY CRITERIA FOR MACROPHAGE ACTIVATION SYNDROME ASSOCIATED WITH CORONAVII DISEASE-19. Annals of the Rheumatic Diseases, 2021, 80, 191.1-192.	RUS 0.9	0
29	AB0369â€EVALUATION OF BASELINE POSITRON EMISSION TOMOGRAPHY IN THE DIAGNOSIS AND ASSESSMEN OF GIANT CELL ARTERITIS. Annals of the Rheumatic Diseases, 2021, 80, 1210.2-1211.	IT _{0.9}	0
30	POS1257â€HYPOGAMMAGLOBULINEMIA IS A SIGNIFICANT RISK FACTOR FOR MORTALITY IN PATIENTS WITH ANCA ASSOCIATED VASCULITIS AND COVID-19. Annals of the Rheumatic Diseases, 2021, 80, 912-913.	0.9	3
31	Reinfection with SARSâ€CoVâ€2 in a kidney transplant recipient. Transplant Infectious Disease, 2021, 23, e13695.	1.7	2
32	Inflammatory status might direct ILC and NK cells to IL-17 expressing ILC3 and NK subsets in Behcet's disease. Immunology Letters, 2021, 235, 1-8.	2.5	8
33	Budd–Chiari syndrome in Behçet's disease: a retrospective multicenter study. Clinical Rheumatology, 2021, , 1.	2.2	7
34	A single center survey study of systemic vasculitis and COVID-19 during the first months of pandemic. Turkish Journal of Medical Sciences, 2021, 51, 2243-2247.	0.9	3
35	Initial complete blood count score and predicting disease progression in COVID-19 patients. American Journal of Blood Research, 2021, 11, 77-83.	0.6	0
36	Real life data of secukinumab in ankylosing spondylitis and psoriatic arthritis: Analysis of 44 patients in a single center. Ulusal Romatoloji Dergisi, 2021, 13, 116-120.	0.0	0

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37	Clinical characteristics of female patients with axial spondyloarthritis treated with tumor necrosis factor-alpha inhibitors and disease related outcomes after therapy. Ulusal Romatoloji Dergisi, 2021, 13, 100-106.	0.0	0
38	Monogenic lupus due to spondyloenchondrodysplasia with spastic paraparesis and intracranial calcification: case-based review. Rheumatology International, 2020, 40, 1903-1910.	3.0	19
39	Molecular dynamics simulations provide molecular insights into the role of HLAâ€B51 in Behçet's disease pathogenesis. Chemical Biology and Drug Design, 2020, 96, 644-658.	3.2	8
40	P26â \in Serum BAFF and APRIL as candidate biomarkers in systemic lupus erythematosus (SLE): a prospective follow-up study. , 2020, , .		0
41	P69 Risk factors for adverse pregnancy outcome in patients with SLE. , 2020, , .		0
42	P73â€The relationship between pregnancy, disease activity and adverse pregnancy outcomes in systemic lupus erythematosus. , 2020, , .		0
43	P181â€Comparison of SLEDAI-2K and SLEDAI-2KG (glucocorticoid) indexes in patients with systemic lupus erythematosus (SLE). , 2020, , .		0
44	P9â€Validation of the adjusted global antiphospholipid syndrome score and correlation with extra-criteria manifestations. , 2020, , .		0
45	Letter: is pneumococcal vaccination safe during the COVIDâ€19 pandemic?. Alimentary Pharmacology and Therapeutics, 2020, 52, 919-920.	3.7	2
46	COVIDâ€19 is a Real Headache!. Headache, 2020, 60, 1415-1421.	3.9	204
47	Common genetic susceptibility loci link PFAPA syndrome, Behçet's disease, and recurrent aphthous stomatitis. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 14405-14411.	7.1	52
