

Norberto Ortego-Centeno

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138
papers

3,725
citations

32
h-index

56
g-index

171
ext. papers

4,411
ext. citations

3.5
avg, IF

4.09
L-index

#	Paper	IF	Citations
138	Genome-wide association study of systemic sclerosis identifies CD247 as a new susceptibility locus. <i>Nature Genetics</i> , 2010 , 42, 426-9	36.3	301
137	Association of a functional single-nucleotide polymorphism of PTPN22, encoding lymphoid protein phosphatase, with rheumatoid arthritis and systemic lupus erythematosus. <i>Arthritis and Rheumatism</i> , 2005 , 52, 219-24		234
136	Transancestral mapping and genetic load in systemic lupus erythematosus. <i>Nature Communications</i> , 2017 , 8, 16021	17.4	171
135	Identification of novel genetic markers associated with clinical phenotypes of systemic sclerosis through a genome-wide association strategy. <i>PLoS Genetics</i> , 2011 , 7, e1002178	6	164
134	ImmunoChip analysis identifies multiple susceptibility loci for systemic sclerosis. <i>American Journal of Human Genetics</i> , 2014 , 94, 47-61	11	151
133	A large-scale genetic analysis reveals a strong contribution of the HLA class II region to giant cell arteritis susceptibility. <i>American Journal of Human Genetics</i> , 2015 , 96, 565-80	11	96
132	A loss-of-function variant of PTPN22 is associated with reduced risk of systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2009 , 18, 569-79	5.6	92
131	Clinical Spectrum Time Course in Anti Jo-1 Positive Antisynthetase Syndrome: Results From an International Retrospective Multicenter Study. <i>Medicine (United States)</i> , 2015 , 94, e1144	1.8	91
130	Somatic NLRP3 mosaicism in Muckle-Wells syndrome. A genetic mechanism shared by different phenotypes of cryopyrin-associated periodic syndromes. <i>Annals of the Rheumatic Diseases</i> , 2015 , 74, 603-10	2.4	83
129	A systemic sclerosis and systemic lupus erythematosus pan-meta-GWAS reveals new shared susceptibility loci. <i>Human Molecular Genetics</i> , 2013 , 22, 4021-9	5.6	81
128	Identification of CSK as a systemic sclerosis genetic risk factor through Genome Wide Association Study follow-up. <i>Human Molecular Genetics</i> , 2012 , 21, 2825-35	5.6	79
127	A rare polymorphism in Toll Like Receptor 2 is associated with systemic sclerosis phenotype and increases production of inflammatory mediators. <i>Journal of Translational Medicine</i> , 2011 , 9,	8.5	78
126	Treatment of therapy-resistant sarcoidosis with adalimumab. <i>Clinical Rheumatology</i> , 2006 , 25, 596-7	3.9	75
125	Registry of the Spanish network for systemic sclerosis: clinical pattern according to cutaneous subsets and immunological status. <i>Seminars in Arthritis and Rheumatism</i> , 2012 , 41, 789-800	5.3	69
124	A replication study confirms the association of TNFSF4 (OX40L) polymorphisms with systemic sclerosis in a large European cohort. <i>Annals of the Rheumatic Diseases</i> , 2011 , 70, 638-41	2.4	61
123	Anti-TNF- α therapy in refractory uveitis associated with sarcoidosis: Multicenter study of 17 patients. <i>Seminars in Arthritis and Rheumatism</i> , 2015 , 45, 361-8	5.3	58
122	Confirmation of TNIP1 but not RHOB and PSORS1C1 as systemic sclerosis risk factors in a large independent replication study. <i>Annals of the Rheumatic Diseases</i> , 2013 , 72, 602-7	2.4	51

