Norberto Ortego-Centeno

List of Publications by Year in Descending Order

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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.

138 56 3,725 32 h-index g-index citations papers 4.09 171 4,411 3.5 avg, IF L-index ext. citations ext. papers

#	Paper	IF	Citations
138	Role of MUC1 rs4072037 polymorphism and serum KL-6 levels in patients with antisynthetase syndrome. <i>Scientific Reports</i> , 2021 , 11, 22574	4.9	O
137	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
136	HLA association with the susceptibility to anti-synthetase syndrome. <i>Joint Bone Spine</i> , 2021 , 88, 105115	2.9	3
135	Influence of MUC5B gene on antisynthetase syndrome. Scientific Reports, 2020, 10, 1415	4.9	7
134	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	
133	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
132	Eosinophilia, pruritic exanthema and digital necrosis. <i>Medicina Claica</i> , 2018 , 150, e17	1	
131	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
130	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
129	Timing of onset affects arthritis presentation pattern in antisyntethase syndrome. <i>Clinical and Experimental Rheumatology</i> , 2018 , 36, 44-49	2.2	29
128	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
127	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
126	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
125	Analysis of ATP8B4 F436L Missense Variant in a Large Systemic Sclerosis Cohort. <i>Arthritis and Rheumatology</i> , 2017 , 69, 1337-1338	9.5	9
124	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
123	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
122	Very early and early systemic sclerosis in the Spanish scleroderma Registry (RESCLE) cohort. <i>Autoimmunity Reviews</i> , 2017 , 16, 796-802	13.6	11

(2016-2017)

121	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
120	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
119	Mutational profile of rare variants in inflammasome-related genes in Behlet disease: A Next Generation Sequencing approach. <i>Scientific Reports</i> , 2017 , 7, 8453	4.9	22
118	Transancestral mapping and genetic load in systemic lupus erythematosus. <i>Nature Communications</i> , 2017 , 8, 16021	17.4	171
117	Venooclussive disease and systemic sclerosis. <i>Medicina Claica</i> , 2017 , 149, 320	1	
116	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
115	T Cell Large Granular Lymphocyte Leukaemia with Cutaneous Infiltration. <i>Sultan Qaboos University Medical Journal</i> , 2017 , 17, e489-e490	0.9	1
114	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
113	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
112	Tocilizumab as an Adjuvant Therapy for Hemophagocytic Lymphohistiocytosis Associated With Visceral Leishmaniasis. <i>American Journal of Therapeutics</i> , 2016 , 23, e1193-6	1	15
111	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
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. Arthritis and Rheumatology, 2016 , 68, 2338-44	9.5	35
109	Pulmonary Langerhans Histiocytosis: an uncommon cause of interstitial pneumonia in a patient with Sjigren syndrome. <i>Clinical Rheumatology</i> , 2016 , 35, 825-8	3.9	4
108	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
107	Groove sign. European Journal of Internal Medicine, 2016 , 28, e3-4	3.9	1
106	Sclerostin serum levels in patients with systemic autoimmune diseases. <i>BoneKEy Reports</i> , 2016 , 5, 775		3
105	Genetic Analysis with the Immunochip Platform in Behlet 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
104	Interleukin 1 beta (IL1) rs16944 genetic variant as a genetic marker of severe renal manifestations and renal sequelae in Henoch-Schilein purpura. Clinical and Experimental Rheumatology, 2016, 34, S84-	8 ^{2.2}	12

