

# Marco A Yamazaki-Nakashimada

## List of Publications by Year in descending order

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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	Perineal Erythema in Kawasaki Disease and MIS-C. Indian Journal of Pediatrics, 2022, 89, 87-87.	0.8	2
2	Kawasaki disease shock syndrome in the COVID-19 pandemic. Cardiology in the Young, 2022, 32, 506-507.	0.8	5
3	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
4	Bronchiolitis Obliterans With Anti-Epiplakin Antibodies in a Boy With Paraneoplastic Pemphigus. Pediatrics, 2022, , .	2.1	1
5	Atypical patterns of STAT3 phosphorylation in subpopulations B cells in patients with common variable immunodeficiency. Human Immunology, 2022, , .	2.4	0
6	Amaurosis as an initial presentation of Takayasu arteritis in children. Rheumatology International, 2022, , 1.	3.0	0
7	Inmunodeficiencia combinada debida a deficiencia de DOCK8. Lo que sabemos hasta ahora. Revista Alergia Mexico, 2022, 69, 31-47.	0.1	0
8	Kawasaki disease mimickers. Pediatrics International, 2021, 63, 880-888.	0.5	10
9	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
10	A male infant with COVID-19 in the context of ARPC1B deficiency. Pediatric Allergy and Immunology, 2021, 32, 199-201.	2.6	17
11	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
12	Adenoviral-induced rash and mucositis: Expanding the spectrum of reactive infectious mucocutaneous eruption. Pediatric Dermatology, 2021, 38, 306-308.	0.9	11
13	Kawasaki disease presenting with hoarseness: A multinational study of the REKAMLATINA network. Pediatrics International, 2021, 63, 643-648.	0.5	4
14	IL-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
15	Diagnostic and therapeutic caveats in Griscelli syndrome. Scandinavian Journal of Immunology, 2021, 93, e13034.	2.7	9
16	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
17	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
18	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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19	Juvenile Dermatomyositis Triggered by SARS-CoV-2. <i>Pediatric Neurology</i> , 2021, 121, 26-27.	2.1	14
20	Pulmonary Geotrichosis in Chronic Granulomatous Disease. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2021, 32, 0.	1.3	0
21	Use of Infliximab in the Treatment of Macrophage Activation Syndrome Complicating Kawasaki Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, 43, e448-e451.	0.6	3
22	Infections With Enterohepatic Non-H. pylori Helicobacter Species in X-Linked Agammaglobulinemia: Clinical Cases and Review of the Literature. <i>Frontiers in Cellular and Infection Microbiology</i> , 2021, 11, 807136.	3.9	4
23	Erythema Multiforme. <i>Journal of Clinical Rheumatology</i> , 2020, 26, e181-e182.	0.9	2
24	Kawasaki Disease in Infants in the First 3 Months of Age in a Mexican Population: A Cautionary Tale. <i>Frontiers in Pediatrics</i> , 2020, 8, 397.	1.9	4
25	Genetic, Immunological, and Clinical Features of the First Mexican Cohort of Patients with Chronic Granulomatous Disease. <i>Journal of Clinical Immunology</i> , 2020, 40, 475-493.	3.8	45
26	BCG: a vaccine with multiple faces. <i>Human Vaccines and Immunotherapeutics</i> , 2020, 16, 1841-1850.	3.3	49
27	A Teenager With Rash and Fever: Juvenile Systemic Lupus Erythematosus or Kawasaki Disease?. <i>Frontiers in Pediatrics</i> , 2020, 8, 149.	1.9	3
28	Multiresistant Kawasaki Disease Complicated With Facial Nerve Palsy, Bilateral Giant Coronary Artery Aneurysms, and Stenosis of the Right Coronary Artery in an Infant. <i>Journal of Clinical Rheumatology</i> , 2020, Publish Ahead of Print, .	0.9	3
29	Edema of Hands and Hypopigmented Lesions on Her Neck and Cheeks. , 2020, , 97-102.		0
30	Introduction to Autoimmunity, Secondary Immunodeficiency, and Transplantation. , 2020, , 1-15.		0
31	Fever and Cervical Lymphadenopathy. , 2020, , 17-24.		0
32	Fever, Anasarca and Arthralgia. , 2020, , 61-64.		0
33	Malaise, Weight Loss and Intermittent Fever. , 2020, , 53-56.		0
34	Síndrome inflamatorio multisistémico asociado a COVID-19 en niños y adolescentes: un llamado al diagnóstico. <i>Revista Chilena De Infectología</i> , 2020, 37, 199-201.	0.1	5
35	Kawasaki disease and immunodeficiencies in children: case reports and literature review. <i>Rheumatology International</i> , 2019, 39, 1829-1838.	3.0	9
36	Expanding the clinical features of autoinflammation and phospholipase C2-associated antibody deficiency and immune dysregulation by description of a novel patient. <i>Journal of the European Academy of Dermatology and Venereology</i> , 2019, 33, 2334-2339.	2.4	25

