Francisco J Espinosa-Rosales

List of Publications by Year
in descending order

77 papers	4,199 citations	212478 28 h-index	129628 63 g-index
82 all docs	82 docs citations	82 times ranked	6059 citing authors

#	Article	IF	Citations
1	Improved HUMARA for the Detection of X-Linked Agammaglobulinemia Carriers. Genetic Testing and Molecular Biomarkers, 2022, , .	0.3	0
2	Coronavirus disease 2019, allergic diseases, and allergen immunotherapy: Possible favorable mechanisms of interaction. Allergy and Asthma Proceedings, 2021, 42, 187-197.	1.0	13
3	Infections With Enterohepatic Non-H. pylori Helicobacter Species in X-Linked Agammaglobulinemia: Clinical Cases and Review of the Literature. Frontiers in Cellular and Infection Microbiology, 2021, 11, 807136.	1.8	4
4	Genetic, Immunological, and Clinical Features of the First Mexican Cohort of Patients with Chronic Granulomatous Disease. Journal of Clinical Immunology, 2020, 40, 475-493.	2.0	45
5	InmunopatologÃa de la infecciÃ³n por virus SARS-CoV-2. Acta Pediatrica De Mexico, 2020, 41, 42.	0.2	3
6	Changing the Lives of People With Primary Immunodeficiencies (PI) With Early Testing and Diagnosis. Frontiers in Immunology, 2018, 9, 1439.	2.2	24
7	Failing to Make Ends Meet: The Broad Clinical Spectrum of DNA Ligase IV Deficiency. Case Series and Review of the Literature. Frontiers in Pediatrics, 2018, 6, 426.	0.9	31
8	Consenso Mexicano para la prescripciÃ³n de inmunoglobulina G como tratamiento de reemplazo e inmunomodulaciÃ³n. Acta Pediatrica De Mexico, 2018, 39, 134.	0.2	2
9	Proteomics: a tool to develop novel diagnostic methods and unravel molecular mechanisms of pediatric diseases. BoletÃn MÃ©dico Del Hospital Infantil De MÃ©xico, 2017, 74, 233-240.	0.2	1
10	Primary immunodeficiency diseases: Genomic approaches delineate heterogeneous Mendelian disorders. Journal of Allergy and Clinical Immunology, 2017, 139, 232-245.	1.5	261
11	Proteomics: a tool to develop novel diagnostic methods and unravel molecular mechanisms of pediatric diseases. BoletÃn MÃ©dico Del Hospital Infantil De MÃ©xico (English Edition), 2017, 74, 233-240.	0.0	0
12	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	2.2	50
13	Multifocal Recurrent Osteomyelitis and Hemophagocytic Lymphohistiocytosis in a Boy with Partial Dominant IFN-Î³R1 Deficiency: Case Report and Review of the Literature. Frontiers in Pediatrics, 2017, 5, 75.	0.9	24
14	Respuesta a la carta al Editor. Acta Pediatrica De Mexico, 2017, 38, 136.	0.2	1
15	Functional characterization of two new STAT3 mutations associated with hyperâ€IgE syndrome in a Mexican cohort. Clinical Genetics, 2016, 89, 217-221.	1.0	10
16	Variations of B cell subpopulations in peripheral blood of healthy Mexican population according to age: Relevance for diagnosis of primary immunodeficiencies. Allergologia Et Immunopathologia, 2016, 44, 571-579.	1.0	16
17	Clinical and mutational features of X-linked agammaglobulinemia in Mexico. Clinical Immunology, 2016, 165, 38-44.	1.4	16
18	Into Action: Improving Access to Optimum Care for all Primary Immunodeficiency Patients. Journal of Clinical Immunology, 2016, 36, 415-417.	2.0	9