103	PTPN22 is not associated with Behllt@ 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
102	HLA-DRB1 association with Henoch-Schonlein purpura. <i>Arthritis and Rheumatology</i> , 2015 , 67, 823-827	9.5	27
101	Anti-TNF-Itherapy 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
100	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
99	Variants of the IFI16 gene affecting the levels of expression of mRNA are associated with susceptibility to Beh∄t disease. <i>Journal of Rheumatology</i> , 2015 , 42, 695-701	4.1	14
98	Role of PTPN22 and CSK gene polymorphisms as predictors of susceptibility and clinical heterogeneity in patients with Henoch-Schilein purpura (IgA vasculitis). <i>Arthritis Research and Therapy</i> , 2015 , 17, 286	5.7	8
97	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
96	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
95	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
94	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
93	Association of HLA-B*41:02 with Henoch-Schilein Purpura (IgA Vasculitis) in Spanish individuals irrespective of the HLA-DRB1 status. <i>Arthritis Research and Therapy</i> , 2015 , 17, 102	5.7	21
92	Specific association of IL17A genetic variants with panuveitis. <i>British Journal of Ophthalmology</i> , 2015 , 99, 566-70	5.5	4
91	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
90	Lack of association of TNFAIP3 and JAK1 with Behlet@ disease in the European population. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, S36-9	2.2	4
89	Association of CCR5B2 and BehBt@ 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
88	Association of haplotypes of the TLR8 locus with susceptibility to Crohn@and Beh@t@diseases. Clinical and Experimental Rheumatology, 2015 , 33, S117-22	2.2	22
87	Health-related internet use by lupus patients in southern Spain. Clinical Rheumatology, 2014, 33, 567-73	3.9	8
86	Influence of psychological stress on headache in patients with systemic lupus erythematosus. Journal of Rheumatology, 2014 , 41, 453-7	4.1	2

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85	Immunochip analysis identifies multiple susceptibility loci for systemic sclerosis. <i>American Journal of Human Genetics</i> , 2014 , 94, 47-61	11	151	
84	GIMAP and Behät disease: no association in the European population. <i>Annals of the Rheumatic Diseases</i> , 2014 , 73, 1433-4	2.4	13	
83	Epistatic interaction of ERAP1 and HLA-B in Beh@t disease: a replication study in the Spanish population. <i>PLoS ONE</i> , 2014 , 9, e102100	3.7	23	
82	Refractory subacute cutaneous lupus erythematous 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	
81	Mesenteric Inflammatory Venoocclusive Disease in a Patient with Sjgren@Syndrome. <i>Case Reports in Medicine</i> , 2014 , 2014, 423420	0.7	3	
80	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		
79	Identification of IL12RB1 as a novel systemic sclerosis susceptibility locus. <i>Arthritis and Rheumatology</i> , 2014 , 66, 3521-3	9.5	27	
78	Analysis of ancestral and functionally relevant CD5 variants in systemic lupus erythematosus patients. <i>PLoS ONE</i> , 2014 , 9, e113090	3.7	12	
77	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	
76	Lack of association between IL6 gene and Henoch-Schilein purpura. <i>Clinical and Experimental Rheumatology</i> , 2014 , 32, S141-2	2.2	7	
75	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	
74	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	
73	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	
72	Influence of the STAT3 genetic variants in the susceptibility to psoriatic arthritis and Behcet@ disease. <i>Human Immunology</i> , 2013 , 74, 230-3	2.3	24	
71	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	
70	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	
69	Off-label uses of anti-TNF therapy in three frequent disorders: Beh@t@ disease, sarcoidosis, and noninfectious uveitis. <i>Mediators of Inflammation</i> , 2013 , 2013, 286857	4.3	32	
68	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	

67	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
66	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
65	HLA and non-HLA genes in Behlet odisease: a multicentric study in the Spanish population. <i>Arthritis Research and Therapy</i> , 2013 , 15, R145	5.7	36
64	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
63	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
62	Evidence of new risk genetic factor to systemic lupus erythematosus: the UBASH3A gene. <i>PLoS ONE</i> , 2013 , 8, e60646	3.7	22
61	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	3 -2 43	6
60	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
59	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
58	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
57	Novel association of acid phosphatase locus 1*C allele with systemic lupus erythematosus. <i>Human Immunology</i> , 2012 , 73, 107-10	2.3	7
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, 1778-91	3.9	18
55	Otolaryngologic manifestations of systemic vasculitis. <i>Acta Otorrinolaringol</i> gica Espa@la, 2012 , 63, 303-10	0.9	6
54	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
53	Tongue infarction as first symptom of temporal arteritis. Rheumatology International, 2012, 32, 799-800	3.6	8
52	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
51	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
50	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