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37	Orange-brown chromonychia: A valid sign in Kawasaki disease in children of different ethnicities. <i>International Journal of Rheumatic Diseases</i> , 2019, 22, 1160-1161.	1.9	6
38	Clinical Manifestations in Carriers of X-Linked Chronic Granulomatous Disease in Mexico. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2019, 29, 134-136.	1.3	5
39	Systemic Autoimmunity in a Patient With CANDLE Syndrome. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2019, 29, 75-76.	1.3	13
40	FRI0570-CLINICAL CHARACTERISTICS IN PATIENTS WITH PEDIATRIC SYSTEMIC LUPUS ERYTHEMATOSUS DIAGNOSED BEFORE 6 -YEARS OLD IN A THIRD-LEVEL HOSPITAL IN MEXICO. , 2019, , .		0
41	FRI0577-ANTIPHOSPHOLIPID SYNDROME SECONDARY TO PEDIATRIC SYSTEMIC LUPUS ERYTHEMATOSUS. EXPERIENCE IN A THIRD-LEVEL HOSPITAL IN MEXICO CITY. , 2019, , .		0
42	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. <i>Allergologia Et Immunopathologia</i> , 2019, 47, 457-466.	1.7	18
43	B subset cells in patients with chronic granulomatous disease in a Mexican population. <i>Allergologia Et Immunopathologia</i> , 2019, 47, 372-377.	1.7	5
44	IgG levels in Kawasaki disease and its association with clinical outcomes. <i>Clinical Rheumatology</i> , 2019, 38, 749-754.	2.2	10
45	Skewed X-inactivation in a Female Carrier with X-linked Chronic Granulomatous Disease. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2019, 18, 447-451.	0.4	4
46	Mycobacterial Infection, Ectodermal Dysplasia and Thrombocytopenic Purpura. , 2019, , 777-780.		0
47	Subcutaneous immunoglobulin for the treatment of deep morphea in a child. <i>Clinical and Experimental Dermatology</i> , 2018, 43, 303-305.	1.3	4
48	Giant coronary artery aneurysms complicating Kawasaki disease in Mexican children. <i>Cardiology in the Young</i> , 2018, 28, 386-390.	0.8	3
49	Delayed diagnosis in X-linked agammaglobulinemia and its relationship to the occurrence of mutations in BTK non-kinase domains. <i>Expert Review of Clinical Immunology</i> , 2018, 14, 83-93.	3.0	8
50	A Variety of Alu-Mediated Copy Number Variations Can Underlie IL-12R $\beta$ 1 Deficiency. <i>Journal of Clinical Immunology</i> , 2018, 38, 617-627.	3.8	45
51	Kawasaki disease shock syndrome: Unique and severe subtype of Kawasaki disease. <i>Pediatrics International</i> , 2018, 60, 781-790.	0.5	87
52	Failing to Make Ends Meet: The Broad Clinical Spectrum of DNA Ligase IV Deficiency. Case Series and Review of the Literature. <i>Frontiers in Pediatrics</i> , 2018, 6, 426.	1.9	31
53	Kawasaki Disease Complicated With Macrophage Activation Syndrome: A Systematic Review. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, 445-451.	0.6	91
54	BCG and Kawasaki disease in Mexico and Japan. <i>Human Vaccines and Immunotherapeutics</i> , 2017, 13, 1091-1093.	3.3	14

