Marco A Yamazaki-Nakashimada

List of Publications by Year
in descending order

91 papers	1,789 citations	331670 21 h-index	302126 39 g-index
99 all docs	99 docs citations	99 times ranked	2970 citing authors

#	Article	IF	Citations
1	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	2.9	278
2	DOCK8 deficiency impairs CD8 T cell survival and function in humans and mice. Journal of Experimental Medicine, 2011, 208, 2305-2320.	8.5	175
3	Hematopoietic stem cell transplantation in 29 patients hemizygous for hypomorphic IKBKG/NEMO mutations. Blood, 2017, 130, 1456-1467.	1.4	95
4	Kawasaki Disease Complicated With Macrophage Activation Syndrome: A Systematic Review. Journal of Pediatric Hematology/Oncology, 2017, 39, 445-451.	0.6	91
5	Kawasaki disease shock syndrome: Unique and severe subtype of Kawasaki disease. Pediatrics International, 2018, 60, 781-790.	0.5	87
6	Clinical manifestations associated with Kawasaki disease shock syndrome in Mexican children. European Journal of Pediatrics, 2013, 172, 337-342.	2.7	67
7	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.	1.0	53
8	Intravenous Immunoglobulin Treatment for Macrophage Activation Syndrome Complicating Chronic Granulomatous Disease. Journal of Clinical Immunology, 2012, 32, 207-211.	3.8	53
9	BCG: a vaccine with multiple faces. Human Vaccines and Immunotherapeutics, 2020, 16, 1841-1850.	3.3	49
10	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12RÎ²1 Deficiency. Journal of Clinical Immunology, 2018, 38, 617-627.	3.8	45
11	Genetic, Immunological, and Clinical Features of the First Mexican Cohort of Patients with Chronic Granulomatous Disease. Journal of Clinical Immunology, 2020, 40, 475-493.	3.8	45
12	COVID-19 in the Context of Inborn Errors of Immunity: a Case Series of 31 Patients from Mexico. Journal of Clinical Immunology, 2021, 41, 1463-1478.	3.8	40
13	Intravenous Immunoglobulin Therapy for Hypocomplementemic Urticarial Vasculitis Associated with Systemic Lupus Erythematosus in a Child. Pediatric Dermatology, 2009, 26, 445-447.	0.9	34
14	Partial IFN-Î³R2 deficiency is due to protein misfolding and can be rescued by inhibitors of glycosylation. Blood, 2013, 122, 2390-2401.	1.4	34
15	Clinical and immunological features of common variable immunodeficiency in Mexican patients. Allergologia Et Immunopathologia, 2014, 42, 235-240.	1.7	34
16	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.	1.9	31
17	Increased Pro-inflammatory Cytokine Production After Lipopolysaccharide Stimulation in Patients with X-linked Agammaglobulinemia. Journal of Clinical Immunology, 2012, 32, 967-974.	3.8	28
18	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.	3.8	28

