Raffaele Manna

List of Publications by Year
in descending order

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143 papers	3,303 citations	172457 29 h-index	175258 52 g-index
150 all docs	150 docs citations	150 times ranked	3637 citing authors

#	Article	IF	Citations
1	Heterogeneity among patients with tumor necrosis factor receptor-associated periodic syndrome phenotypes. Arthritis and Rheumatism, 2003, 48, 2632-2644.	6.7	173
2	Management and treatment of lactose malabsorption. World Journal of Gastroenterology, 2006, 12, 187.	3.3	159
3	Pharmacological and Clinical Basis of Treatment of Familial Mediterranean Fever (FMF) with Colchicine or Analogues: An Update. Inflammation and Allergy: Drug Targets, 2005, 4, 117-124.	3.1	157
4	Familial Mediterranean fever is no longer a rare disease in Italy. European Journal of Human Genetics, 2003, 11, 50-56.	2.8	115
5	Probiotic Treatment Increases Salivary Counts of Lactobacilli: A Double-Blind, Randomized, Controlled Study. Digestion, 2004, 69, 53-56.	2.3	104
6	Familial Mediterranean Fever: A review for clinical management. Joint Bone Spine, 2009, 76, 227-233.	1.6	99
7	Fibrolamellar Carcinoma of the Liver: The Malignant Counterpart of Focal Nodular Hyperplasia with Oncocytic Change. American Journal of Clinical Pathology, 1984, 81, 521-526.	0.7	98
8	Human papillomavirus vaccine and systemic lupus erythematosus. Clinical Rheumatology, 2013, 32, 1301-1307.	2.2	98
9	Insulin and glucagon concentrations in portal and peripheral veins in patients with hepatic cirrhosis. Diabetologia, 1979, 17, 23-28.	6.3	95
10	Response to Interleukin-1 Inhibitors in 140 Italian Patients with Adult-Onset Stillâ€™s Disease: A Multicentre Retrospective Observational Study. Frontiers in Pharmacology, 2017, 8, 369.	3.5	89
11	Schizophrenic symptoms and SPECT abnormalities in a coeliac patient: regression after a glutenâ€free diet. Journal of Internal Medicine, 1997, 242, 421-423.	6.0	86
12	Familial Mediterranean fever: New phenotypes. Autoimmunity Reviews, 2012, 12, 31-37.	5.8	83
13	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
14	Assessment of precision, concordance, specificity, and sensitivity of islet cell antibody measurement in 41 assays. Diabetologia, 1990, 33, 731-736.	6.3	70
15	Delayed Hypersensitivity to Aminopenicillins Is Related to Major Histocompatibility Complex Genes. Annals of Allergy, Asthma and Immunology, 1998, 80, 433-437.	1.0	69
16	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
17	Autoinflammatory gene mutations in Behcet's disease. Annals of the Rheumatic Diseases, 2007, 66, 832-834.	0.9	63
18	Giant cell arteritis and polymyalgia rheumatica after influenza vaccination: report of 10 cases and review of the literature. Lupus, 2012, 21, 153-157.	1.6	61

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19	Effect of exogenous Î²-galactosidase in patients with lactose malabsorption and intolerance: a crossover double-blind placebo-controlled study. European Journal of Clinical Nutrition, 2005, 59, 489-493.	2.9	54
20	Antiâ€Endothelial Autoantibodies in Patients With Sudden Hearing Loss. Laryngoscope, 1999, 109, 1084-1087.	2.0	52
21	IL-1Î² Biological Treatment of Familial Mediterranean Fever. Clinical Reviews in Allergy and Immunology, 2013, 45, 117-130.	6.5	52
22	Current Therapeutic Options for the Main Monogenic Autoinflammatory Diseases and PFAPA Syndrome: Evidence-Based Approach and Proposal of a Practical Guide. Frontiers in Immunology, 2020, 11, 865.	4.8	48
23	Nicotinamide Increases Câ€peptide Secretion in Patients with Recent Onset Type 1 Diabetes. Diabetic Medicine, 1989, 6, 568-572.	2.3	47