48	Ancient familial Mediterranean fever mutations in human pyrin and resistance to Yersinia pestis. Nature Immunology, 2020, 21, 857-867.	14.5	90
49	The efficacy of anti- IL-1 treatment in three patients with coexisting familial Mediterranean fever and multiple sclerosis. Multiple Sclerosis and Related Disorders, 2020, 45, 102332.	2.0	11
50	Serum and urine TNF-like weak inducer of apoptosis, monocyte chemoattractant protein-1 and neutrophil gelatinase-associated lipocalin as biomarkers of disease activity in patients with systemic lupus erythematosus. Lupus, 2020, 29, 379-388.	1.6	11
51	Genetics of Behçet's Disease. , 2020, , 223-233.		0
52	AB0485â€INVESTIGATION OF PERMANENT ORGAN DAMAGE IN GIANT CELL ARTERITIS: DISEASE FLARES ARE ASSOCIATED WITH INCREASED DAMAGE SCORES. Annals of the Rheumatic Diseases, 2020, 79, 1540.2-1540.	0.9	0
53	SAT0178â€HYDROXYCHLOROQUINE CONTROLS DISEASE ACTIVITY IN SLE AND MULTIMODAL IMAGING TECHNIQUES SHOULD BE USED TO DETECT OCULAR TOXICITY. Annals of the Rheumatic Diseases, 2020, 79, 1030.2-1031.	0.9	0
54	ABO486â€ANALYSIS OF 89 PATIENTS WITH GIANT CELL ARTERITIS FROM TURKEY: PET-CT AS AN EMERGING METHOD FOR DIAGNOSIS AND HIGH FLARE RATE WITH STANDARD CARE. Annals of the Rheumatic Diseases, 2020, 79, 1540.1-1541.	0.9	0

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55	ABO480â€AN OPEN-LABEL, EXPLORATORY STUDY TO ESTABLISH THE EFFICACY AND SAFETY OF 1-YEAR CANAKINUMAB TREATMENT IN BEHÇET'S DISEASE PATIENTS WITH NEUROLOGIC OR VASCULAR INVOLVEMENT. Annals of the Rheumatic Diseases, 2020, 79, 1537.3-1538.	0.9	0
56	FRIO160â€THE CORRELATION BETWEEN PREGNANCY, DISEASE ACTIVITY AND ADVERSE PREGNANCY OUTCOMI IN PATIENTS WITH SYSTEMIC LUPUS ERYTHEMATOSUS. Annals of the Rheumatic Diseases, 2020, 79, 664.1-664.	ES 0.9	0
57	SAT0261â€FEATURES AND RISK FACTORS OF SERIOUS INFECTIONS IN ANCA ASSOCIATED VASCULITIS: LONG TERM FOLLOW UP OF 186 PATIENTS. Annals of the Rheumatic Diseases, 2020, 79, 1073.1-1073.	0.9	0
58	THU0369â€EVALUATION OF DIFFERENT CLASSIFICATION CRITERIA IN SYSTEMIC SCLEROSIS IN A TURKISH COHORT: THE IMPORTANCE OF NON-SKIN MANIFESTATIONS, SEROLOGY AND CAPILLAROSCOPY. Annals of the Rheumatic Diseases, 2020, 79, 416.1-417.	0.9	O
59	SAT0200â€RISK FACTORS FOR ADVERSE PREGNANCY OUTCOMES IN SYSTEMIC LUPUS ERYTHEMATOSUS PATIENTS. Annals of the Rheumatic Diseases, 2020, 79, 1042.1-1042.	0.9	0
60	SATO346â€THE EFFICACY AND SAFETY OF RITUXIMAB IN 27 CASES OF TREATMENT RESISTANT SYSTEMIC SCLEROSIS WITH SEVERE DISEASE ASSESSED BY ACTIVITY SCORES. Annals of the Rheumatic Diseases, 2020, 79, 1119.2-1119.	0.9	O
61	SAT0238â€VALIDATION OF THE ADJUSTED GLOBAL ANTIPHOSPHOLIPID SYNDROME SCORE AND CORRELATION WITH EXTRA-CRITERIA MANIFESTATIONS. Annals of the Rheumatic Diseases, 2020, 79, 1062.2-1062.	N _{0.9}	0
62	OP0274â€CLINICAL ASPECTS, LABORATORY CHARACTERISTICS AND TREATMENT RESPONSES OF AA AMYLOIDOSIS: SINGLE CENTER EXPERIENCE WITH 163 PATIENTS. Annals of the Rheumatic Diseases, 2020, 79, 171.1-171.	0.9	0
63	THU0349â€THE RELATIONSHIP BETWEEN DISEASE ACTIVITY AND SEVERITY IN SYSTEMIC SCLEROSIS: A PROSPECTIVE ANALYSIS OF 278 PATIENTS. Annals of the Rheumatic Diseases, 2020, 79, 406.2-406.	0.9	O
64	AB0461â€ANCA-ASSOCIATED VASCULITIS: CLINICAL FEATURES, RELAPSE, ORGAN DAMAGE AND SURVIVAL IN 1 PATIENTS. Annals of the Rheumatic Diseases, 2020, 79, 1529.1-1529.	97 0.9	0
