## Donato Rigante

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/972773/publications.pdf

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297 papers

9,485 citations

52 h-index 80 g-index

302 all docs 302 docs citations

times ranked

302

7943 citing authors

#	Article	IF	CITATIONS
1	EULAR/PRINTO/PRES criteria for Henoch-Schonlein purpura, childhood polyarteritis nodosa, childhood Wegener granulomatosis and childhood Takayasu arteritis: Ankara 2008. Part II: Final classification criteria. Annals of the Rheumatic Diseases, 2010, 69, 798-806.	0.9	1,073
2	Treatment of autoinflammatory diseases: results from the Eurofever Registry and a literature review. Annals of the Rheumatic Diseases, 2013, 72, 678-685.	0.9	350
3	Evidence-based provisional clinical classification criteria for autoinflammatory periodic fevers. Annals of the Rheumatic Diseases, 2015, 74, 799-805.	0.9	215
4	Pediatric Antiphospholipid Syndrome: Clinical and Immunologic Features of 121 Patients in an International Registry. Pediatrics, 2008, 122, e1100-e1107.	2.1	193
5	The Phenotype and Genotype of Mevalonate Kinase Deficiency: A Series of 114 Cases From the Eurofever Registry. Arthritis and Rheumatology, 2016, 68, 2795-2805.	5.6	168
6	International periodic fever, aphthous stomatitis, pharyngitis, cervical adenitis syndrome cohort: description of distinct phenotypes in 301 patients. Rheumatology, 2014, 53, 1125-1129.	1.9	155
7	Phenotypic variability and disparities in treatment and outcomes of childhood arthritis throughout the world: an observational cohort study. The Lancet Child and Adolescent Health, 2019, 3, 255-263.	5.6	120
8	Anakinra treatment in drug-resistant Behcet's disease: a case series. Clinical Rheumatology, 2015, 34, 1293-1301.	2.2	114
9	Follow-Up and Quality of Life of Patients with Cryopyrin-Associated Periodic Syndromes Treated with Anakinra. Journal of Pediatrics, 2010, 157, 310-315.e1.	1.8	105
10	Performance of Current Guidelines for Diagnosis of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Arthritis and Rheumatology, 2014, 66, 2871-2880.	5.6	101
11	Is there a crossroad between infections, genetics, and Henoch–Schönlein purpura?. Autoimmunity Reviews, 2013, 12, 1016-1021.	5.8	100
12	The cryptic interplay between systemic lupus erythematosus and infections. Autoimmunity Reviews, 2014, 13, 96-102.	5.8	100
13	Clues to detect tumor necrosis factor receptor-associated periodic syndrome (TRAPS) among patients with idiopathic recurrent acute pericarditis: results of a multicentre study. Clinical Research in Cardiology, 2012, 101, 525-531.	3.3	97
14	The role of infection in Kawasaki syndrome. Journal of Infection, 2013, 67, 1-10.	3.3	92
15	Caveats and truths in genetic, clinical, autoimmune and autoinflammatory issues in Blau syndrome and early onset sarcoidosis. Autoimmunity Reviews, 2014, 13, 1220-1229.	5.8	86
16	Infections and systemic lupus erythematosus. European Journal of Clinical Microbiology and Infectious Diseases, 2014, 33, 1467-1475.	2.9	83
17	Predictive factors of renal involvement or relapsing disease in children with Henoch-Schï¿⅓2nlein purpura. Rheumatology International, 2005, 25, 45-48.	3.0	79
18	Interleukin-1 as a Common Denominator from Autoinflammatory to Autoimmune Disorders: Premises, Perils, and Perspectives. Mediators of Inflammation, 2015, 2015, 1-21.	3.0	79