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49	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
48	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
47	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
46	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
45	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
44	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
43	Analysis of the REL polymorphism rs13031237 in autoimmune diseases. <i>Annals of the Rheumatic Diseases</i> , 2011 , 70, 711-2	2.4	15
42	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
41	The functional polymorphism 844 A>G in FcRI (CD89) does not contribute to systemic sclerosis or rheumatoid arthritis susceptibility. <i>Journal of Rheumatology</i> , 2011 , 38, 446-9	4.1	4
40	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
39	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
38	Role of the rs6822844 gene polymorphism at the IL2-IL21 region in biopsy-proven giant cell arteritis. Clinical and Experimental Rheumatology, 2011 , 29, S12-6	2.2	4
37	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
36	Vitamin D deficiency in a cohort of patients with systemic scleroderma from the south of Spain. Journal of Rheumatology, 2010 , 37, 1355; author reply 1356	4.1	33
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	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
33	Role of BANK1 gene polymorphisms in biopsy-proven giant cell arteritis. <i>Journal of Rheumatology</i> , 2010 , 37, 1502-4	4.1	2
32	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

31	Adalimumab treatment for SAPHO syndrome. Acta Dermato-Venereologica, 2010, 90, 301-2	2.2	22
30	Long-term evolution of cytophagic histiocytic panniculitis. <i>Journal of Cutaneous Medicine and Surgery</i> , 2010 , 14, 136-40	1.6	2
29	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
28	Role of the C8orf13-BLK region in biopsy-proven giant cell arteritis. <i>Human Immunology</i> , 2010 , 71, 525-	92.3	8
27	Recurrent telangiectasias on the cheek: angiolupoid sarcoidosis. <i>American Journal of Medicine</i> , 2010 , 123, e7-8	2.4	7
26	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
25	Effectiveness of mycophenolic acid in refractory pyoderma gangrenosum. <i>Journal of Clinical Rheumatology</i> , 2010 , 16, 346-7	1.1	5
24	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
23	Lupus pernio or chilblain lupus?: two different entities. <i>Chest</i> , 2009 , 136, 946-947	5.3	5
22	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
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, 3739-48	5.6	45
20	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
19	Alterations in episodic memory in patients with systemic lupus erythematosus. <i>Archives of Clinical Neuropsychology</i> , 2008 , 23, 157-64	2.7	6
18	Omalizumab as a therapeutic alternative for chronic urticaria. <i>Annals of Allergy, Asthma and Immunology</i> , 2008 , 101, 556	3.2	7
17	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
16	Tumor necrosis factor-alpha inhibitor treatment for sarcoidosis. <i>Therapeutics and Clinical Risk Management</i> , 2008 , 4, 1305-13	2.9	32
15	Prevalence of exercise pulmonary arterial hypertension in scleroderma. <i>Journal of Rheumatology</i> , 2008 , 35, 1812-6	4.1	19
14	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		

LIST OF PUBLICATIONS

13	Association of a CD24 gene polymorphism with susceptibility to systemic lupus erythematosus. Arthritis and Rheumatism, 2007 , 56, 3080-6		41	
12	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	
11	Use of adalimumab in poststreptococcal reactive arthritis. <i>Journal of Clinical Rheumatology</i> , 2007 , 13, 176	1.1	7	
10	MYO9B gene polymorphisms are associated with autoimmune diseases in Spanish population. Human Immunology, 2007 , 68, 610-5	2.3	26	
9	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	
8	Pulmonary hypertension and exercise echocardiography. <i>European Journal of Echocardiography</i> , 2006 , 7, 261-2; author reply 263		2	
7	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	
6	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	
5	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	
4	Treatment of therapy-resistant sarcoidosis with adalimumab. Clinical Rheumatology, 2006 , 25, 596-7	3.9	75	
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, 219-24		234	
2	Etidronate and glucocorticoid induced osteoporosis. <i>Journal of Rheumatology</i> , 2005 , 32, 199-200	4.1	2	
1	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	