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55	Afebrile Kawasaki disease is not a benign form of the disease. <i>Pediatrics International</i> , 2017, 59, 1128-1129.	0.5	1
56	Hematopoietic stem cell transplantation in 29 patients hemizygous for hypomorphic IKBKG/NEMO mutations. <i>Blood</i> , 2017, 130, 1456-1467.	1.4	95
57	Clinical and mutational features of X-linked agammaglobulinemia in Mexico. <i>Clinical Immunology</i> , 2016, 165, 38-44.	3.2	16
58	Hemophagocytic Lymphohistiocytosis as a Complication in Patients with MSMD. <i>Journal of Clinical Immunology</i> , 2016, 36, 420-422.	3.8	14
59	Systemic Lupus Erythematosus: Is It One Disease?. <i>ReumatologĀa ClĀnica (English Edition)</i> , 2016, 12, 274-281.	0.3	10
60	Amphotericin B Associated Pulmonary Complications in Chronic Granulomatous Disease Patients. <i>Pediatric Blood and Cancer</i> , 2016, 63, 1871-1872.	1.5	4
61	Use of corticosteroids as an alternative to surgical treatment for liver abscesses in chronic granulomatous disease. <i>Pediatric Blood and Cancer</i> , 2016, 63, 2254-2255.	1.5	5
62	Kawasaki disease after all. <i>Seminars in Arthritis and Rheumatism</i> , 2016, 45, e24.	3.4	0
63	Lupus eritematoso sistĀmico: Āes una sola enfermedad?. <i>ReumatologĀa ClĀnica</i> , 2016, 12, 274-281.	0.5	25
64	Clinical Features, Non-Infectious Manifestations and Survival Analysis of 161 Children with Primary Immunodeficiency in Mexico: A Single Center Experience Over two Decades. <i>Journal of Clinical Immunology</i> , 2016, 36, 56-65.	3.8	28
65	Calidad de vida de los pacientes con inmunodeficiencias primarias de anticuerpos. <i>Acta Pediatrica De Mexico</i> , 2016, 37, 17.	0.2	0
66	Successful adjunctive immunoglobulin treatment in patients affected by leukocyte adhesion deficiency type 1 (LAD-1). <i>Immunologic Research</i> , 2015, 61, 260-268.	2.9	12
67	A novel CD40LG deletion causes the hyper-IgM syndrome with normal CD40L expression in a 6-month-old child. <i>Immunologic Research</i> , 2015, 62, 89-94.	2.9	7
68	Novel hypomorphic mutation in IKBKG impairs NEMO-ubiquitylation causing ectodermal dysplasia, immunodeficiency, incontinentia pigmenti, and immune thrombocytopenic purpura. <i>Clinical Immunology</i> , 2015, 160, 163-171.	3.2	11
69	Tumor germinal mixto con componentes de disgerminoma y coriocarcinoma de ovario en mujer adolescente con ataxiatelangiectasia. <i>Acta Pediatrica De Mexico</i> , 2015, 36, 464.	0.2	1
70	VI nerve palsy after intravenous immunoglobulin in Kawasaki disease. <i>Allergologia Et Immunopathologia</i> , 2014, 42, 82-83.	1.7	8
71	Successful stem cell transplantation in a child with chronic granulomatous disease associated with contiguous gene deletion syndrome and complicated by macrophage activation syndrome. <i>Clinical Immunology</i> , 2014, 154, 112-115.	3.2	6
72	Clinical and immunological features of common variable immunodeficiency in Mexican patients. <i>Allergologia Et Immunopathologia</i> , 2014, 42, 235-240.	1.7	34