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19	Effectiveness and tuberculosis-related safety profile of interleukin-1 blocking agents in the management of Behçet's disease. Autoimmunity Reviews, 2015, 14, 1-9.	5.8	79
20	Treatment with anakinra in the hyperimmunoglobulinemia D/periodic fever syndrome. Rheumatology International, 2006, 27, 97-100.	3.0	77
21	Diagnosis and classification of relapsing polychondritis. Journal of Autoimmunity, 2014, 48-49, 53-59.	6.5	77
22	Kawasaki disease: guidelines of the Italian Society of Pediatrics, part I - definition, epidemiology, etiopathogenesis, clinical expression and management of the acute phase. Italian Journal of Pediatrics, 2018, 44, 102.	2.6	76
23	Inhibition of Interleukin-1 by Canakinumab as a Successful Mono-Drug Strategy for the Treatment of Refractory BehÃSet's Disease: A Case Series. Dermatology, 2014, 228, 211-214.	2.1	74
24	Development and initial validation of the MS score for diagnosis of macrophage activation syndrome in systemic juvenile idiopathic arthritis. Annals of the Rheumatic Diseases, 2019, 78, 1357-1362.	0.9	74
25	A Snapshot on the On-Label and Off-Label Use of the Interleukin-1 Inhibitors in Italy among Rheumatologists and Pediatric Rheumatologists: A Nationwide Multi-Center Retrospective Observational Study. Frontiers in Pharmacology, 2016, 7, 380.	3.5	72
26	The expanding spectrum of low-penetrance TNFRSF1A gene variants in adults presenting with recurrent inflammatory attacks: Clinical manifestations and long-term follow-up. Seminars in Arthritis and Rheumatism, 2014, 43, 818-823.	3.4	71
27	Biological Treatments in Behçet's Disease: Beyond Anti-TNF Therapy. Mediators of Inflammation, 2014, 2014, 1-14.	3.0	68
28	Development of the autoinflammatory disease damage index (ADDI). Annals of the Rheumatic Diseases, 2017, 76, 821-830.	0.9	68
29	Overview of immune abnormalities in lysosomal storage disorders. Immunology Letters, 2017, 188, 79-85.	2.5	68
30	Monogenic Autoinflammatory Syndromes: State of the Art on Genetic, Clinical, and Therapeutic Issues. International Journal of Rheumatology, 2013, 2013, 1-15.	1.6	67
31	First report of macrophage activation syndrome in hyperimmunoglobulinemia D with periodic fever syndrome. Arthritis and Rheumatism, 2007, 56, 658-661.	6.7	65
32	Biological Treatments: New Weapons in the Management of Monogenic Autoinflammatory Disorders. Mediators of Inflammation, 2013, 2013, 1-16.	3.0	64
33	Critical Overview of the Risk Scoring Systems to Predict Non-Responsiveness to Intravenous Immunoglobulin in Kawasaki Syndrome. International Journal of Molecular Sciences, 2016, 17, 278.	4.1	64
34	A comprehensive comparison between pediatric and adult patients with periodic fever, aphthous stomatitis, pharyngitis, and cervical adenopathy (PFAPA) syndrome. Clinical Rheumatology, 2017, 36, 463-468.	2.2	64
35	Safety profile of the interleukin-1 inhibitors anakinra and canakinumab in real-life clinical practice: a nationwide multicenter retrospective observational study. Clinical Rheumatology, 2018, 37, 2233-2240.	2.2	64
36	Autoinflammatory diseases and cardiovascular manifestations. Annals of Medicine, 2011, 43, 341-346.	3.8	61