24	Disappearance of antiphospholipid antibodies syndrome after Helicobacter pylori eradication. American Journal of Medicine, 2001, 111, 163-164.	1.5	43
25	Autoimmunity in Sudden Sensorineural Hearing Loss: Possible Role of Anti-endothelial Cell Autoantibodies. Acta Oto-Laryngologica, 2002, 122, 30-33.	0.9	41
26	ABO blood groups and cancer of the pancreas. International Journal of Gastrointestinal Cancer, 1990, 6, 81-8.	0.4	40
27	Clinical Associations of Serum Antiendothelial Cell Antibodies in Patients With Sudden Sensorineural Hearing Loss. Laryngoscope, 2003, 113, 797-801.	2.0	38
28	Diagnostic Criteria for Adult-Onset Periodic Fever, Aphthous Stomatitis, Pharyngitis, and Cervical Adenitis (PFAPA) Syndrome. Frontiers in Immunology, 2017, 8, 1018.	4.8	37
29	EPIDERMAL GROWTH FACTOR, SOMATOSTATIN, AND PSORIASIS. Lancet, The, 1983, 321, 65.	13.7	35
30	Long-Term Retention Rate of Anakinra in Adult Onset Stillâ€™s Disease and Predictive Factors for Treatment Response. Frontiers in Pharmacology, 2019, 10, 296.	3.5	35
31	Identification of an autosomal recessive mode of inheritance in paediatric BehÃ§et's families by segregation analysis. , 2003, 122A, 115-118.		32
32	Right Ventricular Hypertrophy, Systolic Function, and Disease Severity in Anderson-Fabry Disease: An Echocardiographic Study. Journal of the American Society of Echocardiography, 2017, 30, 282-291.	2.8	31
33	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
34	Fecal Calprotectin in First-Degree Relatives of Patients with Ulcerative Colitis. American Journal of Gastroenterology, 2007, 102, 132-136.	0.4	29
35	Fabry disease: polymorphic haplotypes and a novel missense mutation in the GLA gene. Clinical Genetics, 2012, 81, 224-233.	2.0	28
36	Non-specific gastrointestinal features: Could it be Fabry disease?. Digestive and Liver Disease, 2018, 50, 429-437.	0.9	28

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37	Pharmacokinetics of Cyclosporin Microemulsion in Patients with Inflammatory Bowel Disease. Clinical Pharmacokinetics, 2001, 40, 473-483.	3.5	27
38	Efficacy of etanercept in the treatment of a patient with BehÃ§etâ€™s disease. Clinical Rheumatology, 2008, 27, 933-936.	2.2	27
39	Anti-β-Adrenoceptors Autoimmunity Causing `Idiopathic' Arrhythmias and Cardiomyopathy. Circulation Journal, 2012, 76, 1345-1353.	1.6	27
40	Clinical hints to diagnosis of attenuated forms of Mucopolysaccharidoses. Italian Journal of Pediatrics, 2018, 44, 132.	2.6	25
41	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
42	Clinical Features at Onset and Genetic Characterization of Pediatric and Adult Patients with TNF-Î± Receptorâ€”Associated Periodic Syndrome (TRAPS): A Series of 80 Cases from the AIDA Network. Mediators of Inflammation, 2020, 2020, 1-12.	3.0	24
43	Macular Impairment in Fabry Disease: A Morpho-functional Assessment by Swept-Source OCT Angiography and Focal Electroretinography. , 2019, 60, 2667.		23
44	Wegener's granulomatosis: A challenging disease for otorhinolaryngologists. Acta Oto-Laryngologica, 2005, 125, 1105-1110.	0.9	21
45	Parapelvic cysts, a distinguishing feature of renal Fabry disease. Nephrology Dialysis Transplantation, 2018, 33, 318-323.	0.7	21
46	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
47	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
48	Behavior of pancreatic glucagon, insulin, and HGH in liver cirrhosis, after arginine and i.v. glucose. Acta Diabetologica Latina, 1974, 11, 330-339.	0.2	20
49	Home infusion program with enzyme replacement therapy for Fabry disease: The experience of a large Italian collaborative group. Molecular Genetics and Metabolism Reports, 2017, 12, 85-91.	1.1	20