121	Serum Jo-1 Autoantibody and Isolated Arthritis in the Antisynthetase Syndrome: Review of the Literature and Report of the Experience of AENEAS Collaborative Group. <i>Clinical Reviews in Allergy and Immunology</i> , 2017 , 52, 71-80	12.3	48
120	Novel identification of the IRF7 region as an anticentromere autoantibody propensity locus in systemic sclerosis. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 114-9	2.4	47
119	Rates of, and risk factors for, severe infections in patients with systemic autoimmune diseases receiving biological agents off-label. <i>Arthritis Research and Therapy</i> , 2011 , 13, R112	5.7	46
118	Identification of a new putative functional IL18 gene variant through an association study in systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2009 , 18, 3739-48	5.6	45
117	Proteomic analysis of plasma from patients with systemic lupus erythematosus: increased presence of haptoglobin alpha2 polypeptide chains over the alpha1 isoforms. <i>Proteomics</i> , 2006 , 6 Suppl 1, S282-92	4.8	45
116	A Genome-wide Association Study Identifies Risk Alleles in Plasminogen and P4HA2 Associated with Giant Cell Arteritis. <i>American Journal of Human Genetics</i> , 2017 , 100, 64-74	11	43
115	Usefulness of adalimumab in the treatment of refractory uveitis associated with juvenile idiopathic arthritis. <i>Mediators of Inflammation</i> , 2013 , 2013, 560632	4.3	42
114	Association of a CD24 gene polymorphism with susceptibility to systemic lupus erythematosus. <i>Arthritis and Rheumatism</i> , 2007 , 56, 3080-6		41
113	A combined large-scale meta-analysis identifies COG6 as a novel shared risk locus for rheumatoid arthritis and systemic lupus erythematosus. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 286-294	2.4	39
112	Clinical follow-up predictors of disease pattern change in anti-Jo1 positive anti-synthetase syndrome: Results from a multicenter, international and retrospective study. <i>Autoimmunity Reviews</i> , 2017 , 16, 253-257	13.6	37
111	HLA and non-HLA genes in Behçet disease: a multicentric study in the Spanish population. <i>Arthritis Research and Therapy</i> , 2013 , 15, R145	5.7	36
110	Brief Report: IRF4 Newly Identified as a Common Susceptibility Locus for Systemic Sclerosis and Rheumatoid Arthritis in a Cross-Disease Meta-Analysis of Genome-Wide Association Studies. <i>Arthritis and Rheumatology</i> , 2016 , 68, 2338-44	9.5	35
109	Vitamin D deficiency in a cohort of patients with systemic scleroderma from the south of Spain. <i>Journal of Rheumatology</i> , 2010 , 37, 1355; author reply 1356	4.1	33
108	Off-label uses of anti-TNF therapy in three frequent disorders: Behçet disease, sarcoidosis, and noninfectious uveitis. <i>Mediators of Inflammation</i> , 2013 , 2013, 286857	4.3	32
107	Tumor necrosis factor-alpha inhibitor treatment for sarcoidosis. <i>Therapeutics and Clinical Risk Management</i> , 2008 , 4, 1305-13	2.9	32
106	Genetic Analysis with the ImmunoChip Platform in Behçet Disease. Identification of Residues Associated in the HLA Class I Region and New Susceptibility Loci. <i>PLoS ONE</i> , 2016 , 11, e0161305	3.7	32
105	A genome-wide association study suggests the HLA Class II region as the major susceptibility locus for IgA vasculitis. <i>Scientific Reports</i> , 2017 , 7, 5088	4.9	31
104	Promoter insertion/deletion in the IRF5 gene is highly associated with susceptibility to systemic lupus erythematosus in distinct populations, but exerts a modest effect on gene expression in peripheral blood mononuclear cells. <i>Journal of Rheumatology</i> , 2010 , 37, 574-8	4.1	31