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37	Unveiling the Efficacy, Safety, and Tolerability of Anti-Interleukin-1 Treatment in Monogenic and Multifactorial Autoinflammatory Diseases. International Journal of Molecular Sciences, 2019, 20, 1898.	4.1	60
38	A spectrum of inflammation and demyelination in acute disseminated encephalomyelitis (ADEM) of children. Autoimmunity Reviews, 2015, 14, 923-929.	5.8	59
39	Key facts and hot spots on tumor necrosis factor receptorâ€associated periodic syndrome. Clinical Rheumatology, 2014, 33, 1197-1207.	2.2	58
40	Infections and Systemic Lupus Erythematosus: Binding or Sparring Partners?. International Journal of Molecular Sciences, 2015, 16, 17331-17343.	4.1	58
41	Hydrocephalus in CINCA syndrome treated with anakinra. Child's Nervous System, 2006, 22, 334-337.	1.1	57
42	Responsiveness to intravenous immunoglobulins and occurrence of coronary artery abnormalities in a single-center cohort of Italian patients with Kawasaki syndrome. Rheumatology International, 2010, 30, 841-846.	3.0	57
43	Vitamin D levels and effects of vitamin D replacement in children with periodic fever, aphthous stomatitis, pharyngitis, and cervical adenitis (PFAPA) syndrome. International Journal of Pediatric Otorhinolaryngology, 2014, 78, 964-968.	1.0	57
44	A systematic approach to autoinflammatory syndromes: a spelling booklet for the beginner. Expert Review of Clinical Immunology, 2017, 13, 571-597.	3.0	57
45	The fresco of autoinflammatory diseases from the pediatric perspective. Autoimmunity Reviews, 2012, 11, 348-356.	5.8	56
46	Macrophage activation syndrome in the course of monogenic autoinflammatory disorders. Clinical Rheumatology, 2015, 34, 1333-1339.	2.2	56
47	Comparative efficacy between adalimumab and infliximab in the treatment of non-infectious intermediate uveitis, posterior uveitis, and panuveitis: a retrospective observational study of 107 patients. Clinical Rheumatology, 2019, 38, 407-415.	2.2	56
48	Cardiac Structural Involvement in Mucopolysaccharidoses. Cardiology, 2002, 98, 18-20.	1.4	55
49	Helicobacter pylori, Gastrointestinal Symptoms, and Metabolic Control in Young Type 1 Diabetes Mellitus Patients. Pediatrics, 2003, 111, 800-803.	2.1	55
50	Autoimmunity and autoinflammation as the yin and yang of idiopathic recurrent acute pericarditis. Autoimmunity Reviews, 2015, 14, 90-97.	5.8	55
51	Relapsing Polychondritis: an Update on Pathogenesis, Clinical Features, Diagnostic Tools, and Therapeutic Perspectives. Current Rheumatology Reports, 2016, 18, 3.	4.7	55
52	A national cohort study on pediatric Behçet's disease: cross-sectional data from an Italian registry. Pediatric Rheumatology, 2017, 15, 84.	2.1	55
53	From the Mediterranean to the Sea of Japan: The Transcontinental Odyssey of Autoinflammatory Diseases. BioMed Research International, 2013, 2013, 1-8.	1.9	54
54	Impact of obesity on the clinical outcome of rheumatologic patients in biotherapy. Autoimmunity Reviews, 2016, 15, 447-450.	5.8	54

