

Alberto Tommasini

List of Publications by Citations

Source: <https://exaly.com/author-pdf/5718892/alberto-tommasini-publications-by-citations.pdf>

Version: 2024-04-27

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

161
papers

4,106
citations

34
h-index

59
g-index

179
ext. papers

4,855
ext. citations

4.8
avg, IF

4.94
L-index

#	Paper	IF	Citations
161	Zonulin, a newly discovered modulator of intestinal permeability, and its expression in coeliac disease. <i>Lancet, The</i> , 2000 , 355, 1518-9	4.0	412
160	Clinical and molecular profile of a new series of patients with immune dysregulation, polyendocrinopathy, enteropathy, X-linked syndrome: inconsistent correlation between forkhead box protein 3 expression and disease severity. <i>Journal of Allergy and Clinical Immunology</i> , 2008 , 122, 1105-1112.e1	11.5	165
159	Mass screening for coeliac disease using antihuman transglutaminase antibody assay. <i>Archives of Disease in Childhood</i> , 2004 , 89, 512-5	2.2	161
158	Human recombinant tissue transglutaminase ELISA: an innovative diagnostic assay for celiac disease. <i>American Journal of Gastroenterology</i> , 2000 , 95, 1253-7	0.7	156
157	Molecular dissection of the tissue transglutaminase autoantibody response in celiac disease. <i>Journal of Immunology</i> , 2001 , 166, 4170-6	5.3	148
156	Human IL2RA null mutation mediates immunodeficiency with lymphoproliferation and autoimmunity. <i>Clinical Immunology</i> , 2013 , 146, 248-61	9	141
155	ADA2 deficiency (DADA2) as an unrecognised cause of early onset polyarteritis nodosa and stroke: a multicentre national study. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1648-1656	2.4	139
154	Type I interferon-mediated autoinflammation due to DNase II deficiency. <i>Nature Communications</i> , 2017 , 8, 2176	17.4	111
153	Differentiating PFAPA syndrome from monogenic periodic fevers. <i>Pediatrics</i> , 2009 , 124, e721-8	7.4	109
152	Undiagnosed coeliac disease and risk of autoimmune disorders in subjects with Type I diabetes mellitus. <i>Diabetologia</i> , 2001 , 44, 151-5	10.3	108
151	The efficacy of anakinra in an adolescent with colchicine-resistant familial Mediterranean fever. <i>European Journal of Pediatrics</i> , 2008 , 167, 695-6	4.1	101
150	Follow-up and quality of life of patients with cryopyrin-associated periodic syndromes treated with Anakinra. <i>Journal of Pediatrics</i> , 2010 , 157, 310-315.e1	3.6	88
149	Genetics of inflammatory bowel disease from multifactorial to monogenic forms. <i>World Journal of Gastroenterology</i> , 2015 , 21, 12296-310	5.6	87
148	Long-term clinical profile of children with the low-penetrance R92Q mutation of the TNFRSF1A gene. <i>Arthritis and Rheumatism</i> , 2011 , 63, 1141-50		85
147	X-chromosome inactivation analysis in a female carrier of FOXP3 mutation. <i>Clinical and Experimental Immunology</i> , 2002 , 130, 127-30	6.2	70
146	Development of a novel rapid non-invasive screening test for coeliac disease. <i>Gut</i> , 2000 , 47, 628-31	19.2	64
145	Functional type 1 regulatory T cells develop regardless of FOXP3 mutations in patients with IPEX syndrome. <i>European Journal of Immunology</i> , 2011 , 41, 1120-31	6.1	59