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73	Detection of inheritance pattern in thirty-three Mexican males with chronic granulomatous disease through 123 dihydrorhodamine assay. <i>Allergologia Et Immunopathologia</i> , 2014, 42, 580-585.	1.7	6
74	Clinical manifestations associated with Kawasaki disease shock syndrome in Mexican children. <i>European Journal of Pediatrics</i> , 2013, 172, 337-342.	2.7	67
75	Partial IFN- $\beta$ 2 deficiency is due to protein misfolding and can be rescued by inhibitors of glycosylation. <i>Blood</i> , 2013, 122, 2390-2401.	1.4	34
76	Increased Pro-inflammatory Cytokine Production After Lipopolysaccharide Stimulation in Patients with X-linked Agammaglobulinemia. <i>Journal of Clinical Immunology</i> , 2012, 32, 967-974.	3.8	28
77	Fever is not always present in Kawasaki disease. <i>Rheumatology International</i> , 2012, 32, 2953-2954.	3.0	6
78	Intravenous Immunoglobulin Treatment for Macrophage Activation Syndrome Complicating Chronic Granulomatous Disease. <i>Journal of Clinical Immunology</i> , 2012, 32, 207-211.	3.8	53
79	Autoimmune Thrombocytopenic Purpura in Partial DiGeorge Syndrome. <i>Journal of Pediatric Hematology/Oncology</i> , 2011, 33, 465-466.	0.6	12
80	Treatment of Kimura Disease With Intravenous Immunoglobulin. <i>Pediatrics</i> , 2011, 128, e1633-e1635.	2.1	20
81	DOCK8 deficiency impairs CD8 T cell survival and function in humans and mice. <i>Journal of Experimental Medicine</i> , 2011, 208, 2305-2320.	8.5	175
82	Intravenous Immunoglobulin Therapy for Hypocomplementemic Urticarial Vasculitis Associated with Systemic Lupus Erythematosus in a Child. <i>Pediatric Dermatology</i> , 2009, 26, 445-447.	0.9	34
83	Intestinal Pseudoobstruction Associated With Eosinophilic Enteritis as the Initial Presentation of Systemic Lupus Erythematosus in Children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2009, 48, 482-486.	1.8	19
84	Chronic Granulomatous Disease Associated with Atypical Kawasaki Disease. <i>Pediatric Cardiology</i> , 2008, 29, 169-171.	1.3	18
85	Corticosteroid therapy for refractory infections in chronic granulomatous disease: case reports and review of the literature. <i>Annals of Allergy, Asthma and Immunology</i> , 2006, 97, 257-261.	1.0	53
86	Pediatric Churg-Strauss syndrome in Mexico. <i>Pediatric Pulmonology</i> , 2006, 41, 379-382.	2.0	4
87	Interferon alpha-2B in juvenile hyaline fibromatosis. <i>Clinical and Experimental Dermatology</i> , 2006, 31, 478-479.	1.3	12
88	Hyper-IgE syndrome and autoimmunity in Mexican children. <i>Pediatric Nephrology</i> , 2006, 21, 1200-1205.	1.7	24
89	Catastrophic Kawasaki Disease or Juvenile Polyarteritis Nodosa?. <i>Seminars in Arthritis and Rheumatism</i> , 2006, 35, 349-354.	3.4	23
90	Thrombotic microangiopathy involving the gallbladder as an unusual manifestation of systemic lupus erythematosus and antiphospholipid syndrome: Case report and review of the literature. <i>World Journal of Gastroenterology</i> , 2006, 12, 7206.	3.3	14

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91	Periorbital erythema and edema in multisystemic inflammatory syndrome in children, an important diagnostic clue. International Journal of Rheumatic Diseases, 0, , .	1.9	0