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55	The hereditary autoinflammatory disorders uncovered. Autoimmunity Reviews, 2014, 13, 892-900.	5.8	53
56	Performance of Different Diagnostic Criteria for Familial Mediterranean Fever in Children with Periodic Fevers: Results from a Multicenter International Registry. Journal of Rheumatology, 2016, 43, 154-160.	2.0	52
57	Intra-articular corticosteroids versus intra-articular corticosteroids plus methotrexate in oligoarticular juvenile idiopathic arthritis: a multicentre, prospective, randomised, open-label trial. Lancet, The, 2017, 389, 909-916.	13.7	52
58	Adalimumab effectiveness in Behçet's disease: short and long-term data from a multicenter retrospective observational study. Clinical Rheumatology, 2017, 36, 451-455.	2.2	52
59	Paradoxical mucocutaneous flare in a case of Behçet's disease treated with tocilizumab. Clinical Rheumatology, 2015, 34, 1141-1143.	2.2	51
60	Epigenetic control of the immune system: a lesson from Kabuki syndrome. Immunologic Research, 2016, 64, 345-359.	2.9	51
61	West Syndrome: A Review and Guide for Paediatricians. Clinical Drug Investigation, 2018, 38, 113-124.	2.2	51
62	Development and Initial Validation of the Macrophage Activation Syndrome/Primary Hemophagocytic Lymphohistiocytosis Score, a Diagnostic Tool that Differentiates Primary Hemophagocytic Lymphohistiocytosis from Macrophage Activation Syndrome. Journal of Pediatrics, 2017, 189, 72-78.e3.	1.8	50
63	Diagnosis of PFAPA syndrome applied to a cohort of 17 adults with unexplained recurrent fevers. Clinical and Experimental Rheumatology, 2012, 30, 269-71.	0.8	48
64	The Effect of Bisphenol A on Puberty: A Critical Review of the Medical Literature. International Journal of Environmental Research and Public Health, 2017, 14, 1044.	2.6	47
65	Kawasaki syndrome: an intriguing disease with numerous unsolved dilemmas. Pediatric Rheumatology, 2011, 9, 17.	2.1	46
66	The Presence of Uveitis Is Associated with a Sustained Response to the Interleukin (IL)-1 Inhibitors Anakinra and Canakinumab in Behçet's Disease. Ocular Immunology and Inflammation, 2020, 28, 298-304.	1.8	46
67	Treatment options for periodic fever, aphthous stomatitis, pharyngitis, and cervical adenitis (PFAPA) syndrome in children and adults: a narrative review. Clinical Rheumatology, 2019, 38, 11-17.	2.2	45
68	Cumulative retention rate of adalimumab in patients with Behçet's disease-related uveitis: a four-year follow-up study. British Journal of Ophthalmology, 2018, 102, 637-641.	3.9	44
69	An evolving redefinition of autoimmune encephalitis. Autoimmunity Reviews, 2019, 18, 155-163.	5.8	44
70	The Gut Microbiota-Host Partnership as a Potential Driver of Kawasaki Syndrome. Frontiers in Pediatrics, 2019, 7, 124.	1.9	43
71	The Etiology of Juvenile Idiopathic Arthritis. Clinical Reviews in Allergy and Immunology, 2015, 49, 253-261.	6.5	42
72	A Comprehensive Overview of the Hereditary Periodic Fever Syndromes. Clinical Reviews in Allergy and Immunology, 2018, 54, 446-453.	6.5	42

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73	Basic Characteristics of Adults with Periodic Fever, Aphthous Stomatitis, Pharyngitis, and Adenopathy Syndrome in Comparison with the Typical Pediatric Expression of Disease. Mediators of Inflammation, 2015, 2015, 1-11.	3.0	41
74	Live attenuated MMR/V booster vaccines in children with rheumatic diseases on immunosuppressive therapy are safe: Multicenter, retrospective data collection. Vaccine, 2020, 38, 2198-2201.	3.8	41
75	Autoinflammatory syndromes behind the scenes of recurrent fevers in children. Medical Science Monitor, 2009, 15, RA179-87.	1.1	41
76	Kawasaki disease: guidelines of Italian Society of Pediatrics, part II - treatment of resistant forms and cardiovascular complications, follow-up, lifestyle and prevention of cardiovascular risks. Italian Journal of Pediatrics, 2018, 44, 103.	2.6	40
77	Rapid and Sustained Efficacy of Golimumab in the Treatment of Multirefractory Uveitis Associated with Behçet's Disease. Ocular Immunology and Inflammation, 2019, 27, 58-63.	1.8	40
78	Do children's upper respiratory tract infections benefit from probiotics?. BMC Infectious Diseases, 2014, 14, 194.	2.9	38
79	Anakinra treatment in patients with gout and type 2 diabetes. Clinical Rheumatology, 2015, 34, 981-984.	2.2	38
80	The labyrinth of autoinflammatory disorders: a snapshot on the activity of a third-level center in Italy. Clinical Rheumatology, 2015, 34, 17-28.	2.2	37
81	Severe vitamin D deficiency in patients with Kawasaki disease: a potential role in the risk to develop heart vascular abnormalities?. Clinical Rheumatology, 2016, 35, 1865-1872.	2.2	37
82	Diagnostic Criteria for Adult-Onset Periodic Fever, Aphthous Stomatitis, Pharyngitis, and Cervical Adenitis (PFAPA) Syndrome. Frontiers in Immunology, 2017, 8, 1018.	4.8	37
83	New therapeutic solutions for Behçet's syndrome. Expert Opinion on Investigational Drugs, 2016, 25, 827-840.	4.1	36
84	Long-term efficacy and safety of golimumab in the treatment of multirefractory Behçet's disease. Clinical Rheumatology, 2017, 36, 2063-2069.	2.2	36
85	Free light chains and autoimmunity. Autoimmunity Reviews, 2019, 18, 484-492.	5.8	36
86	Current Advances in the Understanding and Treatment of Mevalonate Kinase Deficiency. International Journal of Immunopathology and Pharmacology, 2014, 27, 491-498.	2.1	35
87	Disease status, reasons for discontinuation and adverse events in 1038 Italian children with juvenile idiopathic arthritis treated with etanercept. Pediatric Rheumatology, 2016, 14, 68.	2.1	35
88	The emerging role of interleukin (IL)-1 in the pathogenesis and treatment of inflammatory and degenerative eye diseases. Clinical Rheumatology, 2017, 36, 2307-2318.	2.2	35
89	The safety of live-attenuated vaccines in patients using IL-1 or IL-6 blockade: an international survey. Pediatric Rheumatology, 2018, 16, 19.	2.1	35
90	Efficacy of adalimumab and infliximab in recalcitrant retinal vasculitis inadequately responsive to other immunomodulatory therapies. Clinical Rheumatology, 2018, 37, 2805-2809.	2.2	35