144	MVK mutations and associated clinical features in Italian patients affected with autoinflammatory disorders and recurrent fever. <i>European Journal of Human Genetics</i> , 2005 , 13, 314-20	5.3	58
143	Post-COVID-19 symptoms 6 months after acute infection among hospitalized and non-hospitalized patients. <i>Clinical Microbiology and Infection</i> , 2021 , 27, 1507-1513	9.5	58
142	Usefulness of screening program for celiac disease in autoimmune thyroiditis. <i>Digestive Diseases and Sciences</i> , 2000 , 45, 403-6	4	57
141	Describing Kawasaki shock syndrome: results from a retrospective study and literature review. <i>Clinical Rheumatology</i> , 2017 , 36, 223-228	3.9	55
140	Clinical impact of MEFV mutations in children with periodic fever in a prevalent western European Caucasian population. <i>Annals of the Rheumatic Diseases</i> , 2012 , 71, 1961-5	2.4	51
139	Forkhead box protein 3 (FOXP3) mutations lead to increased TH17 cell numbers and regulatory T-cell instability. <i>Journal of Allergy and Clinical Immunology</i> , 2011 , 128, 1376-1379.e1	11.5	47
138	Natural isoprenoids are able to reduce inflammation in a mouse model of mevalonate kinase deficiency. <i>Pediatric Research</i> , 2008 , 64, 177-82	3.2	47
137	Wharton's jelly derived mesenchymal stromal cells: Biological properties, induction of neuronal phenotype and current applications in neurodegeneration research. <i>Acta Histochemica</i> , 2015 , 117, 329-38		45
136	Regulatory T-cell function is impaired in celiac disease. <i>Digestive Diseases and Sciences</i> , 2009 , 54, 1513-9	4	44
135	Altered germinal center reaction and abnormal B cell peripheral maturation in PI3KR1-mutated patients presenting with HIGM-like phenotype. <i>Clinical Immunology</i> , 2015 , 159, 33-6	9	43
134	Looking for celiac disease: diagnostic accuracy of two rapid commercial assays. <i>American Journal of Gastroenterology</i> , 2006 , 101, 1597-600	0.7	42
133	A reliable screening procedure for coeliac disease in clinical practice. <i>Scandinavian Journal of Gastroenterology</i> , 2002 , 37, 679-84	2.4	42
132	Next-generation sequencing and its initial applications for molecular diagnosis of systemic auto-inflammatory diseases. <i>Annals of the Rheumatic Diseases</i> , 2016 , 75, 1550-7	2.4	41
131	Wild-type FOXP3 is selectively active in CD4+CD25(hi) regulatory T cells of healthy female carriers of different FOXP3 mutations. <i>Blood</i> , 2009 , 114, 4138-41	2.2	37
130	Long noncoding RNA GAS5: a novel marker involved in glucocorticoid response. <i>Current Molecular Medicine</i> , 2015 , 15, 94-9	2.5	36
129	Therapeutic strategy in p47-phox deficient chronic granulomatous disease presenting as inflammatory bowel disease. <i>Journal of Allergy and Clinical Immunology</i> , 2010 , 125, 943-946.e1	11.5	36
128	The immunosuppressive effect of Wharton's jelly stromal cells depends on the timing of their licensing and on lymphocyte activation. <i>Cytotherapy</i> , 2010 , 12, 154-60	4.8	34
127	Natural isoprenoids inhibit LPS-induced-production of cytokines and nitric oxide in aminobisphosphonate-treated monocytes. <i>International Immunopharmacology</i> , 2010 , 10, 639-42	5.8	32