50	Effect of Fenfluramine on Growth Hormone and Prolactin Secretion in Obese Subjects. Hormone Research, 1987, 27, 190-194.	1.8	19
51	Polymyalgia rheumatica in 2011. Best Practice and Research in Clinical Rheumatology, 2012, 26, 91-104.	3.3	19
52	Somatostatin infusion in liver cirrhosis: Glucagon control of glucose homeostasis. Diabetologia, 1980, 18, 187-191.	6.3	18
53	Relapsing polychondritis with severe aortic insufficiency. Clinical Rheumatology, 1985, 4, 474-480.	2.2	17
54	Association of Myasthenia Gravis and Antisynthetase Syndrome: A Case Report. International Journal of Immunopathology and Pharmacology, 2004, 17, 395-399.	2.1	17

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55	GH Secretion in Open-Angle Glaucoma. Ophthalmologica, 1979, 179, 168-172.	1.9	15
56	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
57	Insulin Adsorption to Threeâ€Liter Ethylen Vinyl Acetate Bags during 24â€Hour Infusion. Journal of Parenteral and Enteral Nutrition, 1989, 13, 539-541.	2.6	14
58	Autoinflammatory diseases: a possible cause of thrombosis?. Thrombosis Journal, 2015, 13, 19.	2.1	14
59	Small Intestinal Bacterial Overgrowth Affects the Responsiveness to Colchicine in Familial Mediterranean Fever. Mediators of Inflammation, 2017, 2017, 1-6.	3.0	14
60	Prognostic significance of right ventricular hypertrophy and systolic function in Andersonâ€“Fabry disease. ESC Heart Failure, 2020, 7, 1605-1614.	3.1	14
61	Allogenic bone marrow transplantation: not a treatment yet for familial Mediterranean fever. Blood, 2003, 102, 409-409.	1.4	13
62	Conventional and intravenous immunoglobulin therapy in paediatric antiphospholipid antibodies-related chorea. Lupus, 2014, 23, 1449-1451.	1.6	13
63	Functional and pharmacological evaluation of novel GLA variants in Fabry disease identifies six (two) Tj ETQq1 1 0.784314 rgBT /Over 2018, 481, 25-33.	1.1	13
64	Coronary Artery Aneurysms Presenting as Acute Coronary Syndrome: An Unusual Case of IgG4-Related Disease Vascular Involvement. Canadian Journal of Cardiology, 2018, 34, 1088.e7-1088.e10.	1.7	13
65	The everchanging framework of autoinflammation. Internal and Emergency Medicine, 2021, 16, 1759-1770.	2.0	13
66	Y688X, the first nonsense mutation in familial Mediterranean fever (FMF). Human Mutation, 2001, 17, 79-79.	2.5	11
67	Systemic Inflammatory Diseases and Silicone Breast Prostheses: Report of a Case of Adult Still Disease and Review of the Literature. American Journal of the Medical Sciences, 2004, 327, 102-104.	1.1	11
68	Chronic active hepatitis and BehÃ§et's syndrome. Clinical Rheumatology, 1985, 4, 93-96.	2.2	10
69	Right ventricular strain in Anderson-Fabry disease. International Journal of Cardiology, 2021, 330, 84-90.	1.7	10
70	Psychiatric Manifestations as a Primary Symptom in Antiphospholipid Syndrome. International Journal of Immunopathology and Pharmacology, 2006, 19, 915-917.	2.1	9
71	Sjogren's Syndrome in a Celiac Patient: Searching for Environmental Triggers. International Journal of Immunopathology and Pharmacology, 2006, 19, 445-448.	2.1	9
72	Association between Familial Mediterranean Fever and Retroperitoneal Fibrosis: Retroperitoneal Fibrosis Regression after Colchicine Therapy. International Journal of Immunopathology and Pharmacology, 2009, 22, 521-524.	2.1	9

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73	The impact of fever/hyperthermia in the diagnosis of Fabry: A retrospective analysis. European Journal of Internal Medicine, 2016, 32, 26-30.	2.2	9
74	FETAL ORIGIN OF AMNIOTIC FLUID INSULIN IN THE HUMAN MOTHER. Clinical Endocrinology, 1980, 12, 67-70.	2.4	8