103	Influence of TYK2 in systemic sclerosis susceptibility: a new locus in the IL-12 pathway. <i>Annals of the Rheumatic Diseases</i> , 2016 , 75, 1521-6	2.4	29
102	Timing of onset affects arthritis presentation pattern in antisynthetase syndrome. <i>Clinical and Experimental Rheumatology</i> , 2018 , 36, 44-49	2.2	29
101	Implication of IL-2/IL-21 region in systemic sclerosis genetic susceptibility. <i>Annals of the Rheumatic Diseases</i> , 2013 , 72, 1233-8	2.4	28
100	Fine mapping and conditional analysis identify a new mutation in the autoimmunity susceptibility gene BLK that leads to reduced half-life of the BLK protein. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 1219-26	2.4	28
99	HLA-DRB1 association with Henoch-Schonlein purpura. <i>Arthritis and Rheumatology</i> , 2015 , 67, 823-827	9.5	27
98	Identification of IL12RB1 as a novel systemic sclerosis susceptibility locus. <i>Arthritis and Rheumatology</i> , 2014 , 66, 3521-3	9.5	27
97	MYO9B gene polymorphisms are associated with autoimmune diseases in Spanish population. <i>Human Immunology</i> , 2007 , 68, 610-5	2.3	26
96	Increased CD38 expression in T cells and circulating anti-CD38 IgG autoantibodies differentially correlate with distinct cytokine profiles and disease activity in systemic lupus erythematosus patients. <i>Cytokine</i> , 2013 , 62, 232-43	4	25
95	Influence of the STAT3 genetic variants in the susceptibility to psoriatic arthritis and Behçet disease. <i>Human Immunology</i> , 2013 , 74, 230-3	2.3	24
94	Epistatic interaction of ERAP1 and HLA-B in Behçet disease: a replication study in the Spanish population. <i>PLoS ONE</i> , 2014 , 9, e102100	3.7	23
93	Mutational profile of rare variants in inflammasome-related genes in Behçet disease: A Next Generation Sequencing approach. <i>Scientific Reports</i> , 2017 , 7, 8453	4.9	22
92	Adalimumab treatment for SAPHO syndrome. <i>Acta Dermato-Venereologica</i> , 2010 , 90, 301-2	2.2	22
91	Evidence of new risk genetic factor to systemic lupus erythematosus: the UBASH3A gene. <i>PLoS ONE</i> , 2013 , 8, e60646	3.7	22
90	Association of haplotypes of the TLR8 locus with susceptibility to Crohn and Behçet diseases. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, S117-22	2.2	22
89	Association of HLA-B*41:02 with Henoch-Schönlein Purpura (IgA Vasculitis) in Spanish individuals irrespective of the HLA-DRB1 status. <i>Arthritis Research and Therapy</i> , 2015 , 17, 102	5.7	21
88	First clinical symptom as a prognostic factor in systemic sclerosis: results of a retrospective nationwide cohort study. <i>Clinical Rheumatology</i> , 2018 , 37, 999-1009	3.9	21
87	PXK locus in systemic lupus erythematosus: fine mapping and functional analysis reveals novel susceptibility gene ABHD6. <i>Annals of the Rheumatic Diseases</i> , 2015 , 74, e14	2.4	20
86	Altered AKT1 and MAPK1 gene expression on peripheral blood mononuclear cells and correlation with T-helper-transcription factors in systemic lupus erythematosus patients. <i>Mediators of Inflammation</i> , 2012 , 2012, 495934	4.3	20