126	Recent Insight into SARS-CoV2 Immunopathology and Rationale for Potential Treatment and Preventive Strategies in COVID-19. <i>Vaccines</i> , 2020 , 8,	5.3	31
125	Celiac disease in patients with sporadic and inherited cardiomyopathies and in their relatives. <i>European Heart Journal</i> , 2003 , 24, 1455-61	9.5	30
124	Dealing with Chronic Non-Bacterial Osteomyelitis: a practical approach. <i>Pediatric Rheumatology</i> , 2017 , 15, 87	3.5	29
123	Diagnostics and therapeutic insights in a severe case of mevalonate kinase deficiency. <i>Pediatrics</i> , 2007 , 119, e523-7	7.4	29
122	Off-Label Use of Sirolimus and Everolimus in a Pediatric Center: A Case Series and Review of the Literature. <i>Paediatric Drugs</i> , 2019 , 21, 185-193	4.2	27
121	Role of the Long Non-Coding RNA Growth Arrest-Specific 5 in Glucocorticoid Response in Children with Inflammatory Bowel Disease. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2018 , 122, 87-93	3.1	27
120	Medium-term survival without haematopoietic stem cell transplantation in a case of IPEX: insights into nutritional and immunosuppressive therapy. <i>European Journal of Pediatrics</i> , 2007 , 166, 1195-7	4.1	24
119	Severe inflammatory bowel disease associated with congenital alteration of transforming growth factor beta signaling. <i>Journal of Crohns and Colitis</i> , 2014 , 8, 770-4	1.5	22
118	The diagnostic challenge of very early-onset enterocolitis in an infant with XIAP deficiency. <i>BMC Pediatrics</i> , 2015 , 15, 208	2.6	22
117	Prevalence of autoimmune disorders in relatives of patients with celiac disease. <i>Digestive Diseases and Sciences</i> , 2002 , 47, 1427-31	4	22
116	Clinical Characteristics of Patients Carrying the Q703K Variant of the NLRP3 Gene: A 10-year Multicentric National Study. <i>Journal of Rheumatology</i> , 2016 , 43, 1093-100	4.1	22
115	The MDM2 inhibitor Nutlin-3 modulates dendritic cell-induced T cell proliferation. <i>Human Immunology</i> , 2012 , 73, 342-5	2.3	21
114	Curcumin Anti-Apoptotic Action in a Model of Intestinal Epithelial Inflammatory Damage. <i>Nutrients</i> , 2017 , 9,	6.7	20
113	Variant mannose-binding lectin alleles are associated with celiac disease. <i>Immunogenetics</i> , 2002 , 54, 596-82	3.2	20
112	Characterization of the anti-tissue transglutaminase antibody response in nonobese diabetic mice. <i>Journal of Immunology</i> , 2005 , 174, 5830-6	5.3	20
111	Clinical, Immunological, and Molecular Features of Typical and Atypical Severe Combined Immunodeficiency: Report of the Italian Primary Immunodeficiency Network. <i>Frontiers in Immunology</i> , 2019 , 10, 1908	8.4	19
110	Genetic profile of patients with early onset inflammatory bowel disease. <i>Gene</i> , 2018 , 645, 18-29	3.8	19
109	Cryopyrin-associated Periodic Syndromes in Italian Patients: Evaluation of the Rate of Somatic NLRP3 Mosaicism and Phenotypic Characterization. <i>Journal of Rheumatology</i> , 2017 , 44, 1667-1673	4.1	19

