

Norberto Ortego-Centeno

List of Publications by Year in descending order

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139
papers

4,714
citations

109137

35
h-index

118652

62
g-index

171
all docs

171
docs citations

171
times ranked

6671
citing authors

#	ARTICLE	IF	CITATIONS
1	Genome-wide association study of systemic sclerosis identifies CD247 as a new susceptibility locus. <i>Nature Genetics</i> , 2010, 42, 426-429.	9.4	351
2	Transancestral mapping and genetic load in systemic lupus erythematosus. <i>Nature Communications</i> , 2017, 8, 16021.	5.8	314
3	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, 219-224.	6.7	275
4	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.	1.5	201
5	ImmunoChip Analysis Identifies Multiple Susceptibility Loci for Systemic Sclerosis. <i>American Journal of Human Genetics</i> , 2014, 94, 47-61.	2.6	182
6	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, 565-580.	2.6	144
7	Clinical Spectrum Time Course in Anti Jo-1 Positive Antisynthetase Syndrome. <i>Medicine (United States)</i> , 2015, 94, e1144.	0.4	133
8	A loss-of-function variant of PTPN22 is associated with reduced risk of systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2008, 18, 569-579.	1.4	106
9	A systemic sclerosis and systemic lupus erythematosus pan-meta-GWAS reveals new shared susceptibility loci. <i>Human Molecular Genetics</i> , 2013, 22, 4021-4029.	1.4	104
10	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, 603-610.	0.5	104
11	Identification of CSK as a systemic sclerosis genetic risk factor through Genome Wide Association Study follow-up. <i>Human Molecular Genetics</i> , 2012, 21, 2825-2835.	1.4	98
12	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.	1.6	92
13	Treatment of therapy-resistant sarcoidosis with adalimumab. <i>Clinical Rheumatology</i> , 2006, 25, 596-597.	1.0	91
14	Anti-TNF- α therapy in refractory uveitis associated with sarcoidosis: Multicenter study of 17 patients. <i>Seminars in Arthritis and Rheumatism</i> , 2015, 45, 361-368.	1.6	78
15	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.	2.6	78
16	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, 638-641.	0.5	63
17	Novel identification of the IRF7 region as an anticentromere autoantibody propensity locus in systemic sclerosis. <i>Annals of the Rheumatic Diseases</i> , 2012, 71, 114-119.	0.5	62
18	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.	2.9	60

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19	A combined large-scale meta-analysis identifies <i>COG6</i> as a novel shared risk locus for rheumatoid arthritis and systemic lupus erythematosus. <i>Annals of the Rheumatic Diseases</i> , 2017, 76, 286-294.	0.5	58
20	Confirmation of <i>TNIP1</i> but not <i>RHOB</i> and <i>PSORS1C1</i> as systemic sclerosis risk factors in a large independent replication study. <i>Annals of the Rheumatic Diseases</i> , 2013, 72, 602-607.	0.5	56
21	Identification of a new putative functional IL18 gene variant through an association study in systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2009, 18, 3739-3748.	1.4	54
22	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.	1.6	53
23	Proteomic analysis of plasma from patients with systemic lupus erythematosus: Increased presence of haptoglobin 1±2 polypeptide chains over the 1±1 isoforms. <i>Proteomics</i> , 2006, 6, S282-S292.	1.3	51
24	HLA and non-HLA genes in Behçet's disease: a multicentric study in the Spanish population. <i>Arthritis Research and Therapy</i> , 2013, 15, R145.	1.6	50
25	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.	1.1	48
26	Association of a <i>CD24</i> gene polymorphism with susceptibility to systemic lupus erythematosus. <i>Arthritis and Rheumatism</i> , 2007, 56, 3080-3086.	6.7	47
27	Usefulness of Adalimumab in the Treatment of Refractory Uveitis Associated with Juvenile Idiopathic Arthritis. <i>Mediators of Inflammation</i> , 2013, 2013, 1-6.	1.4	47
28	Brief Report: <i>IRF4</i> 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, 2338-2344.	2.9	46
29	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.	2.5	46
30	Off-Label Uses of Anti-TNF Therapy in Three Frequent Disorders: Behçet's Disease, Sarcoidosis, and Noninfectious Uveitis. <i>Mediators of Inflammation</i> , 2013, 2013, 1-10.	1.4	45
31	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.	1.6	44
32	Influence of <i>TYK2</i> in systemic sclerosis susceptibility: a new locus in the IL-12 pathway. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 1521-1526.	0.5	41
33	Tumor necrosis factor-alpha inhibitor treatment for sarcoidosis. <i>Therapeutics and Clinical Risk Management</i> , 2008, Volume 4, 1305-1313.	0.9	39
34	Vitamin D Deficiency in a Cohort of Patients with Systemic Scleroderma from the South of Spain. <i>Journal of Rheumatology</i> , 2010, 37, 1355-1355.	1.0	38
35	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, 232-243.	1.4	37
36	Brief Report: Association of HLA-DRB1*01 With IgA Vasculitis (Henoch-Schönlein). <i>Arthritis and Rheumatology</i> , 2015, 67, 823-827.	2.9	35