85	Prevalence of exercise pulmonary arterial hypertension in scleroderma. <i>Journal of Rheumatology</i> , 2008 , 35, 1812-6	4.1	19
84	Increased expression and phosphorylation of the two S100A9 isoforms in mononuclear cells from patients with systemic lupus erythematosus: a proteomic signature for circulating low-density granulocytes. <i>Journal of Proteomics</i> , 2012 , 75, 1778-91	3.9	18
83	Influence of CD40 rs1883832 polymorphism in susceptibility to and clinical manifestations of biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2010 , 37, 2076-80	4.1	17
82	Adalimumab therapy for refractory uveitis: a pilot study. <i>Journal of Ocular Pharmacology and Therapeutics</i> , 2008 , 24, 613-4; author reply 614	2.6	17
81	Evidence of association of the NLRP1 gene with giant cell arteritis. <i>Annals of the Rheumatic Diseases</i> , 2013 , 72, 628-30	2.4	16
80	Increased association of CD38 with lipid rafts in T cells from patients with systemic lupus erythematosus and in activated normal T cells. <i>Molecular Immunology</i> , 2006 , 43, 1029-39	4.3	16
79	Tocilizumab as an Adjuvant Therapy for Hemophagocytic Lymphohistiocytosis Associated With Visceral Leishmaniasis. <i>American Journal of Therapeutics</i> , 2016 , 23, e1193-6	1	15
78	Association between -174 interleukin-6 gene polymorphism and biological response to rituximab in several systemic autoimmune diseases. <i>DNA and Cell Biology</i> , 2012 , 31, 1486-91	3.6	15
77	Analysis of the REL polymorphism rs13031237 in autoimmune diseases. <i>Annals of the Rheumatic Diseases</i> , 2011 , 70, 711-2	2.4	15
76	Identification of HAVCR1 gene haplotypes associated with mRNA expression levels and susceptibility to autoimmune diseases. <i>Human Genetics</i> , 2010 , 128, 221-9	6.3	15
75	Development of tuberculosis in a patient treated with infliximab who had received prophylactic therapy with isoniazid. <i>Journal of Rheumatology</i> , 2003 , 30, 1657-8	4.1	15
74	New insights into the genetic component of non-infectious uveitis through an Immunochip strategy. <i>Journal of Medical Genetics</i> , 2017 , 54, 38-46	5.8	14
73	Variants of the IFI16 gene affecting the levels of expression of mRNA are associated with susceptibility to Behçet disease. <i>Journal of Rheumatology</i> , 2015 , 42, 695-701	4.1	14
72	A candidate gene approach identifies an IL33 genetic variant as a novel genetic risk factor for GCA. <i>PLoS ONE</i> , 2014 , 9, e113476	3.7	14
71	Bone mass and vitamin D in patients with systemic sclerosis from two Spanish regions. <i>Clinical and Experimental Rheumatology</i> , 2012 , 30, 905-11	2.2	14
70	GIMAP and Behçet disease: no association in the European population. <i>Annals of the Rheumatic Diseases</i> , 2014 , 73, 1433-4	2.4	13
69	Association of the FCGR3A-158F/V gene polymorphism with the response to rituximab treatment in Spanish systemic autoimmune disease patients. <i>DNA and Cell Biology</i> , 2012 , 31, 1671-7	3.6	13
68	Oral Calcidiol Is More Effective Than Cholecalciferol Supplementation to Reach Adequate 25(OH)D Levels in Patients with Autoimmune Diseases Chronically Treated with Low Doses of Glucocorticoids: A "Real-Life" Study. <i>Journal of Osteoporosis</i> , 2015 , 2015, 729451	2.8	12