108	Failure of interferon- γ -pre-treated mesenchymal stem cell treatment in a patient with Crohn's disease. <i>World Journal of Gastroenterology</i> , 2015 , 21, 4379-84	5.6	19
107	Dual sugar gut-permeability testing on blood drop in animal models. <i>Clinica Chimica Acta</i> , 2005 , 352, 191-7	6.2	18
106	The Complex Interplay between Lipids, Immune System and Interleukins in Cardio-Metabolic Diseases. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	18
105	The farnesyltransferase inhibitors tipifarnib and lonafarnib inhibit cytokines secretion in a cellular model of mevalonate kinase deficiency. <i>Pediatric Research</i> , 2011 , 70, 78-82	3.2	17
104	Diagnostic Approach to Monogenic Inflammatory Bowel Disease in Clinical Practice: A Ten-Year Multicentric Experience. <i>Inflammatory Bowel Diseases</i> , 2020 , 26, 720-727	4.5	17
103	Genetic profiling of autoinflammatory disorders in patients with periodic fever: a prospective study. <i>Pediatric Rheumatology</i> , 2015 , 13, 11	3.5	16
102	F402L variant in NLRP12 in subjects with undiagnosed periodic fevers and in healthy controls. <i>Clinical and Experimental Rheumatology</i> , 2014 , 32, 993-4	2.2	16
101	A common genetic background could explain early-onset Crohn's disease. <i>Medical Hypotheses</i> , 2012 , 78, 520-2	3.8	15
100	Targeting farnesyl-transferase as a novel therapeutic strategy for mevalonate kinase deficiency: in vitro and in vivo approaches. <i>Pharmacological Research</i> , 2010 , 61, 506-10	10.2	15
99	Testing for anti-human transglutaminase antibodies in saliva is not useful for diagnosis of celiac disease. <i>Clinical Chemistry</i> , 2004 , 50, 216-9	5.5	15
98	One-step cloning of anti tissue transglutaminase scFv from subjects with celiac disease. <i>Journal of Autoimmunity</i> , 2004 , 22, 65-72	15.5	15
97	Fate of lymphocytes after withdrawal of tofacitinib treatment. <i>PLoS ONE</i> , 2014 , 9, e85463	3.7	15
96	Long term outcome of eight patients with type 1 Leukocyte Adhesion Deficiency (LAD-1): Not only infections, but high risk of autoimmune complications. <i>Clinical Immunology</i> , 2018 , 191, 75-80	9	14
95	Phylloid pattern of hypomelanosis closely related to chromosomal abnormalities in the 13q detected by SNP array analysis. <i>Dermatology</i> , 2012 , 225, 294-7	4.4	14
94	Decreased cholesterol levels reflect a consumption of anti-inflammatory isoprenoids associated with an impaired control of inflammation in a mouse model of mevalonate kinase deficiency. <i>Inflammation Research</i> , 2010 , 59, 335-8	7.2	14
93	Inhibition of mesenchymal stromal cells by pre-activated lymphocytes and their culture media. <i>Stem Cell Research and Therapy</i> , 2014 , 5, 3	8.3	13
92	Combined Analysis of Methylation and Gene Expression Profiles in Separate Compartments of Small Bowel Mucosa Identified Celiac Disease Patients' Signatures. <i>Scientific Reports</i> , 2019 , 9, 10020	4.9	12
91	Inflammation profile of four early onset Crohn patients. <i>Gene</i> , 2012 , 493, 282-5	3.8	12