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37	MYO9B gene polymorphisms are associated with autoimmune diseases in Spanish population. <i>Human Immunology</i> , 2007, 68, 610-615.	1.2	33
38	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, 1219-1226.	0.5	33
39	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.	1.6	33
40	Promoter Insertion/Deletion in the <i>IRF5</i> 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, 574-578.	1.0	32
41	Influence of the STAT3 genetic variants in the susceptibility to psoriatic arthritis and Behçet's disease. <i>Human Immunology</i> , 2013, 74, 230-233.	1.2	30
42	Implication of <i>IL-2/IL-21</i> region in systemic sclerosis genetic susceptibility. <i>Annals of the Rheumatic Diseases</i> , 2013, 72, 1233-1238.	0.5	30
43	Epistatic Interaction of ERAP1 and HLA-B in Behçet Disease: A Replication Study in the Spanish Population. <i>PLoS ONE</i> , 2014, 9, e102100.	1.1	30
44	Timing of onset affects arthritis presentation pattern in antisynthetase syndrome. <i>Clinical and Experimental Rheumatology</i> , 2018, 36, 44-49.	0.4	30
45	Identification of <i>IL12RB1</i> as a Novel Systemic Sclerosis Susceptibility Locus. <i>Arthritis and Rheumatology</i> , 2014, 66, 3521-3523.	2.9	29
46	Mutational profile of rare variants in inflammasome-related genes in Behçet disease: A Next Generation Sequencing approach. <i>Scientific Reports</i> , 2017, 7, 8453.	1.6	29
47	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.	1.0	27
48	Evidence of New Risk Genetic Factor to Systemic Lupus Erythematosus: The UBASH3A Gene. <i>PLoS ONE</i> , 2013, 8, e60646.	1.1	27
49	Adalimumab Treatment for SAPHO Syndrome. <i>Acta Dermato-Venereologica</i> , 2010, 90, 301-302.	0.6	26
50	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, 1-14.	1.4	26
51	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-e14.	0.5	24
52	Association of haplotypes of the TLR8 locus with susceptibility to Crohn's and Behçet's diseases. <i>Clinical and Experimental Rheumatology</i> , 2015, 33, S117-22.	0.4	24
53	Evidence of association of the <i>NLRP1</i> gene with giant cell arteritis. <i>Annals of the Rheumatic Diseases</i> , 2013, 72, 628-630.	0.5	23
54	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, 1029-1039.	1.0	21