65	Cardiovascular risk in antiphospholipid syndrome: A comparison of thrombotic and obstetric disease subsets. Ulusal Romatoloji Dergisi, 2020, 12, 83-87.	0.0	O
66	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. Seminars in Arthritis and Rheumatism, 2019, 48, 752-762.	3.4	56
67	Autoimmune haemolytic anaemia and thrombocytopaenia in a single-centre cohort of patients with systemic lupus erythematosus from Turkey: clinical associations and effect on disease damage and survival. Lupus, 2019, 28, 1480-1487.	1.6	8
68	Classification criteria for autoinflammatory recurrent fevers. Annals of the Rheumatic Diseases, 2019, 78, 1025-1032.	0.9	300
69	Update on Outcome Measure Development in Large-vessel Vasculitis: Report from OMERACT 2018. Journal of Rheumatology, 2019, 46, 1198-1201.	2.0	24
70	Clinical course of abdominal aortic aneurysms in Beh $\tilde{\text{A}}$ §et disease: a retrospective analysis. Rheumatology International, 2019, 39, 1061-1067.	3.0	9
71	Behçet Disease. , 2019, , 647-665.		3
72	AB0524â€SERUM BAFF AND APRIL AS CANDIDATE BIOMARKERS IN SYSTEMİC LUPUS ERYTHEMATOSUS (SLE) PROSPECTIVE FOLLOW-UP STUDY. , 2019, , .	: A	0

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7 3	THU0303â€THE OMERACT CORE DOMAIN SET FOR CLINICAL TRIALS IN BEHÇET'S SYNDROME. , 2019, , .		2
74	281â€Lupus nephritis biomarkers. , 2019, , .		0
75	SATO184â€ASSOCIATION OF SERUM AND URINE LEVELS OF TWEAK, MCP-1 AND NGAL WITH DISEASE ACTIVITY SYSTEMIC LUPUS ERYTHEMATOSUS. , 2019, , .	Y IN	0
76	Effect of Interferon alfa-2a Treatment on Adaptive and Innate Immune Systems in Patients With Beh \tilde{A} Set Disease Uveitis., 2019, 60, 52.		29
77	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. Journal of Rheumatology, 2019, 46, 429-436.	2.0	16
78	Improvement of Libman-Sacks endocarditis after combination of warfarin and immunosuppressive therapy. Turk Kardiyoloji Dernegi Arsivi, 2019, 47, 687-690.	0.5	4
79	Pseudodominance of autoinflammatory disease in a single Turkish family explained by co-inheritance of haploinsufficiency of A20 and familial Mediterranean fever. Clinical and Experimental Rheumatology, 2019, 37 Suppl 121, 89-92.	0.8	3
80	Phenotypic variability including Behçet's disease-like manifestations in DADA2 patients due to a homozygous c.973-2A>G splice site mutation. Clinical and Experimental Rheumatology, 2019, 37 Suppl 121, 142-146.	0.8	7
81	<i>IL1RN</i> Variation Influences Both Disease Susceptibility and Response to Recombinant Human Interleukin†Receptor Antagonist Therapy in Systemic Juvenile Idiopathic Arthritis. Arthritis and Rheumatology, 2018, 70, 1319-1330.	5.6	40
82	2018 update of the EULAR recommendations for the management of Behçet's syndrome. Annals of the Rheumatic Diseases, 2018, 77, annrheumdis-2018-213225.	0.9	442
83	Use of Gevokizumab in Patients with Behçet's Disease Uveitis: An International, Randomized, Double-Masked, Placebo-Controlled Study and Open-Label Extension Study. Ocular Immunology and Inflammation, 2018, 26, 1023-1033.	1.8	82
