

# Pilar Nozal

## List of Publications by Year in descending order

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Version: 2024-02-01

22  
papers

385  
citations

840776

11  
h-index

752698

20  
g-index

24  
all docs

24  
docs citations

24  
times ranked

565  
citing authors

#	ARTICLE	IF	CITATIONS
1	Nephritic Factors: An Overview of Classification, Diagnostic Tools and Clinical Associations. <i>Frontiers in Immunology</i> , 2019, 10, 886.	4.8	52
2	Autoantibodies to complement components in C3 glomerulopathy and atypical hemolytic uremic syndrome. <i>Immunology Letters</i> , 2014, 160, 163-171.	2.5	50
3	Complement factor I deficiency: a not so rare immune defect. Characterization of new mutations and the first large gene deletion. <i>Orphanet Journal of Rare Diseases</i> , 2012, 7, 42.	2.7	48
4	Heterogeneity but individual constancy of epitopes, isotypes and avidity of factor H autoantibodies in atypical hemolytic uremic syndrome. <i>Molecular Immunology</i> , 2016, 70, 47-55.	2.2	33
5	Complement as a diagnostic tool in immunopathology. <i>Seminars in Cell and Developmental Biology</i> , 2019, 85, 86-97.	5.0	33
6	Testing the Activity of Complement Convertases in Serum/Plasma for Diagnosis of C4NeF-Mediated C3 Glomerulonephritis. <i>Journal of Clinical Immunology</i> , 2016, 36, 517-527.	3.8	26
7	Anti-factor H antibody affecting factor H cofactor activity in a patient with dense deposit disease. <i>CKJ: Clinical Kidney Journal</i> , 2012, 5, 133-136.	2.9	20
8	An ELISA assay with two monoclonal antibodies allows the estimation of free factor H and identifies patients with acquired deficiency of this complement regulator. <i>Molecular Immunology</i> , 2014, 58, 194-200.	2.2	20
9	Case report: lupus nephritis with autoantibodies to complement alternative pathway proteins and C3 gene mutation. <i>BMC Nephrology</i> , 2015, 16, 40.	1.8	18
10	Complement Genetic Variants and FH Desialylation in <i>S. pneumoniae</i> -Haemolytic Uraemic Syndrome. <i>Frontiers in Immunology</i> , 2021, 12, 641656.	4.8	14
11	The effect of methotrexate versus other disease-modifying anti-rheumatic drugs on serum drug levels and clinical response in patients with rheumatoid arthritis treated with tumor necrosis factor inhibitors. <i>Clinical Rheumatology</i> , 2019, 38, 949-954.	2.2	13
12	Immunological features of patients affected by Barraquer-Simons syndrome. <i>Orphanet Journal of Rare Diseases</i> , 2020, 15, 9.	2.7	11
13	Autoantibodies against alternative complement pathway proteins in renal pathologies. <i>Nefrologia</i> , 2016, 36, 489-495.	0.4	8
14	Autoanticuerpos frente a proteínas de la vía alternativa del complemento en enfermedad renal. <i>Nefrologia</i> , 2016, 36, 489-495.	0.4	8
15	Chilblain-like lesions and COVID-19 infection: A prospective observational study at Spain's ground zero. <i>Journal of the American Academy of Dermatology</i> , 2021, 84, 507-509.	1.2	7
16	Complement Factor D (adipsin) Levels Are Elevated in Acquired Partial Lipodystrophy (Barraquer-Simons syndrome). <i>International Journal of Molecular Sciences</i> , 2021, 22, 6608.	4.1	7
17	Evidence of ongoing complement activation on adipose tissue from an 11-year-old girl with Barraquer-Simons syndrome. <i>Journal of Dermatology</i> , 2020, 47, 1439-1444.	1.2	6
18	Blood Lymphocyte Subsets for Early Identification of Non-Remission to TNF Inhibitors in Rheumatoid Arthritis. <i>Frontiers in Immunology</i> , 2020, 11, 1913.	4.8	5

#	ARTICLE	IF	CITATIONS
19	BAFF predicts immunogenicity in older patients with rheumatoid arthritis treated with TNF inhibitors. Scientific Reports, 2021, 11, 11632.	3.3	5
20	Infliximab concentrations in two non-switching cohorts of patients with inflammatory bowel disease: originator vs. biosimilar. Scientific Reports, 2020, 10, 17099.	3.3	1
21	Reduction in antidrug antibody levels after switching to rituximab in patients with rheumatoid arthritis with prior infliximab or adalimumab secondary failure. Seminars in Arthritis and Rheumatism, 2020, 50, E1-E2.	3.4	0
22	Characterization of hypersensitivity reactions to polysulfone hemodialysis membranes. Annals of Allergy, Asthma and Immunology, 2022, , .	1.0	0