67	Analysis of ancestral and functionally relevant CD5 variants in systemic lupus erythematosus patients. <i>PLoS ONE</i> , 2014 , 9, e113090	3.7	12
66	Interleukin 1 beta (IL1 β)rs16944 genetic variant as a genetic marker of severe renal manifestations and renal sequelae in Henoch-Schönlein purpura. <i>Clinical and Experimental Rheumatology</i> , 2016 , 34, S84-8	2.2	12
65	Very early and early systemic sclerosis in the Spanish scleroderma Registry (RESCLE) cohort. <i>Autoimmunity Reviews</i> , 2017 , 16, 796-802	13.6	11
64	IL2/IL21 region polymorphism influences response to rituximab in systemic lupus erythematosus patients. <i>Molecular Biology Reports</i> , 2013 , 40, 4851-6	2.8	11
63	Successful treatment of severe portopulmonary hypertension in a patient with Child C cirrhosis by sildenafil. <i>Liver Transplantation</i> , 2006 , 12, 690-1	4.5	11
62	Hepatobiliary involvement in systemic sclerosis and the cutaneous subsets: Characteristics and survival of patients from the Spanish RESCLE Registry. <i>Seminars in Arthritis and Rheumatism</i> , 2018 , 47, 849-857	5.3	10
61	Analysis of ATP8B4 F436L Missense Variant in a Large Systemic Sclerosis Cohort. <i>Arthritis and Rheumatology</i> , 2017 , 69, 1337-1338	9.5	9
60	Applying the ACR/EULAR Systemic Sclerosis Classification Criteria to the Spanish Scleroderma Registry Cohort. <i>Journal of Rheumatology</i> , 2015 , 42, 2327-31	4.1	9
59	Serosal involvement in IgG4-related disease: report of two cases and review of the literature. <i>Rheumatology International</i> , 2016 , 36, 1033-41	3.6	9
58	Evaluation of the IL2/IL21, IL2RA and IL2RB genetic variants influence on the endogenous non-anterior uveitis genetic predisposition. <i>BMC Medical Genetics</i> , 2013 , 14, 52	2.1	9
57	Analyses of hair and salivary cortisol for evaluating hypothalamic-pituitary-adrenal axis activation in patients with autoimmune disease. <i>Stress</i> , 2017 , 20, 541-548	3	9
56	Functional variants of Fc gamma receptor (FCGR2A) and FCGR3A are not associated with susceptibility to systemic sclerosis in a large European Study (EUSTAR). <i>Journal of Rheumatology</i> , 2010 , 37, 1673-9	4.1	9
55	Role of PTPN22 and CSK gene polymorphisms as predictors of susceptibility and clinical heterogeneity in patients with Henoch-Schönlein purpura (IgA vasculitis). <i>Arthritis Research and Therapy</i> , 2015 , 17, 286	5.7	8
54	Health-related internet use by lupus patients in southern Spain. <i>Clinical Rheumatology</i> , 2014 , 33, 567-73	3.9	8
53	Refractory subacute cutaneous lupus erythematosus responding to a single course of belimumab: a new anti-BLyS human monoclonal antibody. <i>Indian Journal of Dermatology, Venereology and Leprology</i> , 2014 , 80, 477-8	0.8	8
52	Polymorphisms in the interleukin 4, interleukin 13, and corresponding receptor genes are not associated with systemic sclerosis and do not influence gene expression. <i>Journal of Rheumatology</i> , 2012 , 39, 112-8	4.1	8
51	Tongue infarction as first symptom of temporal arteritis. <i>Rheumatology International</i> , 2012 , 32, 799-800	3.6	8
50	Role of the C8orf13-BLK region in biopsy-proven giant cell arteritis. <i>Human Immunology</i> , 2010 , 71, 525-9	2.3	8