84	A20 haploinsufficiency (HA20): clinical phenotypes and disease course of patients with a newly recognised NF-kB-mediated autoinflammatory disease. Annals of the Rheumatic Diseases, 2018, 77, 728-735.	0.9	176
85	Binding stability of peptides on major histocompatibility complex class I proteins: role of entropy and dynamics. Physical Biology, 2018, 15, 026005.	1.8	2
86	The association between P selectin glycoprotein ligand 1 gene variable number of tandem repeats polymorphism and risk of thrombosis in Behçet's disease. International Journal of Rheumatic Diseases, 2018, 21, 2175-2179.	1.9	0
87	Dynamics of Inflammatory Response in Autoinflammatory Disorders: Autonomous and Hyperinflammatory States. Frontiers in Immunology, 2018, 9, 2422.	4.8	19
88	Canakinumab for the Treatment of Autoinflammatory Recurrent Fever Syndromes. New England Journal of Medicine, 2018, 378, 1908-1919.	27.0	327
89	Behçet Disease. , 2018, , 365-370.		0
90	Consensus proposal for taxonomy and definition of the autoinflammatory diseases (AIDs): a Delphi study. Annals of the Rheumatic Diseases, 2018, 77, 1558-1565.	0.9	114

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91	Management of major organ involvement of Behçet's syndrome: a systematic review for update of the EULAR recommendations. Rheumatology, 2018, 57, 2200-2212.	1.9	89
92	Molecular Diagnosis Experience in Familial Mediterranean Fever: The Most Frequent Mutations in the MEFV Gene. Haseki Tip Bulteni, 2018, 56, 42-49.	0.3	1
93	FRIO441 Idiopathic inflammatory myopathies: clinical characteristics, survivial and poor prognostic factors of 110 patients from turkey. , 2018, , .		0
94	FRIO675â€Rabiopred, an innovative theranostic tool to assist clinicians select an optimal anti-tnf alpha biological therapy for rheumatoid arthritis patients. , 2018, , .		1
95	SAT0489â€Assesment of persistent organ damage according to imacs (INTERNATIONAL MYOSITIS) Tj ETQq1 1 myositis. , 2018, , .	0.784314	4 rgBT /Overl O
96	SAT0597â€New autoinflammatory phenotype manifesting as hypocomplementemic urticarial vasculitis and associated with homozygous agbl3 variant. , 2018, , .		0
97	Safety and Efficacy of Gevokizumab in Patients with Behçet's Disease Uveitis: Results of an Exploratory Phase 2 Study. Ocular Immunology and Inflammation, 2017, 25, 62-70.	1.8	52
98	Dense genotyping of immune-related loci implicates host responses to microbial exposure in Behçet's disease susceptibility. Nature Genetics, 2017, 49, 438-443.	21.4	129
99	Developing a Core Set of Outcome Measures for Behçet Disease: Report from OMERACT 2016. Journal of Rheumatology, 2017, 44, 1750-1753.	2.0	25
100	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. Journal of Rheumatology, 2017, 44, 619-625.	2.0	41
101	Colchicine resistance and intolerance in familial mediterranean fever: Definition, causes, and alternative treatments. Seminars in Arthritis and Rheumatism, 2017, 47, 115-120.	3.4	108
102	Diagnostic criteria for cryopyrin-associated periodic syndrome (CAPS). Annals of the Rheumatic Diseases, 2017, 76, 942-947.	0.9	175
103	Genetic architecture distinguishes systemic juvenile idiopathic arthritis from other forms of juvenile idiopathic arthritis: clinical and therapeutic implications. Annals of the Rheumatic Diseases, 2017, 76, 906-913.	0.9	123