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55	Adalimumab Therapy for Refractory Uveitis: A Pilot Study. <i>Journal of Ocular Pharmacology and Therapeutics</i> , 2008, 24, 613-614.	0.6	21
56	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, 1778-1791.	1.2	21
57	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.	1.0	21
58	Prevalence of exercise pulmonary arterial hypertension in scleroderma. <i>Journal of Rheumatology</i> , 2008, 35, 1812-6.	1.0	21
59	Influence of <i>CD40</i> rs1883832 Polymorphism in Susceptibility to and Clinical Manifestations of Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2010, 37, 2076-2080.	1.0	19
60	Identification of HAVCR1 gene haplotypes associated with mRNA expression levels and susceptibility to autoimmune diseases. <i>Human Genetics</i> , 2010, 128, 221-229.	1.8	18
61	Analysis of the <i>REL</i> polymorphism rs13031237 in autoimmune diseases. <i>Annals of the Rheumatic Diseases</i> , 2011, 70, 711-712.	0.5	18
62	Association of the <i>FCGR3A</i> -158F/V Gene Polymorphism with the Response to Rituximab Treatment in Spanish Systemic Autoimmune Disease Patients. <i>DNA and Cell Biology</i> , 2012, 31, 1671-1677.	0.9	18
63	New insights into the genetic component of non-infectious uveitis through an Immuchip strategy. <i>Journal of Medical Genetics</i> , 2017, 54, 38-46.	1.5	18
64	Association Between α 174 <i>Interleukin-6</i> Gene Polymorphism and Biological Response to Rituximab in Several Systemic Autoimmune Diseases. <i>DNA and Cell Biology</i> , 2012, 31, 1486-1491.	0.9	17
65	GMAP and Behçet disease: no association in the European population: Table 1. <i>Annals of the Rheumatic Diseases</i> , 2014, 73, 1433-1434.	0.5	17
66	Variants of the <i>IFI16</i> Gene Affecting the Levels of Expression of mRNA Are Associated with Susceptibility to Behçet Disease. <i>Journal of Rheumatology</i> , 2015, 42, 695-701.	1.0	17
67	Tocilizumab as an Adjuvant Therapy for Hemophagocytic Lymphohistiocytosis Associated With Visceral Leishmaniasis. <i>American Journal of Therapeutics</i> , 2016, 23, e1193-e1196.	0.5	17
68	A Candidate Gene Approach Identifies an IL33 Genetic Variant as a Novel Genetic Risk Factor for GCA. <i>PLoS ONE</i> , 2014, 9, e113476.	1.1	17
69	Recurrent Telangiectasias on the Cheek: Angiolupoid Sarcoidosis. <i>American Journal of Medicine</i> , 2010, 123, e7-e8.	0.6	16
70	Very early and early systemic sclerosis in the Spanish scleroderma Registry (RESCLE) cohort. <i>Autoimmunity Reviews</i> , 2017, 16, 796-802.	2.5	16
71	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.	1.6	16
72	IL2/IL21 region polymorphism influences response to rituximab in systemic lupus erythematosus patients. <i>Molecular Biology Reports</i> , 2013, 40, 4851-4856.	1.0	15

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73	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.	0.8	15
74	Analysis of Ancestral and Functionally Relevant CD5 Variants in Systemic Lupus Erythematosus Patients. <i>PLoS ONE</i> , 2014, 9, e113090.	1.1	15
75	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.	0.2	14
76	Bone mass and vitamin D in patients with systemic sclerosis from two Spanish regions. <i>Clinical and Experimental Rheumatology</i> , 2012, 30, 905-11.	0.4	14
77	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, 1-7.	0.1	13
78	Applying the ACR/EULAR Systemic Sclerosis Classification Criteria to the Spanish Scleroderma Registry Cohort. <i>Journal of Rheumatology</i> , 2015, 42, 2327-2331.	1.0	13
79	Tongue infarction as first symptom of temporal arteritis. <i>Rheumatology International</i> , 2012, 32, 799-800.	1.5	12
80	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	12
81	Serosal involvement in IgG4-related disease: report of two cases and review of the literature. <i>Rheumatology International</i> , 2016, 36, 1033-1041.	1.5	12
82	Influence of MUC5B gene on antisyndetase syndrome. <i>Scientific Reports</i> , 2020, 10, 1415.	1.6	12
83	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.	0.4	12
84	Transient global amnesia in a patient with high and persistent levels of antiphospholipid antibodies. <i>Clinical Rheumatology</i> , 2006, 25, 407-408.	1.0	11
85	Successful treatment of severe portopulmonary hypertension in a patient with Child C cirrhosis by Sildenafil. <i>Liver Transplantation</i> , 2006, 12, 690-691.	1.3	11
86	Health-related Internet use by lupus patients in southern Spain. <i>Clinical Rheumatology</i> , 2014, 33, 567-573.	1.0	11
87	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.	1.6	11
88	Use of Adalimumab in Poststreptococcal Reactive Arthritis. <i>Journal of Clinical Rheumatology</i> , 2007, 13, 176.	0.5	10
89	OMALIZUMAB AS A THERAPEUTIC ALTERNATIVE FOR CHRONIC URTICARIA. <i>Annals of Allergy, Asthma and Immunology</i> , 2008, 101, 556.	0.5	9
90	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, 1673-1679.	1.0	9