49	Influence of antibody profile in clinical features and prognosis in a cohort of Spanish patients with systemic sclerosis. <i>Clinical and Experimental Rheumatology</i> , 2017 , 35 Suppl 106, 98-105	2.2	8
48	Novel association of acid phosphatase locus 1*C allele with systemic lupus erythematosus. <i>Human Immunology</i> , 2012 , 73, 107-10	2.3	7
47	Recurrent telangiectasias on the cheek: angiolutoid sarcoidosis. <i>American Journal of Medicine</i> , 2010 , 123, e7-8	2.4	7
46	Omalizumab as a therapeutic alternative for chronic urticaria. <i>Annals of Allergy, Asthma and Immunology</i> , 2008 , 101, 556	3.2	7
45	Use of adalimumab in poststreptococcal reactive arthritis. <i>Journal of Clinical Rheumatology</i> , 2007 , 13, 176	1.1	7
44	Transient global amnesia in a patient with high and persistent levels of antiphospholipid antibodies. <i>Clinical Rheumatology</i> , 2006 , 25, 407-8	3.9	7
43	Influence of MUC5B gene on antisynthetase syndrome. <i>Scientific Reports</i> , 2020 , 10, 1415	4.9	7
42	Lack of association between IL6 gene and Henoch-Schönlein purpura. <i>Clinical and Experimental Rheumatology</i> , 2014 , 32, S141-2	2.2	7
41	Association between perceived level of stress, clinical characteristics and psychopathological symptoms in women with systemic lupus erythematosus. <i>Clinical and Experimental Rheumatology</i> , 2018 , 36, 434-441	2.2	7
40	Otolaryngologic manifestations of systemic vasculitis. <i>Acta Otorrinolaringológica Española</i> , 2012 , 63, 303-10	0.9	6
39	Lack of association between STAT4 gene polymorphism and biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2009 , 36, 1021-5	4.1	6
38	Alterations in episodic memory in patients with systemic lupus erythematosus. <i>Archives of Clinical Neuropsychology</i> , 2008 , 23, 157-64	2.7	6
37	Lack of association between the protein tyrosine phosphatase non-receptor type 22 R263Q and R620W functional genetic variants and endogenous non-anterior uveitis. <i>Molecular Vision</i> , 2013 , 19, 638-43	2.3	6
36	Changes in the pattern of death of 987 patients with systemic sclerosis from 1990 to 2009 from the nationwide Spanish Scleroderma Registry (RESCLE). <i>Clinical and Experimental Rheumatology</i> , 2017 , 35 Suppl 106, 40-47	2.2	6
35	Lack of association between TRAF1/C5 gene polymorphisms and biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2010 , 37, 131-5	4.1	5
34	Lupus pernio or chilblain lupus?: two different entities. <i>Chest</i> , 2009 , 136, 946-947	5.3	5
33	Effectiveness of mycophenolic acid in refractory pyoderma gangrenosum. <i>Journal of Clinical Rheumatology</i> , 2010 , 16, 346-7	1.1	5
32	Pulmonary Langerhans Histiocytosis: an uncommon cause of interstitial pneumonia in a patient with Sjögren syndrome. <i>Clinical Rheumatology</i> , 2016 , 35, 825-8	3.9	4

31	Specific association of IL17A genetic variants with panuveitis. <i>British Journal of Ophthalmology</i> , 2015 , 99, 566-70	5.5	4
30	The functional polymorphism 844 A>G in FcBI (CD89) does not contribute to systemic sclerosis or rheumatoid arthritis susceptibility. <i>Journal of Rheumatology</i> , 2011 , 38, 446-9	4.1	4
29	Role of the rs6822844 gene polymorphism at the IL2-IL21 region in biopsy-proven giant cell arteritis. <i>Clinical and Experimental Rheumatology</i> , 2011 , 29, S12-6	2.2	4
28	Lack of association of TNFAIP3 and JAK1 with Behçet disease in the European population. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, S36-9	2.2	4
27	Mesenteric Inflammatory Venocclusive Disease in a Patient with Sjögren Syndrome. <i>Case Reports in Medicine</i> , 2014 , 2014, 423420	0.7	3
26	Two functional variants of IRF5 influence the development of macular edema in patients with non-anterior uveitis. <i>PLoS ONE</i> , 2013 , 8, e76777	3.7	3
25	A nonsynonymous functional variant of the ITGAM gene is not involved in biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2011 , 38, 2598-601	4.1	3
24	Use of rituximab in Wegener granulomatosis: comment on the article by Wong. <i>Nephrology Dialysis Transplantation</i> , 2007 , 22, 958-9; author reply 959	4.3	3
23	Sclerostin serum levels in patients with systemic autoimmune diseases. <i>BoneKey Reports</i> , 2016 , 5, 775		3
22	HLA association with the susceptibility to anti-synthetase syndrome. <i>Joint Bone Spine</i> , 2021 , 88, 105115	2.9	3
21	Autoimmune disease-associated CD226 gene variants are not involved in giant cell arteritis susceptibility in the Spanish population. <i>Clinical and Experimental Rheumatology</i> , 2012 , 30, S29-33	2.2	3
20	Influence of psychological stress on headache in patients with systemic lupus erythematosus. <i>Journal of Rheumatology</i> , 2014 , 41, 453-7	4.1	2
19	Association study of BAK1 gene polymorphisms in Spanish rheumatoid arthritis and systemic lupus erythematosus cohorts. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 314-6	2.4	2
18	No evidence of association between common autoimmunity STAT4 and IL23R risk polymorphisms and non-anterior uveitis. <i>PLoS ONE</i> , 2013 , 8, e72892	3.7	2
17	Influence of IL2RA rs2104286 polymorphism in the risk of biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2010 , 37, 2331-3	4.1	2
16	Role of BANK1 gene polymorphisms in biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2010 , 37, 1502-4	4.1	2
15	Long-term evolution of cytophagic histiocytic panniculitis. <i>Journal of Cutaneous Medicine and Surgery</i> , 2010 , 14, 136-40	1.6	2
14	Pulmonary hypertension and exercise echocardiography. <i>European Journal of Echocardiography</i> , 2006 , 7, 261-2; author reply 263		2