90	The universe of immune deficiencies in Crohn's disease: a new viewpoint for an old disease?. <i>Scandinavian Journal of Gastroenterology</i> , 2010 , 45, 1141-9	2.4	12
89	Differential expression of GAS5 in rapamycin-induced reversion of glucocorticoid resistance. <i>Clinical and Experimental Pharmacology and Physiology</i> , 2016 , 43, 602-5	3	12
88	Cryptic gluten intolerance in type 1 diabetes: identifying suitable candidates for a gluten free diet. <i>Gut</i> , 2006 , 55, 133-4	19.2	11
87	Diagnostic challenge of hyper-IgD syndrome in four children with inflammatory gastrointestinal complaints. <i>Scandinavian Journal of Gastroenterology</i> , 2006 , 41, 430-6	2.4	11
86	Elective bone marrow transplantation in a child with X-linked hyper-IgM syndrome presenting with acute respiratory distress syndrome. <i>Bone Marrow Transplantation</i> , 2002 , 30, 49-52	4.4	11
85	Plasmacytoid Dendritic Cells Depletion and Elevation of IFN- γ -Dependent Chemokines CXCL9 and CXCL10 in Children With Multisystem Inflammatory Syndrome. <i>Frontiers in Immunology</i> , 2021 , 12, 654587	8.4	11
84	An Easy and Reliable Strategy for Making Type I Interferon Signature Analysis Comparable among Research Centers. <i>Diagnostics</i> , 2019 , 9,	3.8	10
83	CARD15/NOD2 mutations are not related to abdominal PFAPA. <i>Journal of Pediatrics</i> , 2006 , 149, 427	3.6	10
82	Anti-transglutaminase antibodies and age. <i>Clinical Chemistry</i> , 2004 , 50, 1856-60	5.5	10
81	Chronic infantile neurological cutaneous articular syndrome: CD10 over-expression in neutrophils is a possible key to the pathogenesis of the disease. <i>European Journal of Pediatrics</i> , 2003 , 162, 669-73	4.1	10
80	Reappraisal of Antimalarials in Interferonopathies: New Perspectives for Old Drugs. <i>Current Medicinal Chemistry</i> , 2018 , 25, 2797-2810	4.3	10
79	Case Report: Use of Anakinra in Multisystem Inflammatory Syndrome During COVID-19 Pandemic. <i>Frontiers in Pediatrics</i> , 2020 , 8, 624248	3.4	10
78	In vivo detection of polyomaviruses JCV and SV40 in mesenchymal stem cells from human umbilical cords. <i>Pediatric Blood and Cancer</i> , 2014 , 61, 1347-9	3	9
77	Evolutionary hypothesis of the Mevalonate Kinase Deficiency. <i>Medical Hypotheses</i> , 2013 , 80, 67-9	3.8	9
76	Clinical genetic testing of periodic fever syndromes. <i>BioMed Research International</i> , 2013 , 2013, 501305	3	9
75	Higher interferon score and normal complement levels may identify a distinct clinical subset in children with systemic lupus erythematosus. <i>Arthritis Research and Therapy</i> , 2020 , 22, 91	5.7	8
74	Theophylline as a precision therapy in a young girl with PIK3R1 immunodeficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2018 , 6, 2165-2167	5.4	8
73	Novel NOD2 Mutation in Early-Onset Inflammatory Bowel Phenotype. <i>Inflammatory Bowel Diseases</i> , 2018 , 24, 1204-1212	4.5	8

72	Neuronal Dysfunction Associated with Cholesterol Deregulation. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	8
71	Novel missense mutation in the NOD2 gene in a patient with early onset ulcerative colitis: causal or chance association?. <i>International Journal of Molecular Sciences</i> , 2014 , 15, 3834-41	6.3	8
70	T cells stimulated in vitro have a suppressive function but do not contain only regulatory T cells. <i>Clinical and Experimental Immunology</i> , 2007 , 150, 561-6	6.2	8
69	Biological and Clinical Changes in a Pediatric Series Treated with Off-Label JAK Inhibitors. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	8
68	Different presentations of mevalonate kinase deficiency: a case series. <i>Clinical and Experimental Rheumatology</i> , 2015 , 33, 437-42	2.2	8
67	Genetic and functional profiling of Crohn's disease: autophagy mechanism and susceptibility to infectious diseases. <i>BioMed Research International</i> , 2013 , 2013, 297501	3	7
66	Hughes Stovin: Sustained remission and regression of pulmonary aneurysms with anti-tumor necrosis factor treatment. <i>Pediatric Pulmonology</i> , 2019 , 54, E13-E15	3.5	6
65	Immunomodulatory drugs in autoimmune lymphoproliferative syndrome (ALPS). <i>Pediatric Blood and Cancer</i> , 2012 , 58, 310; author reply 311	3	6
64	Differential action of 3-hydroxyanthranilic acid on viability and activation of stimulated lymphocytes. <i>International Immunopharmacology</i> , 2011 , 11, 2242-5	5.8	6
63	Selective resistance to different glucocorticoids in severe autoimmune disorders. <i>Clinical Immunology</i> , 2010 , 134, 313-9	9	6
62	Human tissue transglutaminase ELISA and an old study: a revision of the blood donor screening study for coeliac disease in the USA. <i>Scandinavian Journal of Gastroenterology</i> , 2004 , 39, 195-7	2.4	6
61	Repositioning Drugs for Rare Immune Diseases: Hopes and Challenges for a Precision Medicine. <i>Current Medicinal Chemistry</i> , 2018 , 25, 2764-2782	4.3	6
60	The Italian Registry for Primary Immunodeficiencies (Italian Primary Immunodeficiency Network; IPINet): Twenty Years of Experience (1999-2019). <i>Journal of Clinical Immunology</i> , 2020 , 40, 1026-1037	5.7	6
59	Innovative Target Therapies Are Able to Block the Inflammation Associated with Dysfunction of the Cholesterol Biosynthesis Pathway. <i>International Journal of Molecular Sciences</i> , 2015 , 17,	6.3	6
58	Low-dose sirolimus in two cousins with autoimmune lymphoproliferative syndrome-associated infection. <i>Pediatrics International</i> , 2018 , 60, 315-317	1.2	5
57	Hemophagocytic lymphohistiocytosis in total parenteral nutrition dependent children: description of 5 cases and practical tips for management. <i>Journal of Pediatric Hematology/Oncology</i> , 2014 , 36, e440-2 ^{1,2}		5
56	Patients Induced Pluripotent Stem Cells to Model Drug Induced Adverse Events: A Role in Predicting Thiopurine Induced Pancreatitis?. <i>Current Drug Metabolism</i> , 2015 , 17, 91-8	3.5	5
55	"Hyper-IgD syndrome" or "mevalonate kinase deficiency": an old syndrome needing a new name?. <i>Rheumatology International</i> , 2014 , 34, 423-4	3.6	5