75	Effect of Fenfluramine on Insulin/Growth Hormone Ratio in Obese Subjects. Pharmacology, 1988, 36, 106-111.	2.2	8
76	Massive Coronary Microvascular Dysfunction in Severe Anderson-Fabry Disease Cardiomyopathy. Circulation: Cardiovascular Imaging, 2019, 12, e009104.	2.6	8
77	Evidence of evolution towards left midventricular obstruction in severe Andersonâ€“Fabry cardiomyopathy. ESC Heart Failure, 2021, 8, 725-728.	3.1	8
78	Mesothelioma in Familial Mediterranean Fever With Colchicine Intolerance: A Case Report and Literature Review. Frontiers in Immunology, 2020, 11, 889.	4.8	8
79	Mitral valve endocarditis caused by streptococcus oralis occurring after upper gastrointestinal endoscopy. American Journal of Gastroenterology, 2002, 97, 2149-2150.	0.4	7
80	Early diagnosis and stage-adapted treatment of Wegenerâ€™s granulomatosis. Journal of Laryngology and Otology, 2003, 117, 208-211.	0.8	7
81	Effectiveness of Colchicine Therapy in 4 Cases of Retroperitoneal Fibrosis Associated with Autoinflammatory Diseases. Journal of Rheumatology, 2010, 37, 1971-1972.	2.0	7
82	Recommendations for the inclusion of Fabry disease as a rare febrile condition in existing algorithms for fever of unknown origin. Internal and Emergency Medicine, 2017, 12, 1059-1067.	2.0	7
83	Resolution of femoral metaphyseal dysplasia in CINCA syndrome after long-term treatment with interleukin-1 blockade. Clinical Rheumatology, 2018, 37, 2007-2009.	2.2	7
84	Role of Colchicine Treatment in Tumor Necrosis Factor Receptor Associated Periodic Syndrome (TRAPS): Real-Life Data from the AIDA Network. Mediators of Inflammation, 2020, 2020, 1-6.	3.0	7
85	Normalisation of High Ca 19-9 Values in Autoimmune Hepatitis after Steroidal Treatment. International Journal of Immunopathology and Pharmacology, 2005, 18, 603-607.	2.1	6
86	Serum macrophage migration inhibitory factor (MIF) in the intercritical phase of hereditary periodic fevers and its relationship with theMIFâ€173G/C polymorphism. Scandinavian Journal of Rheumatology, 2007, 36, 307-310.	1.1	6
87	A position for tumor necrosis factor inhibitors in the management of colchicine-resistant familial Mediterranean fever?. Immunology Letters, 2016, 180, 77-78.	2.5	6
88	Kidney involvement in the Schnitzler syndrome, a rare disease. CKJ: Clinical Kidney Journal, 2017, 10, 723-727.	2.9	6
89	Biotechnological Agents for Patients With Tumor Necrosis Factor Receptor Associated Periodic Syndromeâ€Therapeutic Outcome and Predictors of Response: Real-Life Data From the AIDA Network. Frontiers in Medicine, 2021, 8, 668173.	2.6	6
90	Cryoglobulinaemia: A True Internistic Disease?. International Journal of Immunopathology and Pharmacology, 2003, 16, 33-41.	2.1	6

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91	The Use of Chitotriosidase as a Marker of Active Sarcoidosis and in the Diagnosis of Fever of Unknown Origin (FUO). Journal of Clinical Medicine, 2021, 10, 5283.	2.4	6
92	Decreased insulin binding to red blood cells in liver cirrhosis. Acta Diabetologica Latina, 1983, 20, 251-256.	0.2	5
93	Leucoencephalitis after recombinant hepatitis B vaccine. Journal of Hepatology, 1996, 24, 764-765.	3.7	5
94	Rapid resolution of severe pericardial effusion using anakinra in a patient with COVID-19 vaccine-related acute pericarditis relapse: a case report. European Heart Journal - Case Reports, 2022, 6, ytac123.	0.6	5
95	Microscopic haematuria: a diagnostic aid in giant-cell arteritis?. Lancet, The, 1997, 350, 1226.	13.7	4
96	Non-Life-Threatening Sepsis: Report of Two Cases. American Journal of the Medical Sciences, 2004, 327, 275-277.	1.1	4