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19	Reply. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 1019-1020.	2.0	0
20	Use of corticosteroids as an alternative to surgical treatment for liver abscesses in chronic granulomatous disease. Pediatric Blood and Cancer, 2016, 63, 2254-2255.	0.8	5
21	International Consensus Document (ICON): Common Variable Immunodeficiency Disorders. Journal of Allergy and Clinical Immunology: in Practice, 2016, 4, 38-59.	2.0	669
22	Mycobacterial disease in patients with chronic granulomatous disease: AÂretrospective analysis of 71 cases. Journal of Allergy and Clinical Immunology, 2016, 138, 241-248.e3.	1.5	106
23	Primary Immunodeficiency Diseases in Aguascalientes, Mexico: Results from an Educational Program. Journal of Clinical Immunology, 2016, 36, 173-178.	2.0	8
24	Clinical Features, Non-Infectious Manifestations and Survival Analysis of 161 Children with Primary Immunodeficiency in Mexico: A Single Center Experience Over two Decades. Journal of Clinical Immunology, 2016, 36, 56-65.	2.0	28
25	Calidad de vida de los pacientes con inmunodeficiencias primarias de anticuerpos. Acta Pediatrica De Mexico, 2016, 37, 17.	0.2	0
26	En acciÃ³n: mejorando el acceso a la atenciÃ³n Ã³ptima para todos los pacientes con inmunodeficiencias primarias Semana mundial de las Inmunodeficiencias Primarias. Acta Pediatrica De Mexico, 2016, 37, 64.	0.2	1
27	En acciÃ³n: para mejorar el acceso a la atenciÃ³n Ã³ptima para todos los pacientes con inmunodeficiencias primarias. Revista Alergia Mexico, 2016, 63, 109-112.	0.9	0
28	Uso de glucocorticoides sistÃ©micos en PediatrÃa: generalidades. Acta Pediatrica De Mexico, 2016, 37, 349.	0.2	1
29	A novel CD40LG deletion causes the hyper-IgM syndrome with normal CD40L expression in a 6-month-old child. Immunologic Research, 2015, 62, 89-94.	1.3	7
30	Clinical and Genotypic Spectrum of Chronic Granulomatous Disease in 71 Latin American Patients: First Report from the LASID Registry. Pediatric Blood and Cancer, 2015, 62, 2101-2107.	0.8	67
31	Current state and future perspectives of the Latin American Society for Immunodeficiencies (LASID). Allergologia Et Immunopathologia, 2015, 43, 493-497.	1.0	14
32	Enfermedad de Kawasaki: cuadro clÃnico, exÃ¡menes de laboratorio y lesiones coronarias. Acta Pediatrica De Mexico, 2015, 36, 314.	0.2	8
33	Abstract 191: Atherosclerosis risk and Carotid Intima-Media Thickness after Kawasaki Disease in Mexican Children. Circulation, 2015, 131, .	1.6	0
34	Abstract 204: Risk factors for development of coronary artery aneurysms in Kawasaki disease in Mexican Children. Circulation, 2015, 131, .	1.6	0
35	Lymphocytes and B-cell abnormalities in patients with common variable immunodeficiency (CVID). Allergologia Et Immunopathologia, 2014, 42, 35-43.	1.0	18
36	Molecular analysis for patients with <scp>IL</scp>â€12 receptor Î²1 deficiency. Clinical Genetics, 2014, 86, 161-166.	1.0	19

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37	First Report of the Hyper-IgM Syndrome Registry of the Latin American Society for Immunodeficiencies: Novel Mutations, Unique Infections, and Outcomes. Journal of Clinical Immunology, 2014, 34, 146-156.	2.0	70
38	Gastric Adenocarcinoma in the Context of X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2014, 34, 134-137.	2.0	22
39	Attending to Warning Signs of Primary Immunodeficiency Diseases Across the Range of Clinical Practice. Journal of Clinical Immunology, 2014, 34, 10-22.	2.0	86
40	BCG vaccination in patients with severe combined immunodeficiency: Complications, risks, and vaccination policies. Journal of Allergy and Clinical Immunology, 2014, 133, 1134-1141.	1.5	212
41	Clinical and immunological features of common variable immunodeficiency in Mexican patients. Allergologia Et Immunopathologia, 2014, 42, 235-240.	1.0	34
42	Guidelines for the use of human immunoglobulin therapy in patients with primary immunodeficiencies in Latin America. Allergologia Et Immunopathologia, 2014, 42, 245-260.	1.0	22
43	Detection of inheritance pattern in thirty-three Mexican males with chronic granulomatous disease through 123 dihydrorhodamine assay. Allergologia Et Immunopathologia, 2014, 42, 580-585.	1.0	6
44	Next Generation Sequencing Reveals Skewing of the T and B Cell Receptor Repertoires in Patients with WiskottÃ¢â‚¬â€œAldrich Syndrome. Frontiers in Immunology, 2014, 5, 340.	2.2	40
45	IL-12RÎ²1 Deficiency: Mutation Update and Description of theIL12RB1Variation Database. Human Mutation, 2013, 34, 1329-1339.	1.1	81
46	Survival of Mexican patients with paediatric-onset systemic lupus erythematosus and abnormal electroencephalogram. Allergologia Et Immunopathologia, 2013, 41, 108-113.	1.0	3
47	Partial IFN-Î³R2 deficiency is due to protein misfolding and can be rescued by inhibitors of glycosylation. Blood, 2013, 122, 2390-2401.	0.6	34
48	Juvenile rheumatoid arthritis and asthma, but not childhood-onset systemic lupus erythematosus are associated with FCRL3 polymorphisms in Mexicans. Molecular Immunology, 2013, 53, 374-378.	1.0	26
49	Clinical and genetic analysis of patients with Xâ€linked hyperâ€<scp>IgM</scp> syndrome. Clinical Genetics, 2013, 83, 585-587.	1.0	12
50	Bruton's tyrosine kinaseâ€”an integral protein of B cell development that also has an essential role in the innate immune system. Journal of Leukocyte Biology, 2013, 95, 243-250.	1.5	85
51	Sporadic progressive mucinous histiocytosis in a Mexican patient. Skinmed, 2013, 11, 175-8.	0.0	6
52	604â€ƒPhenotypic and Functional Analysis of B Cells in Patients with Common Variable Immunodeficiency. World Allergy Organization Journal, 2012, 5, S191.	1.6	0
53	138â€ƒTransmission Pattern and Carriers Identification in Male Patients with Chronic Granulomatous Disease. World Allergy Organization Journal, 2012, 5, S46.	1.6	0
54	Immunogenicity of A 23-Valent Pneumococcal Polysaccharide Vaccine Among Mexican Children. Archives of Medical Research, 2012, 43, 402-405.	1.5	2