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91	Long-Term Retention Rate of Anakinra in Adult Onset Still's Disease and Predictive Factors for Treatment Response. Frontiers in Pharmacology, 2019, 10, 296.	3.5	35
92	Clinical and biochemical landmarks in systemic autoinflammatory diseases. Annals of Medicine, 2012, 44, 664-673.	3.8	34
93	Immunological Role of IgG Subclasses. Immunological Investigations, 2021, 50, 427-444.	2.0	34
94	IL-6 blockade in the management of non-infectious uveitis. Clinical Rheumatology, 2017, 36, 1459-1469.	2.2	33
95	Timing of uveitis onset in oligoarticular juvenile idiopathic arthritis (JIA) is the main predictor of severe course uveitis. Acta Ophthalmologica, 2012, 90, 91-95.	1.1	31
96	Debate around infection-dependent hemophagocytic syndrome in paediatrics. BMC Infectious Diseases, 2013, 13, 15.	2.9	31
97	Renal involvement in secondary amyloidosis of Muckle-Wells syndrome: marked improvement of renal function and reduction of proteinuria after therapy with human anti-interleukin- $1^2$ monoclonal antibody canakinumab. Clinical Rheumatology, 2015, 34, 1311-1316.	2.2	31
98	Canakinumab efficacy in refractory adultâ€onset <scp>PFAPA</scp> syndrome. International Journal of Rheumatic Diseases, 2017, 20, 1050-1051.	1.9	31
99	FAMILIAL MEDITERRANEAN FEVER: ASSESSING THE OVERALL CLINICAL IMPACT AND FORMULATING TREATMENT PLANS. Mediterranean Journal of Hematology and Infectious Diseases, 2019, 11, e2019027.	1.3	31
100	Revised recommendations of the Italian Society of Pediatrics about the general management of Kawasaki disease. Italian Journal of Pediatrics, 2021, 47, 16.	2.6	31
101	The autoinflammatory side of recurrent pericarditis: Enlightening the pathogenesis for a more rational treatment. Trends in Cardiovascular Medicine, 2021, 31, 265-274.	4.9	31
102	Rare NLRP12 variants associated with the NLRP12-autoinflammatory disorder phenotype: an Italian case series. Clinical and Experimental Rheumatology, 2013, 31, 155-6.	0.8	31
103	Intravenous immunoglobulins (IVIG) in systemic sclerosis: a challenging yet promising future. Immunologic Research, 2015, 61, 326-337.	2.9	30
104	Prognostic Impact of Atypical Presentation in Pediatric Systemic Lupus Erythematosus: Results from a Multicenter Study. Journal of Pediatrics, 2010, 156, 972-977.	1.8	29
105	Long-lasting uveitis remission and hearing loss recovery after rituximab in Vogt-Koyanagi-Harada disease. Clinical Rheumatology, 2015, 34, 1817-1820.	2.2	29
106	Determinants of bone mineral density, bone mineral content, and body composition in a cohort of healthy children: influence of sex, age, puberty, and physical activity. Rheumatology International, 2012, 32, 2737-2743.	3.0	28
107	Spinal involvement in mucopolysaccharidoses: a review. Child's Nervous System, 2015, 31, 203-212.	1.1	28
108	PFAPA syndrome and Behçet's disease: a comparison of two medical entities based on the clinical interviews performed by three different specialists. Clinical Rheumatology, 2016, 35, 501-505.	2.2	28