13	Etidronate and glucocorticoid induced osteoporosis. <i>Journal of Rheumatology</i> , 2005 , 32, 199-200	4.1	2
12	PTPN22 is not associated with Behçet disease. Study spanning the complete gene region in the Spanish population and meta-analysis of the functional variant R620W. <i>Clinical and Experimental Rheumatology</i> , 2016 , 34, S41-S45	2.2	2
11	Effect of ethnicity on clinical presentation and risk of antiphospholipid syndrome in Roma and Caucasian patients with systemic lupus erythematosus: a multicenter cross-sectional study. <i>International Journal of Rheumatic Diseases</i> , 2018 , 21, 2028-2035	2.3	1
10	Groove sign. <i>European Journal of Internal Medicine</i> , 2016 , 28, e3-4	3.9	1
9	T Cell Large Granular Lymphocyte Leukaemia with Cutaneous Infiltration. <i>Sultan Qaboos University Medical Journal</i> , 2017 , 17, e489-e490	0.9	1
8	Association Between FGF-23 Levels and Risk of Fracture in Women With Systemic Sclerosis. <i>Journal of Clinical Densitometry</i> , 2021 , 24, 362-368	3.5	1
7	Association of CCR5 β 2 and Behçet disease: new data from a case-control study in the Spanish population and meta-analysis. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, S96-100	2.2	1
6	Role of MUC1 rs4072037 polymorphism and serum KL-6 levels in patients with antisynthetase syndrome. <i>Scientific Reports</i> , 2021 , 11, 22574	4.9	0
5	Low-Dose Rituximab for Hemolytic Anemia Retreatment in a Patient With Systemic Lupus Erythematosus. <i>American Journal of Therapeutics</i> , 2018 , 25, e577-e578	1	
4	Eosinophilia, pruritic exanthema and digital necrosis. <i>Medicina Clínica</i> , 2018 , 150, e17	1	
3	Venocclusive disease and systemic sclerosis. <i>Medicina Clínica</i> , 2017 , 149, 320	1	
2	Health-related Internet use by patients with systemic sclerosis and other autoimmune diseases: comment on the article by van der Vaart et al. <i>Arthritis Care and Research</i> , 2014 , 66, 334	4.7	
1	Corticosteroids in preventing severe lupus flares: do all patients have the same risk? Comment on the article by Tseng et al. <i>Arthritis and Rheumatism</i> , 2007 , 56, 2098-9; author reply 2099		