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55	Increased Pro-inflammatory Cytokine Production After Lipopolysaccharide Stimulation in Patients with X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2012, 32, 967-974.	2.0	28
56	Advancing the management of primary immunodeficiency diseases in Latin America: Latin American Society for Immunodeficiencies (LASID) Initiatives. Allergologia Et Immunopathologia, 2012, 40, 187-193.	1.0	14
57	Cochrane Review: Immunostimulants for preventing respiratory tract infection in children. Evidence-Based Child Health: A Cochrane Review Journal, 2012, 7, 629-717.	2.0	40
58	Commentary on: â€˜Immunostimulants for preventing respiratory tract infection in childrenâ€™ with a response from the review authors. Evidence-Based Child Health: A Cochrane Review Journal, 2012, 7, 718-720.	2.0	4
59	Consequences of two naturally occurring missense mutations in the structure and function of Bruton agammaglobulinemia tyrosine kinase. IUBMB Life, 2012, 64, 346-353.	1.5	5
60	Advances in primary immunodeficiency diseases in Latin America: epidemiology, research, and perspectives. Annals of the New York Academy of Sciences, 2012, 1250, 62-72.	1.8	34
61	Critical issues and needs in management of primary immunodeficiency diseases in Latin America. Allergologia Et Immunopathologia, 2011, 39, 45-51.	1.0	17
62	Primary immunodeficiency diseases in Latin America: Proceedings of the Second Latin American Society for Immunodeficiencies (LASID) Advisory Board. Allergologia Et Immunopathologia, 2011, 39, 106-110.	1.0	18
63	Gain-of-function human STAT1 mutations impair IL-17 immunity and underlie chronic mucocutaneous candidiasis. Journal of Experimental Medicine, 2011, 208, 1635-1648.	4.2	739
64	Revisiting Human IL-12RÎ²1 Deficiency. Medicine (United States), 2010, 89, 381-402.	0.4	367
65	Bullous lesions as a manifestation of systemic lupus erythematosus in two Mexican teenagers. Pediatric Rheumatology, 2010, 8, 19.	0.9	15
66	Association of TLR7 copy number variation with susceptibility to childhood-onset systemic lupus erythematosus in Mexican population. Annals of the Rheumatic Diseases, 2010, 69, 1861-1865.	0.5	101
67	Tumor necrosis factorâ€“Î± is a common genetic risk factor for asthma, juvenile rheumatoid arthritis, and systemic lupus erythematosus in a Mexican pediatric population. Human Immunology, 2009, 70, 251-256.	1.2	77
68	Characterization of Bruton's tyrosine kinase mutations in Mexican patients with X-linked agammaglobulinemia. Molecular Immunology, 2008, 45, 1094-1098.	1.0	21
69	Association of PDCD1 polymorphisms with childhood-onset systemic lupus erythematosus. European Journal of Human Genetics, 2007, 15, 336-341.	1.4	53
70	Primary Immunodeficiency Diseases in Latin America: The Second Report of the LAGID Registry. Journal of Clinical Immunology, 2007, 27, 101-108.	2.0	119
71	Immunostimulants for preventing respiratory tract infection in children. The Cochrane Library, 2006, , CD004974.	1.5	67
72	Corticosteroid therapy for refractory infections in chronic granulomatous disease: case reports and review of the literature. Annals of Allergy, Asthma and Immunology, 2006, 97, 257-261.	0.5	53

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73	Pediatric Churg-Strauss syndrome in Mexico. Pediatric Pulmonology, 2006, 41, 379-382.	1.0	4
74	Association analysis of the PTPN22 gene in childhood-onset systemic lupus erythematosus in Mexican population. Genes and Immunity, 2006, 7, 693-695.	2.2	36
75	Hyper-IgE syndrome and autoimmunity in Mexican children. Pediatric Nephrology, 2006, 21, 1200-1205.	0.9	24
76	Catastrophic Kawasaki Disease or Juvenile Polyarteritis Nodosa?. Seminars in Arthritis and Rheumatism, 2006, 35, 349-354.	1.6	23
77	Search for poliovirus carriers among people with primary immune deficiency diseases in the United States, Mexico, Brazil, and the United Kingdom. Bulletin of the World Health Organization, 2004, 82, 3-8.	1.5	122