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109	THE BROAD-RANGING PANORAMA OF SYSTEMIC AUTOINFLAMMATORY DISORDERS WITH SPECIFIC FOCUS ON ACUTE PAINFUL SYMPTOMS AND HEMATOLOGIC MANIFESTATIONS IN CHILDREN. Mediterranean Journal of Hematology and Infectious Diseases, 2018, 10, e2018067.	1.3	28
110	In silico validation of the Autoinflammatory Disease Damage Index. Annals of the Rheumatic Diseases, 2018, 77, 1599-1605.	0.9	27
111	Long-term response after 6-year treatment with anakinra and onset of focal bone erosion in neonatal-onset multisystem inflammatory disease (NOMID/CINCA). Rheumatology International, 2011, 31, 1661-1664.	3.0	26
112	Interleukin-1 Inhibition in Behçet's disease. Israel Medical Association Journal, 2016, 18, 171-6.	0.1	26
113	Untangling the Web of Systemic Autoinflammatory Diseases. Mediators of Inflammation, 2014, 2014, 1-15.	3.0	25
114	Pediatric autoimmune neuropsychiatric disorders associated with streptococcal infections: an overview. European Journal of Clinical Microbiology and Infectious Diseases, 2014, 33, 2105-2109.	2.9	25
115	Anakinra Drug Retention Rate and Predictive Factors of Long-Term Response in Systemic Juvenile Idiopathic Arthritis and Adult Onset Still Disease. Frontiers in Pharmacology, 2019, 10, 918.	3.5	25
116	Clinical Features at Onset and Genetic Characterization of Pediatric and Adult Patients with TNF-⟨i⟩α⟨/i⟩ Receptor—Associated Periodic Syndrome (TRAPS): A Series of 80 Cases from the AIDA Network. Mediators of Inflammation, 2020, 2020, 1-12.	3.0	24
117	The most recent advances in pathophysiology and management of tumour necrosis factor receptor-associated periodic syndrome (TRAPS): personal experience and literature review. Clinical and Experimental Rheumatology, 2013, 31, 141-9.	0.8	24
118	Helicobacter pylori Eradication Rate and Glycemic Control in Young Patients With Type 1 Diabetes. Journal of Pediatric Gastroenterology and Nutrition, 2004, 38, 422-425.	1.8	23
119	Incomplete Kawasaki syndrome followed by systemic onset-juvenile idiopathic arthritis mimicking Kawasaki syndrome. Rheumatology International, 2010, 30, 535-539.	3.0	23
120	Persistent Spontaneous Pneumothorax in an Adolescent with Marfan's Syndrome and Pulmonary Bullous Dysplasia. Respiration, 2001, 68, 621-624.	2.6	22
121	Efficacy and safety of anakinra in tumor necrosis factor receptor-associated periodic syndrome (TRAPS) complicated by severe renal failure: a report after long-term follow-up and review of the literature. Clinical Rheumatology, 2017, 36, 1687-1690.	2.2	22
122	Evaluation of autoimmune phenomena in patients with pediatric autoimmune neuropsychiatric disorders associated with streptococcal infections (PANDAS). Autoimmunity Reviews, 2014, 13, 1236-1240.	5.8	21
123	The Enigma of Periodic Fever, Aphthous Stomatitis, Pharyngitis and Adenitis Syndrome. Pediatric Infectious Disease Journal, 2014, 33, 650-652.	2.0	21
124	Bilateral dexamethasone intravitreal implant in a young patient with Vogt-Koyanagi-Harada disease and refractory uveitis. Clinical Rheumatology, 2015, 34, 1145-1148.	2.2	21
125	Predictors of sustained clinical response in patients with Behçet's disease-related uveitis treated with infliximab and adalimumab. Clinical Rheumatology, 2018, 37, 1715-1720.	2.2	21
126	Rare missense variants in the ALPK1 gene may predispose to periodic fever, aphthous stomatitis, pharyngitis and adenitis (PFAPA) syndrome. European Journal of Human Genetics, 2019, 27, 1361-1368.	2.8	21