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19	Lupus eritematoso sistÃ©mico: Â¿es una sola enfermedad?. ReumatologÃa ClÃnica, 2016, 12, 274-281.	0.5	25
20	Expanding the clinical features of autoinflammation and phospholipase Cî²2â€associated antibody deficiency and immune dysregulation by description of a novel patient. Journal of the European Academy of Dermatology and Venereology, 2019, 33, 2334-2339.	2.4	25
21	Hyper-IgE syndrome and autoimmunity in Mexican children. Pediatric Nephrology, 2006, 21, 1200-1205.	1.7	24
22	Catastrophic Kawasaki Disease or Juvenile Polyarteritis Nodosa?. Seminars in Arthritis and Rheumatism, 2006, 35, 349-354.	3.4	23
23	Treatment of Kimura Disease With Intravenous Immunoglobulin. Pediatrics, 2011, 128, e1633-e1635.	2.1	20
24	Intestinal Pseudoobstruction Associated With Eosinophilic Enteritis as the Initial Presentation of Systemic Lupus Erythematosus in Children. Journal of Pediatric Gastroenterology and Nutrition, 2009, 48, 482-486.	1.8	19
25	Chronic Granulomatous Disease Associated with Atypical Kawasaki Disease. Pediatric Cardiology, 2008, 29, 169-171.	1.3	18
26	Low percentages of regulatory T cells in common variable immunodeficiency (CVID) patients with autoimmune diseases and its association with increased numbers of CD4+CD45RO+ T and CD21low B cells. Allergologia Et Immunopathologia, 2019, 47, 457-466.	1.7	18
27	A male infant with COVIDâ€19 in the context of ARPC1B deficiency. Pediatric Allergy and Immunology, 2021, 32, 199-201.	2.6	17
28	Clinical and mutational features of X-linked agammaglobulinemia in Mexico. Clinical Immunology, 2016, 165, 38-44.	3.2	16
29	Hemophagocytic Lymphohistiocytosis as a Complication in Patients with MSMD. Journal of Clinical Immunology, 2016, 36, 420-422.	3.8	14
30	BCG and Kawasaki disease in Mexico and Japan. Human Vaccines and Immunotherapeutics, 2017, 13, 1091-1093.	3.3	14
31	Juvenile Dermatomyositis Triggered by SARS-CoV-2. Pediatric Neurology, 2021, 121, 26-27.	2.1	14
32	Thrombotic microangiopathy involving the gallbladder as an unusual manifestation of systemic lupus erythematosus and antiphospholipid syndrome: Case report and review of the literature. World Journal of Gastroenterology, 2006, 12, 7206.	3.3	14
33	Systemic Autoimmunity in a Patient With CANDLE Syndrome. Journal of Investigational Allergology and Clinical Immunology, 2019, 29, 75-76.	1.3	13
34	Interferon alpha-2B in juvenile hyaline fibromatosis. Clinical and Experimental Dermatology, 2006, 31, 478-479.	1.3	12
35	Autoimmune Thrombocytopenic Purpura in Partial DiGeorge Syndrome. Journal of Pediatric Hematology/Oncology, 2011, 33, 465-466.	0.6	12
36	Successful adjunctive immunoglobulin treatment in patients affected by leukocyte adhesion deficiency type 1 (LAD-1). Immunologic Research, 2015, 61, 260-268.	2.9	12

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37	Novel hypomorphic mutation in IKBKG impairs NEMO-ubiquitylation causing ectodermal dysplasia, immunodeficiency, incontinentia pigmenti, and immune thrombocytopenic purpura. Clinical Immunology, 2015, 160, 163-171.	3.2	11
38	Adenoviralâ€induced rash and mucositis: Expanding the spectrum of reactive infectious mucocutaneous eruption. Pediatric Dermatology, 2021, 38, 306-308.	0.9	11
39	Systemic Lupus Erythematosus: Is It One Disease?. ReumatologÃa ClÃnica (English Edition), 2016, 12, 274-281.	0.3	10
40	IgG levels in Kawasaki disease and its association with clinical outcomes. Clinical Rheumatology, 2019, 38, 749-754.	2.2	10
41	Kawasaki disease mimickers. Pediatrics International, 2021, 63, 880-888.	0.5	10
42	Kawasaki disease and immunodeficiencies in children: case reports and literature review. Rheumatology International, 2019, 39, 1829-1838.	3.0	9
43	Diagnostic and therapeutic caveats in Griscelli syndrome. Scandinavian Journal of Immunology, 2021, 93, e13034.	2.7	9
44	VI nerve palsy after intravenous immunoglobulin in Kawasaki disease. Allergologia Et Immunopathologia, 2014, 42, 82-83.	1.7	8
45	Delayed diagnosis in X-linked agammaglobulinemia and its relationship to the occurrence of mutations in BTK non-kinase domains. Expert Review of Clinical Immunology, 2018, 14, 83-93.	3.0	8
46	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.	2.9	7
47	Fever is not always present in Kawasaki disease. Rheumatology International, 2012, 32, 2953-2954.	3.0	6
48	Successful stem cell transplantation in a child with chronic granulomatous disease associated with contiguous gene deletion syndrome and complicated by macrophage activation syndrome. Clinical Immunology, 2014, 154, 112-115.	3.2	6
49	Detection of inheritance pattern in thirty-three Mexican males with chronic granulomatous disease through 123 dihydrorhodamine assay. Allergologia Et Immunopathologia, 2014, 42, 580-585.	1.7	6
50	Orangeâ€brown chromonychia: A valid sign in Kawasaki disease in children of different ethnicities. International Journal of Rheumatic Diseases, 2019, 22, 1160-1161.	1.9	6
51	Use of corticosteroids as an alternative to surgical treatment for liver abscesses in chronic granulomatous disease. Pediatric Blood and Cancer, 2016, 63, 2254-2255.	1.5	5
52	Clinical Manifestations in Carriers of X-Linked Chronic Granulomatous Disease in Mexico. Journal of Investigational Allergology and Clinical Immunology, 2019, 29, 134-136.	1.3	5
53	B subset cells in patients with chronic granulomatous disease in a Mexican population. Allergologia Et Immunopathologia, 2019, 47, 372-377.	1.7	5
54	Macrophage activation syndrome in two infants with multisystem inflammatory syndrome in children. Pediatric Blood and Cancer, 2021, 68, e29199.	1.5	5