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91	Role of the C8orf13-BLK region in biopsy-proven giant cell arteritis. <i>Human Immunology</i> , 2010, 71, 525-529.	1.2	9
92	Novel association of acid phosphatase locus 1* <i>C</i> allele with systemic lupus erythematosus. <i>Human Immunology</i> , 2012, 73, 107-110.	1.2	9
93	Manifestaciones otorrinolaringol3gicas de las vasculitis sist3micas. <i>Acta Otorrinolaringol3gica Espa±ola</i> , 2012, 63, 303-310.	0.2	9
94	Analysis of <i>ATP8B4</i> F436L Missense Variant in a Large Systemic Sclerosis Cohort. <i>Arthritis and Rheumatology</i> , 2017, 69, 1337-1338.	2.9	9
95	Lack of Association Between <i>STAT4</i> Gene Polymorphism and Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2009, 36, 1021-1025.	1.0	8
96	Lack of Association Between TRAF1/C5 Gene Polymorphisms and Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2010, 37, 131-135.	1.0	8
97	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, 112-118.	1.0	8
98	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.	0.9	8
99	HLA association with the susceptibility to anti-synthetase syndrome. <i>Joint Bone Spine</i> , 2021, 88, 105115.	0.8	8
100	Sclerostin serum levels in patients with systemic autoimmune diseases. <i>BoneKEy Reports</i> , 2016, 5, 775.	2.7	8
101	Lack of association between IL6 gene and Henoch-Sch3nlein purpura. <i>Clinical and Experimental Rheumatology</i> , 2014, 32, S141-2.	0.4	8
102	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.	0.4	8
103	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.	0.4	8
104	Lupus Pernio or Chilblain Lupus?. <i>Chest</i> , 2009, 136, 946-947.	0.4	7
105	Effectiveness of Mycophenolic Acid in Refractory Pyoderma Gangrenosum. <i>Journal of Clinical Rheumatology</i> , 2010, 16, 346-347.	0.5	7
106	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.	1.1	7
107	Alterations in episodic memory in patients with systemic lupus erythematosus†. <i>Archives of Clinical Neuropsychology</i> , 2008, 23, 157-64.	0.3	6
108	Specific association of <i>IL17A</i> genetic variants with panuveitis. <i>British Journal of Ophthalmology</i> , 2015, 99, 566-570.	2.1	6