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127	Real-world effectiveness of apremilast in multirefractory mucosal involvement of Behçet's disease. Annals of the Rheumatic Diseases, 2019, 78, 1736-1737.	0.9	21
128	Comparison of Early vs. Delayed Anakinra Treatment in Patients With Adult Onset Still's Disease and Effect on Clinical and Laboratory Outcomes. Frontiers in Medicine, 2020, 7, 42.	2.6	21
129	Demographic, clinical and therapeutic findings in a monocentric cohort of adult patients with suspected PFAPA syndrome. Clinical and Experimental Rheumatology, 2016, 34, 77-81.	0.8	21
130	Longitudinal study of microvascular involvement by nailfold capillaroscopy in children with Henoch–Schönlein purpura. Clinical Rheumatology, 2009, 28, 1101-1105.	2.2	20
131	Autoimmune/inflammatory syndrome induced by adjuvants (ASIA): clues and pitfalls in the pediatric background. Immunologic Research, 2014, 60, 366-375.	2.9	20
132	Working the endless puzzle of hereditary autoinflammatory disorders. Modern Rheumatology, 2014, 24, 381-389.	1.8	20
133	Non-canonical manifestations of familial Mediterranean fever: a changing paradigm. Clinical Rheumatology, 2015, 34, 1503-1511.	2.2	20
134	Tumor necrosis factor receptor-associated periodic syndrome managed with the couple canakinumab-alendronate. Clinical Rheumatology, 2015, 34, 807-809.	2.2	20
135	Long-Term Effectiveness of Secukinumab in Patients with Axial Spondyloarthritis. Mediators of Inflammation, 2020, 2020, 1-5.	3.0	20
136	Musculo-skeletal phenotype of Costello syndrome and cardio-facio-cutaneous syndrome: insights on the functional assessment status. Orphanet Journal of Rare Diseases, 2021, 16, 43.	2.7	20
137	Weekly oral alendronate in mevalonate kinase deficiency. Orphanet Journal of Rare Diseases, 2013, 8, 196.	2.7	19
138	A developing portrait of hereditary periodic fevers in childhood. Expert Opinion on Orphan Drugs, 2018, 6, 47-55.	0.8	19
139	New mosaic tiles in childhood hereditary autoinflammatory disorders. Immunology Letters, 2018, 193, 67-76.	2.5	19
140	Efficacy and safety of certolizumab pegol and golimumab in the treatment of non-infectious uveitis. Clinical and Experimental Rheumatology, 2019, 37, 680-683.	0.8	19
141	Geoepidemiological hints about Streptococcus pyogenes strains in relationship with acute rheumatic fever. Autoimmunity Reviews, 2015, 14, 616-621.	5.8	18
142	Efficacy of anakinra in refractory Behçet's disease sacroiliitis. Clinical and Experimental Rheumatology, 2014, 32, S171.	0.8	18
143	Interleukin-1: Ariadne's Thread in Autoinflammatory and Autoimmune Disorders. Israel Medical Association Journal, 2015, 17, 93-7.	0.1	18
144	Monogenic autoinflammatory syndromes at a dermatological level. Archives of Dermatological Research, 2011, 303, 375-380.	1.9	17