54	Amantadine in chronic granulomatous disease. <i>Pediatric Hematology and Oncology</i> , 2005 , 22, 147-51	1.7	5
53	Altered pattern of tumor necrosis factor-alpha production in peripheral blood monocytes from Crohn's disease. <i>World Journal of Gastroenterology</i> , 2016 , 22, 9117-9126	5.6	5
52	To Extinguish the Fire from Outside the Cell or to Shutdown the Gas Valve Inside? Novel Trends in Anti-Inflammatory Therapies. <i>International Journal of Molecular Sciences</i> , 2015 , 16, 21277-93	6.3	4
51	Neither hereditary periodic fever nor periodic fever, aphthae, pharyngitis, adenitis: Undifferentiated periodic fever in a tertiary pediatric center. <i>World Journal of Clinical Pediatrics</i> , 2018 , 7, 49-55	2.5	4
50	Genetic and immunologic findings in children with recurrent aphthous stomatitis with systemic inflammation. <i>Pediatric Rheumatology</i> , 2021 , 19, 70	3.5	4
49	Monocyte-predominant engraftment, cytokine levels and early transplant-related complications in pediatric hematopoietic stem cell recipients. <i>Cancer Medicine</i> , 2019 , 8, 890-901	4.8	3
48	Pulmonary arterial hypertension in interferonopathies: a case report and a review of the literature. <i>Pulmonary Circulation</i> , 2019 , 9, 2045894019869837	2.7	3
47	Selective IgA deficiency: ruling out coeliac disease and selective antibody deficiency to polysaccharides. <i>Journal of Clinical Immunology</i> , 2013 , 33, 1149	5.7	3
46	JCV+ Patients with Inflammatory bowel disease show elevated plasma levels of MIG and SCF. <i>Inflammatory Bowel Diseases</i> , 2012 , 18, 1194-6	4.5	3
45	Defective and excessive immunities in pediatric diseases. <i>Current Pharmaceutical Design</i> , 2012 , 18, 5729-34	3.4	3
44	Autoantibodies as predictors of disease. <i>Lancet, The</i> , 2004 , 364, 1403-4	4.0	3
43	High prevalence of rare FBLIM1 gene variants in an Italian cohort of patients with Chronic Non-bacterial Osteomyelitis (CNO). <i>Pediatric Rheumatology</i> , 2020 , 18, 55	3.5	3
42	Putative modifier genes in mevalonate kinase deficiency. <i>Molecular Medicine Reports</i> , 2016 , 13, 3181-9	2.9	3
41	Familial hypogammaglobulinemia with high RTE and naïve T lymphocytes. <i>Inflammation Research</i> , 2019 , 68, 901-904	7.2	2
40	Clinical significance of hyper-IgA in a paediatric laboratory series. <i>Archives of Disease in Childhood</i> , 2014 , 99, 1114-6	2.2	2
39	TNF- β NP rs1800629 and risk of relapse in childhood acute lymphoblastic leukemia: relation to immunophenotype. <i>Pharmacogenomics</i> , 2014 , 15, 619-27	2.6	2
38	A red baby should not be taken too lightly. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2012 , 101, e573-7	3.1	2
37	Clinical immunology Primary Immunodeficiency Diseases in two neighboring pediatric centers: registry data bring out a wide spectrum of diseases with complex clinical presentations. <i>Central-European Journal of Immunology</i> , 2012 , 4, 365-370	1.6	2