104	International multi-centre study of pregnancy outcomes with interleukin-1 inhibitors. Rheumatology, 2017, 56, 2102-2108.	1.9	84
105	Development of a Core Set of Outcome Measures for Large-vessel Vasculitis: Report from OMERACT 2016. Journal of Rheumatology, 2017, 44, 1933-1937.	2.0	33
106	Induced-Pluripotent-Stem-Cell-Derived Primitive Macrophages Provide a Platform for Modeling Tissue-Resident Macrophage Differentiation and Function. Immunity, 2017, 47, 183-198.e6.	14.3	245
107	THU0195â€Consistent efficacy and safety of tofacitinib in rheumatoid arthritis patients with inadequate response or intolerance to NON-MTX csdmards. , 2017, , .		1
108	Behçet's disease (or syndrome) –Âan update. Revue De Medecine Interne, 2017, 38, A8-A9.	1.0	0

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109	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. Arthritis Care and Research, 2017, 69, 578-586.	3.4	75
110	OP0063â€Canakinumab treatment in patients with colchicine-resistant FMF (CRFMF), HIDS/MKD and traps: efficacy in the 16 weeks randomised controlled phase and maintenance of disease control and safety at week 40., 2017,,.		0
111	THU0329â€Budd-chiari syndrome in behÇet's disease: a retrospective multicenter study. , 2017, , .		O
112	Remarkable damage along with poor quality of life in Takayasu arteritis: cross-sectional results of a long-term followed-up multicentre cohort. Clinical and Experimental Rheumatology, 2017, 35 Suppl 103, 77-82.	0.8	7
113	Reply to: Behçet's disease: an MHC-l-opathy?. Clinical and Experimental Rheumatology, 2017, 35 Suppl 104, 6.	0.8	0
114	FRIO479 Efficacy of Colchicine and IL-1 Inhibitors in Amyloidosis Associated with Familial Mediterranean Fever: A Retrospective Analysis. Annals of the Rheumatic Diseases, 2016, 75, 611.3-612.	0.9	3
115	AB0433â€Disease Characteristics, Survival Analysis and Mortality in A Single Centre Cohort of 240 Patients with Antiphospholipid Syndrome. Annals of the Rheumatic Diseases, 2016, 75, 1055.2-1055.	0.9	0
116	A single endoplasmic reticulum aminopeptidase-1 protein allotype is a strong risk factor for Behçet's disease in HLA-B*51 carriers. Annals of the Rheumatic Diseases, 2016, 75, 2208-2211.	0.9	59
117	Tuberculosis and other opportunistic infections in tofacitinib-treated patients with rheumatoid arthritis. Annals of the Rheumatic Diseases, 2016, 75, 1133-1138.	0.9	186
118	FRIO502â€Biologic Agents in Refractory Adult Still's Disease: Better Response Rates and Acceptable Safety with Anakinra and Tocilizumab. Annals of the Rheumatic Diseases, 2016, 75, 620.1-620.	0.9	2
119	Approach to the patients with inadequate response to colchicine in familial Mediterranean fever. Best Practice and Research in Clinical Rheumatology, 2016, 30, 296-303.	3.3	33
120	Evaluation of KIR3DL1/KIR3DS1 polymorphism in Behçet's disease. Genes and Immunity, 2016, 17, 396-399.	4.1	15
121	AB0896â€Cogan Syndrome: Differential Response To Biologic Agents and Role of PET-CT in The Increased Diagnosis of Aortitis. Annals of the Rheumatic Diseases, 2016, 75, 1208.3-1209.	0.9	O
122	SAT0295â€Autoimmune Hemolytic Anemia and Thrombocytopenia in A Single Centre Cohort of Patients with Systemic Lupus Erythematosus from Turkey: Clinical Associations and Effect on Disease Damage and Survival. Annals of the Rheumatic Diseases, 2016, 75, 775.1-775.	0.9	0