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145	Lung Involvement in Children with Hereditary Autoinflammatory Disorders. International Journal of Molecular Sciences, 2016, 17, 2111.	4.1	17
146	Clinical-Serological Characterization and Treatment Outcome of a Large Cohort of Italian Children with Pediatric Autoimmune Neuropsychiatric Disorder Associated with Streptococcal Infection and Pediatric Acute Neuropsychiatric Syndrome. Journal of Child and Adolescent Psychopharmacology, 2019, 29, 608-614.	1.3	17
147	Hints for Genetic and Clinical Differentiation of Adult-Onset Monogenic Autoinflammatory Diseases. Mediators of Inflammation, 2019, 2019, 1-29.	3.0	17
148	Study of vitamin D status and vitamin D receptor polymorphisms in a cohort of Italian patients with juvenile idiopathic arthritis. Scientific Reports, 2020, 10, 17550.	3 <b>.</b> 3	17
149	- KAWASAKI DISEASE AS THE IMMUNE-MEDIATED ECHO OF A VIRAL INFECTION. Mediterranean Journal of Hematology and Infectious Diseases, 2020, 12, e2020039.	1.3	17
150	Discrimination between incomplete and atypical Kawasaki syndrome versus other febrile diseases in childhood: results from an international registry-based study. Clinical and Experimental Rheumatology, 2012, 30, 799-804.	0.8	17
151	A novel MEN1 frameshift germline mutation in two Italian monozygotic twins. Clinical Chemistry and Laboratory Medicine, 2008, 46, 824-6.	2.3	16
152	Mevalonate kinase genotype in children with recurrent fevers and high serum IgD level. Rheumatology International, 2013, 33, 3039-3042.	3.0	16
153	Update on the Medical Management of Gastrointestinal Behçet's Disease. Mediators of Inflammation, 2017, 2017, 1-11.	3.0	16
154	Cross-Sectional Evaluation of Plasma Vitamin D Levels in a Large Cohort of Italian Patients with Pediatric Autoimmune Neuropsychiatric Disorders Associated with Streptococcal Infections. Journal of Child and Adolescent Psychopharmacology, 2018, 28, 124-129.	1.3	16
155	Post-inflammatory retinal dystrophy in CINCA syndrome. Rheumatology International, 2010, 30, 389-393.	3.0	15
156	Typical and severe tumor necrosis factor receptor–associated periodic syndrome in the absence of mutations in the TNFRSF1A gene: a case series. Rheumatology International, 2012, 32, 4015-4018.	3.0	15
157	Fibroblast growth factor 23 (FGF23) gene polymorphism in children with Kawasaki syndrome (KS) and susceptibility to cardiac abnormalities. Italian Journal of Pediatrics, 2013, 39, 69.	2.6	15
158	The protean ocular involvement in monogenic autoinflammatory diseases: state of the art. Clinical Rheumatology, 2015, 34, 1171-1180.	2.2	15
159	Biological therapies for the treatment of Behçet's disease-related uveitis beyond TNF-alpha blockade: a narrative review. Rheumatology International, 2018, 38, 25-35.	3.0	15
160	Drug Retention Rate and Predictive Factors of Drug Survival for Interleukin-1 Inhibitors in Systemic Juvenile Idiopathic Arthritis. Frontiers in Pharmacology, 2018, 9, 1526.	<b>3.</b> 5	15
161	Children and Adults with PFAPA Syndrome: Similarities and Divergences in a Real-Life Clinical Setting. Advances in Therapy, 2021, 38, 1078-1093.	2.9	15
162	New Potential Weapons for Refractory Scleritis in the Era of Targeted Therapy. Mediators of Inflammation, 2020, 2020, 1-6.	3.0	15

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163	Evolving Frontiers in the Treatment of Periodic Fever, Aphthous Stomatitis, Pharyngitis, Cervical Adenitis (PFAPA) Syndrome. Israel Medical Association Journal, 2017, 19, 444-447.	0.1	15
164	Epidemiological profile of non-infectious uveitis from the rheumatologist's perspective: a survey from two tertiary referral centres in Italy. Clinical and Experimental Rheumatology, 2018, 36, 68-73.	0.8	15
165	Kawasaki syndrome and concurrent Coxsackie virus B3 infection. Rheumatology International, 2012, 32, 4037-4040.	3.0	14
166	Serum amyloid-A in Behçet's disease. Clinical Rheumatology, 2014, 33, 1165-1167.	2.2	14
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