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55	Kawasaki disease shock syndrome in the COVID-19 pandemic. Cardiology in the Young, 2022, 32, 506-507.	0.8	5
56	SÃndrome inflamatorio multisistÃ©mico asociado a COVID-19 en niÃ±os y adolescentes: un llamado al diagnÃ³stico. Revista Chilena De Infectologia, 2020, 37, 199-201.	0.1	5
57	Pediatric Churg-Strauss syndrome in Mexico. Pediatric Pulmonology, 2006, 41, 379-382.	2.0	4
58	Amphotericin B Associated Pulmonary Complications in Chronic Granulomatous Disease Patients. Pediatric Blood and Cancer, 2016, 63, 1871-1872.	1.5	4
59	Subcutaneous immunoglobulin for the treatment of deep morphoea in a child. Clinical and Experimental Dermatology, 2018, 43, 303-305.	1.3	4
60	Kawasaki Disease in Infants in the First 3 Months of Age in a Mexican Population: A Cautionary Tale. Frontiers in Pediatrics, 2020, 8, 397.	1.9	4
61	Kawasaki disease presenting with hoarseness: A multinational study of the REKAMLATINA network. Pediatrics International, 2021, 63, 643-648.	0.5	4
62	Skewed X-inactivation in a Female Carrier with X-linked Chronic Granulomatous Disease. Iranian Journal of Allergy, Asthma and Immunology, 2019, 18, 447-451.	0.4	4
63	Severe congenital neutropenia due to G6PC3 deficiency: Case series of five patients and literature review. Scandinavian Journal of Immunology, 2022, 95, e13136.	2.7	4
64	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.	3.9	4
65	Giant coronary artery aneurysms complicating Kawasaki disease in Mexican children. Cardiology in the Young, 2018, 28, 386-390.	0.8	3
66	A Teenager With Rash and Fever: Juvenile Systemic Lupus Erythematosus or Kawasaki Disease?. Frontiers in Pediatrics, 2020, 8, 149.	1.9	3
67	ILâ€1 receptor antagonist defect (DIRA) in a pediatric patient, receiving adalimumab with good clinical response. International Journal of Dermatology, 2021, 60, 639-640.	1.0	3
68	Multiresistant Kawasaki Disease Complicated With Facial Nerve Palsy, Bilateral Giant Coronary Artery Aneurysms, and Stenosis of the Right Coronary Artery in an Infant. Journal of Clinical Rheumatology, 2020, Publish Ahead of Print, .	0.9	3
69	Use of Infliximab in the Treatment of Macrophage Activation Syndrome Complicating Kawasaki Disease. Journal of Pediatric Hematology/Oncology, 2021, 43, e448-e451.	0.6	3
70	Erythema Multiforme. Journal of Clinical Rheumatology, 2020, 26, e181-e182.	0.9	2
71	Perineal Erythema in Kawasaki Disease and MIS-C. Indian Journal of Pediatrics, 2022, 89, 87-87.	0.8	2
72	Clinical Manifestations, Mutational Analysis, and Immunological Phenotype in Patients with RAG1/2 Mutations: First Cases Series from Mexico and Description of Two Novel Mutations. Journal of Clinical Immunology, 2021, 41, 1291-1302.	3.8	2