36	Gastrointestinal Foxp3 expression in normal, inflammatory and neoplastic conditions. <i>Pathology</i> , 2011 , 43, 465-71	1.6	2
35	Is autophagy an elective strategy to protect neurons from dysregulated cholesterol metabolism?. <i>Neural Regeneration Research</i> , 2019 , 14, 582-587	4.5	2
34	Fever tree revisited: From malaria to autoinflammatory diseases. <i>World Journal of Clinical Pediatrics</i> , 2015 , 4, 106-12	2.5	2
33	A Case of Uveitis in a Patient With Juvenile Myelomonocytic Leukemia Successfully Treated With Adalimumab. <i>Journal of Pediatric Hematology/Oncology</i> , 2020 , 42, e373-e376	1.2	2
32	Priming of the cGAS-STING-TBK1 Pathway Enhances LPS-Induced Release of Type I Interferons. <i>Cells</i> , 2021 , 10,	7.9	2
31	Antibodies reacting to mimotopes of Simian virus 40 large T antigen, the viral oncoprotein, in sera from children. <i>Journal of Cellular Physiology</i> , 2019 , 234, 3170-3179	7	2
30	B-cell depletion induces prolonged remission in patients with giant cell hepatitis and autoimmune hemolytic anemia. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2020 , 44, 66-72	2.4	2
29	Case Report: Refractory Autoimmune Gastritis Responsive to Abatacept in LRBA Deficiency. <i>Frontiers in Immunology</i> , 2021 , 12, 619246	8.4	2
28	The Challenge of Next Generation Sequencing in a Boy With Severe Mononucleosis and EBV-related Lymphoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2018 , 40, e323-e326	1.2	2
27	Acute rheumatic fever prophylaxis in high-income countries: clinical observations from an Italian multicentre, retrospective study. <i>Clinical and Experimental Rheumatology</i> , 2020 , 38, 1016-1020	2.2	2
26	Mevalonate kinase deficiency: therapeutic targets, treatments, and outcomes. <i>Expert Opinion on Orphan Drugs</i> , 2017 , 5, 515-524	1.1	1
25	Tregs and Th17 lymphocytes in human DYRK1A haploinsufficiency. <i>Immunology Letters</i> , 2019 , 214, 52-54	4.1	1
24	Biomarkers and Precision Therapy for Primary Immunodeficiencies: An In Vitro Study Based on Induced Pluripotent Stem Cells From Patients. <i>Clinical Pharmacology and Therapeutics</i> , 2020 , 108, 358-367	6.1	1
23	Action of methotrexate and tofacitinib on directly stimulated and bystander-activated lymphocytes. <i>Molecular Medicine Reports</i> , 2016 , 14, 574-82	2.9	1
22	Fecal Calprotectin to Detect Inflammatory Bowel Disease in Juvenile Idiopathic Arthritis. <i>Journal of Rheumatology</i> , 2018 , 45, 1418-1421	4.1	1
21	Two-gene mutation in a single patient: Biochemical and functional analysis for a correct interpretation of exome results. <i>Molecular Medicine Reports</i> , 2015 , 12, 6128-32	2.9	1
20	Heterozygous nucleotide-binding oligomerization domain-2 mutations affect monocyte maturation in Crohn's disease. <i>World Journal of Gastroenterology</i> , 2007 , 13, 6191-6	5.6	1
19	SARS-CoV-2 Infection and Inflammatory Response in a Twin Pregnancy. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	1