97	Tolerance induction to rofecoxib in a patient with Bartter's syndrome. Allergy: European Journal of Allergy and Clinical Immunology, 2004, 59, 788-789.	5.7	4
98	Haemophagocytic Syndrome Associated with Mucormycosis Infection. International Journal of Immunopathology and Pharmacology, 2012, 25, 751-755.	2.1	4
99	Early introduction of anakinra improves acute pericarditis and prevents tamponade in Staphylococcal sepsis. Internal and Emergency Medicine, 2021, 16, 1391-1394.	2.0	4
100	Drug survival of anakinra and canakinumab in monogenic autoinflammatory diseases: observational study from the International AIDA Registry. Rheumatology, 2021, 60, 5705-5712.	1.9	4
101	Therapeutic management of idiopathic recurrent serositis: a retrospective study. European Review for Medical and Pharmacological Sciences, 2020, 24, 3352-3359.	0.7	4
102	Multiple gated nuclear angiography performed at rest to evaluate left ventricular function in clinically asymptomatic diabetic patients. Nuclear Medicine Communications, 1984, 5, 681-688.	1.1	3
103	Stimulatory Effect of Pentagastrin on Growth Hormone and Prolactin Secretion in Normal Subjects. Experimental and Clinical Endocrinology and Diabetes, 1986, 88, 334-338.	1.2	3
104	Massive hepatomegaly following splenectomy for myeloid metaplasia. American Journal of Medicine, 1988, 84, 797.	1.5	3
105	Wegenerâ€™s granulomatosis: an update on diagnosis and therapy. Expert Review of Clinical Immunology, 2008, 4, 481-495.	3.0	3
106	Severe Giant Cell Arteritis Associated with Essential Thrombocythaemia. International Journal of Immunopathology and Pharmacology, 2010, 23, 1271-1274.	2.1	3
107	Epidemiology of FMF Worldwide. Rare Diseases of the Immune System, 2015, , 81-90.	0.1	3
108	Switch from anakinra to canakinumab in a severe case of CINCA syndrome. International Journal of Rheumatic Diseases, 2016, 19, 1354-1356.	1.9	3

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109	Potential resistance to SARS-CoV-2 infection in lysosomal storage disorders. CKJ: Clinical Kidney Journal, 2021, 14, 1488-1490.	2.9	3
110	Use of Intravenous Immunoglobulin Therapy at Unconventional Doses in Refractory Fulminant Systemic Lupus Erythematosus. European Journal of Case Reports in Internal Medicine, 2018, 5, 1.	0.4	3
111	Coronary Microvascular Dysfunction IsÂAssociated With a Worse Cardiac Phenotype in Patients With Fabry Disease. JACC: Cardiovascular Imaging, 2022, 15, 1518-1520.	5.3	3
112	Association of retroperitoneal fibrosis, proctitis and rectal stenosis. Postgraduate Medical Journal, 1981, 57, 674-676.	1.8	2
113	The Role of cAMP and Prostaglandins in Gastric Acid Secretion after Pentagastrin Administration. Experimental and Clinical Endocrinology and Diabetes, 1986, 87, 219-222.	1.2	2
114	Folic acid supplementation during methotrexate treatment: nonsense?. British Journal of Rheumatology, 2005, 44, 563-564.	2.3	2
115	Quantifying the efficacy of influenza vaccines. Lancet Infectious Diseases, The, 2012, 12, 659-660.	9.1	2
116	Giant cell arteritis and polymyalgia rheumatica after influenza vaccination: Comparing different experiences. Journal of Dermatology, 2012, 39, 888-889.	1.2	2
117	Systemic Complications of Esophageal Lichen Planus. International Journal of Immunopathology and Pharmacology, 2013, 26, 575-578.	2.1	2
118	Colchicine trial in PFAPA Syndrome and MEFV-negative patients. Pediatric Rheumatology, 2015, 13, .	2.1	2
119	Scleroderma and liver disease: a case of an association with primary sclerosing cholangitis. Scandinavian Journal of Rheumatology, 2016, 45, 334-335.	1.1	2
120	Improvement of liver involvement in familial Mediterranean fever after introduction of canakinumab: a case report. Mediterranean Journal of Hematology and Infectious Diseases, 2020, 12, e20200059.	1.3	2