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73	Afebrile Kawasaki disease is not a benign form of the disease. Pediatrics International, 2017, 59, 1128-1129.	0.5	1
74	Hair pigment distribution changes after haematopoietic stem cell transplantation in Griscelli syndrome type 2. Journal of the European Academy of Dermatology and Venereology, 2021, 35, e53-e56.	2.4	1
75	Tumor germinal mixto con componentes de disgerminoma y coriocarcinoma de ovario en mujer adolescente con ataxiatelangiectasia. Acta Pediatrica De Mexico, 2015, 36, 464.	0.2	1
76	Bronchiolitis Obliterans With Anti-Epiplakin Antibodies in a Boy With Paraneoplastic Pemphigus. Pediatrics, 2022, , .	2.1	1
77	Kawasaki disease after all. Seminars in Arthritis and Rheumatism, 2016, 45, e24.	3.4	0
78	FRI0570â€…CLINICAL CHARACTERISTICS IN PATIENTS WITH PEDIATRIC SYSTEMIC LUPUS ERYTHEMATOSUSDIAGNOSED BEFORE 6 -YEARS OLD IN A THIRD-LEVEL HOSPITAL IN MEXICO. , 2019, , .		0
79	FRI0577â€…ANTIPHOSPHOLIPID SYNDROME SECONDARY TO PEDIATRIC SYSTEMIC LUPUS ERYTHEMATOSUS. EXPERIENCE IN A THIRD-LEVEL HOSPITAL IN MEXICO CITY. , 2019, , .		0
80	Pulmonary Geotrichosis in Chronic Granulomatous Disease. Journal of Investigational Allergology and Clinical Immunology, 2021, 32, 0.	1.3	0
81	Calidad de vida de los pacientes con inmunodeficiencias primarias de anticuerpos. Acta Pediatrica De Mexico, 2016, 37, 17.	0.2	0
82	Mycobacterial Infection, Ectodermal Dysplasia and Thrombocytopenic Purpura. , 2019, , 777-780.		0
83	Edema of Hands and Hypopigmented Lesions on Her Neck and Cheeks. , 2020, , 97-102.		0
84	Introduction to Autoimmunity, Secondary Immunodeficiency, and Transplantation. , 2020, , 1-15.		0
85	Fever and Cervical Lymphadenopathy. , 2020, , 17-24.		0
86	Fever, Anasarca and Arthralgia. , 2020, , 61-64.		0
87	Malaise, Weight Loss and Intermittent Fever. , 2020, , 53-56.		0
88	Atypical patterns of STAT3 phosphorylation in subpopulations B cells in patients with common variable immunodeficiency. Human Immunology, 2022, , .	2.4	0
89	Amaurosis as an initial presentation of Takayasu arteritis in children. Rheumatology International, 2022, , 1.	3.0	0
90	Periorbital erythema and edema in multisystemic inflammatory syndrome in children, an important diagnostic clue. International Journal of Rheumatic Diseases, 0, , .	1.9	0

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91	Inmunodeficiencia combinada debida a deficiencia de DOCK8. Lo que sabemos hasta ahora. Revista Alergia Mexico, 2022, 69, 31-47.	0.1	0