123	Characteristics Predicting Tuberculosis Risk under Tumor Necrosis Factor-α Inhibitors: Report from a Large Multicenter Cohort with High Background Prevalence. Journal of Rheumatology, 2016, 43, 524-529.	2.0	67
124	Loss-of-function mutations in TNFAIP3 leading to A20 haploinsufficiency cause an early-onset autoinflammatory disease. Nature Genetics, 2016, 48, 67-73.	21.4	513
125	Microscopic colitis in patients with Takayasu's arteritis: a potential association between the two disease entities. Clinical Rheumatology, 2016, 35, 2495-2499.	2.2	7
126	Development and initial validation of international severity scoring system for familial Mediterranean fever (ISSF). Annals of the Rheumatic Diseases, 2016, 75, 1051-1056.	0.9	83

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127	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. Lupus, 2016, 25, 177-184.	1.6	47
128	SAT0343â€Utility of Vascular Findings by PET/CT Scan in The Diagnosis and Activity Assessment of Takayasu Arteritis. Annals of the Rheumatic Diseases, 2016, 75, 791.2-791.	0.9	0
129	Cost of Familial Mediterranean Fever (Fmf) Disease In Turkey. Value in Health, 2015, 18, A666.	0.3	3
130	AB0633â€A Very High Prevalance of Avascular Necrosis in Turkish Patients with Granulomatosis with Polyangiitis (Wegener's): Preliminary Results from a Single Centre: Table 1 Annals of the Rheumatic Diseases, 2015, 74, 1110.3-1111.	0.9	0
131	Generation of integration-free induced pluripotent stem cells from a patient with Familial Mediterranean Fever (FMF). Stem Cell Research, 2015, 15, 694-696.	0.7	14
132	Efficacy and safety of canakinumab in adolescents and adults with colchicine-resistant familial Mediterranean fever. Arthritis Research and Therapy, 2015, 17, 243.	3.5	83
133	THU0561â€Salivary IL-1 Alpha and IL-1 Beta Levels Associated with Oral Mucosal Disease Activity in Behcet's Disease: Table 1 Annals of the Rheumatic Diseases, 2015, 74, 403.3-404.	0.9	0
134	FRIO279â€High Rate and Bimodal Pattern of Severe Infection in a Selected ANCA Associated Vasculitis Cohort:. Annals of the Rheumatic Diseases, 2015, 74, 525.3-526.	0.9	0
135	AB1118â€Validity and Reliability of Medication Adherence Scale in FMF (Adult Version). Annals of the Rheumatic Diseases, 2015, 74, 1274.3-1275.	0.9	0
136	Quality of life changes with canakinumab therapy in adults with colchicine resistant FMF. Pediatric Rheumatology, 2015, 13, .	2.1	0
137	Unified Modeling of Familial Mediterranean Fever and Cryopyrin Associated Periodic Syndromes. Computational and Mathematical Methods in Medicine, 2015, 2015, 1-18.	1.3	11
138	<i>HLA-DRB1*11</i> and variants of the MHC class II locus are strong risk factors for systemic juvenile idiopathic arthritis. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 15970-15975.	7.1	139
139	Reply to Stoimenis et al. European Journal of Human Genetics, 2015, 23, 1280-1280.	2.8	0
140	Pathogenesis of Behçet's disease: autoinflammatory features and beyond. Seminars in Immunopathology, 2015, 37, 413-418.	6.1	107
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