18	Hydroxychloroquine modulates immunological pathways activated by RNA:DNA hybrids in Aicardi-Goutières syndrome patients carrying RNASEH2 mutations. <i>Cellular and Molecular Immunology</i> , 2021 , 18, 1593-1595	15.4	1
17	Immunity and Genetics at the Revolving Doors of Diagnostics in Primary Immunodeficiencies. <i>Diagnostics</i> , 2021 , 11,	3.8	1
16	Carbamazepine-induced thrombocytopenic purpura in a child: Insights from a genomic analysis. <i>Blood Cells, Molecules, and Diseases</i> , 2016 , 59, 97-9	2.1	1
15	Case Report: EBV Chronic Infection and Lymphoproliferation in Four APDS Patients: The Challenge of Proper Characterization, Therapy, and Follow-Up. <i>Frontiers in Pediatrics</i> , 2021 , 9, 703853	3.4	1
14	Mevalonate Kinase Deficiency and Squalene Synthase Inhibitor (TAK-475): The Balance to Extinguish the Inflammation. <i>Biomolecules</i> , 2021 , 11,	5.9	1
13	A Novel ELISA-Based Peptide Biosensor Assay for Screening ABL1 Activity : A Challenge for Precision Therapy in BCR-ABL1 and BCR-ABL1 Like Leukemias. <i>Frontiers in Pharmacology</i> , 2021 , 12, 749381	5.6	0
12	Genetic Variants Assessing Crohn's Disease Pattern in Pediatric Inflammatory Bowel Disease Patients by a Clinical Exome Survey.. <i>Bioinformatics and Biology Insights</i> , 2021 , 15, 11779322211055285	5.3	0
11	Consensus of the Italian Primary Immunodeficiency Network on transition management from pediatric to adult care in patients affected with childhood-onset inborn errors of immunity. <i>Journal of Allergy and Clinical Immunology</i> , 2020 , 146, 967-983	11.5	0
10	response to mucosal bacteria and muramyl dipeptide in inflammatory bowel disease. <i>World Journal of Gastroenterology</i> , 2016 , 22, 9734-9743	5.6	0
9	Clinical and Cytometric Study of Immune Involvement in a Heterogeneous Cohort of Subjects With RASopathies and mTORopathies. <i>Frontiers in Pediatrics</i> , 2021 , 9, 703613	3.4	0
8	Druggable monogenic immune defects hidden in diverse medical specialties: Focus on overlap syndromes.. <i>World Journal of Clinical Pediatrics</i> , 2022 , 11, 136-150	2.5	0
7	Le interferonopatie di tipo I. <i>Medico E Bambino</i> , 2021 , 40, 509-514	0.4	
6	From bone to skin. <i>Journal of Paediatrics and Child Health</i> , 2020 , 56, 1481	1.3	
5	Le sindromi autoinfiammatorie: quando non è solo PFAPA. <i>Medico E Bambino</i> , 2021 , 40, 221-225	0.4	
4	Swollen Ankle with a Hole: Brodie Abscess. <i>Journal of Pediatrics</i> , 2021 , 236, 319-320	3.6	
3	Cutaneous manifestations in mevalonate kinase deficient patients treated with canakinumab. <i>Clinical and Experimental Rheumatology</i> , 2020 , 38, 760-762	2.2	
2	Complex regional pain syndrome of the foot in a girl with post-SARS-CoV-2 chilblains. <i>Clinical and Experimental Rheumatology</i> , 2021 ,	2.2	
1	Cytofluorimetric assay to investigate variability in blinatumomab response.. <i>Frontiers in Bioscience</i> , 2022 , 27, 39		