121	Canakinumab improves patient-reported outcomes in children and adults with autoinflammatory recurrent fever syndromes: results from the CLUSTER trial. Clinical and Experimental Rheumatology, 2021, 39, 51-58.	0.8	2
122	Thromboxane production in diabetes mellitus. Research in Clinic and Laboratory, 1986, 16, 539-542.	0.3	2
123	Circulating levels of cyclosporin A in inflammatory bowel disease: relationships with lymphocyte inhibition and the age of patients. European Journal of Clinical Pharmacology, 2004, 60, 161-164.	1.9	1
124	La maladie pÃ©riodique. Revue Du Rhumatisme (Edition Francaise), 2009, 76, 382-389.	0.0	1
125	Long-term treatment with anakinra and canakinumab resolves patellar subchondral erosion in neonatal-onset multisystem inflammatory disease. Reumatismo, 2019, 71, 53-55.	0.9	1
126	Effect of somatostatin (SRIF) on plasma glucose and insulin response to glucagon in liver cirrhosis. Acta Diabetologica Latina, 1979, 16, 139-145.	0.2	0

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127	Adult Respiratory Distress Syndrome. Archives of Internal Medicine, 1981, 141, 268.	3.8	0
128	Does Secretin Control Insulin Secretion?. Experimental and Clinical Endocrinology and Diabetes, 1984, 84, 81-86.	1.2	0
129	Evidence of Extragastric Gastrin Release in Postoperative Ulcer Patients. Experimental and Clinical Endocrinology and Diabetes, 1986, 87, 223-226.	1.2	0
130	The Effect of Lysine Acetylsalicylate on Somatostatin Inhibition of Insulin Secretion Induced by Arginine. Experimental and Clinical Endocrinology and Diabetes, 1986, 88, 119-122.	1.2	0
131	CASE REPORT: A SWOLLEN AND RED EAR. Journal of the American Geriatrics Society, 2003, 51, 138-139.	2.6	0
132	Globus pharyngis: was it a stroke of lightning?. American Journal of Gastroenterology, 2003, 98, 938-939.	0.4	0
133	FMF revisited. European Journal of Human Genetics, 2004, 12, 255-255.	2.8	0
134	PFAPA syndrome as an hereditary autoinflamatory disorder. Pediatric Rheumatology, 2015, 13, .	2.1	0
135	Do we understand the pathophysiology of gastrointestinal symptoms in patients with Fabry disease?. Molecular Genetics and Metabolism, 2018, 123, S63.	1.1	0
136	Proposal of a rating scale to recognize Fabry disease in patients with nonspecific gastrointestinal symptoms. Molecular Genetics and Metabolism, 2018, 123, S63-S64.	1.1	0
137	AB1305â€…EVALUATION OF SERUM LEVELS OF ASC FOR THE DIAGNOSIS AND MONITORING OF CRYOPYRIN ASSOCIATED PERIODIC SYNDROMES (CAPS). , 2019, , .		0
138	Pediatric Motor Inflammatory Neuropathy: The Role of Antiphospholipid Antibodies. Brain Sciences, 2020, 10, 156.	2.3	0
139	Anderson-Fabryâ€™s Disease: A Rare but Treatable Case of Fever of Unknown Origin. European Journal of Case Reports in Internal Medicine, 2017, 2, 000645.	0.4	0
140	Intravenous immunoglobulin for Pediatric Neuropsychiatric Lupus Triggered by Epstein-Barr Virus Cerebral Infection. Israel Medical Association Journal, 2016, 18, 763-766.	0.1	0
141	Canakinumab improves patient-reported outcomes in children and adults with autoinflammatory recurrent fever syndromes: results from the CLUSTER trial. Clinical and Experimental Rheumatology, 2021, 39 Suppl 132, 51-58.	0.8	0
142	Transitional care management in patients with auto-inflammatory diseases: experience of cooperation of a paediatric and adult centre. Journal of Transition Medicine, 2022, 4, .	0.5	0
143	Mitral valve endocarditis caused by streptococcus oralis occurring after upper gastrointestinal endoscopy. American Journal of Gastroenterology, 2002, 97, 2149-2150.	0.4	0