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109	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.	0.4	6
110	Use of rituximab in Wegener's granulomatosis: comment on the article by Wong. <i>Nephrology Dialysis Transplantation</i> , 2007, 22, 958-959.	0.4	4
111	Role of BANK1 Gene Polymorphisms in Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2010, 37, 1502-1504.	1.0	4
112	The Functional Polymorphism 844 A>G in FcÎ±RI (CD89) Does Not Contribute to Systemic Sclerosis or Rheumatoid Arthritis Susceptibility. <i>Journal of Rheumatology</i> , 2011, 38, 446-449.	1.0	4
113	A Nonsynonymous Functional Variant of the ITCAM Gene Is Not Involved in Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2011, 38, 2598-2601.	1.0	4
114	No Evidence of Association between Common Autoimmunity STAT4 and IL23R Risk Polymorphisms and Non-Anterior Uveitis. <i>PLoS ONE</i> , 2013, 8, e72892.	1.1	4
115	Pulmonary Langerhans Histiocytosis: an uncommon cause of interstitial pneumonia in a patient with Sjögren syndrome. <i>Clinical Rheumatology</i> , 2016, 35, 825-828.	1.0	4
116	Role of MUC1 rs4072037 polymorphism and serum KL-6 levels in patients with antisynthetase syndrome. <i>Scientific Reports</i> , 2021, 11, 22574.	1.6	4
117	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.	0.4	4
118	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.	0.4	4
119	Lack of association of TNFAIP3 and JAK1 with Behçet's disease in the European population. <i>Clinical and Experimental Rheumatology</i> , 2015, 33, S36-9.	0.4	4
120	Pulmonary hypertension and exercise echocardiography. <i>European Journal of Echocardiography</i> , 2006, 7, 261-262.	2.3	3
121	Influence of IL2RA rs2104286 Polymorphism in the Risk of Biopsy-proven Giant Cell Arteritis. <i>Journal of Rheumatology</i> , 2010, 37, 2331-2333.	1.0	3
122	Two Functional Variants of IRF5 Influence the Development of Macular Edema in Patients with Non-Anterior Uveitis. <i>PLoS ONE</i> , 2013, 8, e76777.	1.1	3
123	Mesenteric Inflammatory Venooclusive Disease in a Patient with Sjögren's Syndrome. <i>Case Reports in Medicine</i> , 2014, 2014, 1-3.	0.3	3
124	Etidronate and glucocorticoid induced osteoporosis. <i>Journal of Rheumatology</i> , 2005, 32, 199-200.	1.0	3
125	Long-Term Evolution of Cytophagic Histiocytic Panniculitis. <i>Journal of Cutaneous Medicine and Surgery</i> , 2010, 14, 136-140.	0.6	2
126	Association study of BANK1 gene polymorphisms in Spanish rheumatoid arthritis and systemic lupus erythematosus cohorts. <i>Annals of the Rheumatic Diseases</i> , 2012, 71, 314-316.	0.5	2

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127	Influence of Psychological Stress on Headache in Patients with Systemic Lupus Erythematosus. <i>Journal of Rheumatology</i> , 2014, 41, 453-457.	1.0	2
128	Association Between FGF-23 Levels and Risk of Fracture in Women With Systemic Sclerosis. <i>Journal of Clinical Densitometry</i> , 2021, 24, 362-368.	0.5	2
129	PTPN22 is not associated with Behçet's 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.	0.4	2
130	Groove sign. <i>European Journal of Internal Medicine</i> , 2016, 28, e3-e4.	1.0	1
131	T Cell Large Granular Lymphocyte Leukaemia with Cutaneous Infiltration. <i>Sultan Qaboos University Medical Journal</i> , 2018, 17, 489.	0.3	1
132	Association of CCR5 ^{Δ32} and Behçet's 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.	0.4	1
133	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, 2098-2099.	6.7	0
134	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, .	1.8	0
135	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, 334-334.	1.5	0
136	Enfermedad venooclusiva y esclerosis sistémica. <i>Medicina Clínica</i> , 2017, 149, 320.	0.3	0
137	Low-Dose Rituximab for Hemolytic Anemia Retreatment in a Patient With Systemic Lupus Erythematosus. <i>American Journal of Therapeutics</i> , 2018, 25, e577-e578.	0.5	0
138	Eosinofilia, exantema pruriginoso y necrosis digital. <i>Medicina Clínica</i> , 2018, 150, e17.	0.3	0
139	Heme oxygenase-1 promoter polymorphisms do not influence susceptibility to systemic sclerosis and its clinical phenotypes. <i>Clinical and Experimental Rheumatology</i> , 2013, 31, 186.	0.4	0