

Elena Palmisani

List of Publications by Citations

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

30
papers

504
citations

11
h-index

22
g-index

32
ext. papers

617
ext. citations

4.3
avg, IF

2.72
L-index

#	Paper	IF	Citations
30	The PRINTO criteria for clinically inactive disease in juvenile dermatomyositis. <i>Annals of the Rheumatic Diseases</i> , 2013 , 72, 686-93	2.4	84
29	Development and validation of a preliminary definition of minimal disease activity in patients with juvenile idiopathic arthritis. <i>Arthritis and Rheumatism</i> , 2008 , 59, 1120-7		80
28	Mycophenolate mofetil and Sirolimus as second or further line treatment in children with chronic refractory Primitive or Secondary Autoimmune Cytopenias: a single centre experience. <i>British Journal of Haematology</i> , 2015 , 171, 247-253	4.5	39
27	Factors associated with achievement of inactive disease in children with juvenile idiopathic arthritis treated with etanercept. <i>Journal of Rheumatology</i> , 2013 , 40, 192-200	4.1	39
26	Correlation between juvenile idiopathic arthritis activity and damage measures in early, advanced, and longstanding disease. <i>Arthritis and Rheumatism</i> , 2006 , 55, 843-9		37
25	Temporomandibular Joint Involvement in Association With Quality of Life, Disability, and High Disease Activity in Juvenile Idiopathic Arthritis. <i>Arthritis Care and Research</i> , 2017 , 69, 677-686	4.7	32
24	Mycophenolate mofetil for the treatment of children with immune thrombocytopenia and Evans syndrome. A retrospective data review from the Italian association of paediatric haematology/oncology. <i>British Journal of Haematology</i> , 2016 , 175, 490-495	4.5	30
23	Pearson Syndrome: A Retrospective Cohort Study from the Marrow Failure Study Group of A.I.E.O.P. (Associazione Italiana Emato-Oncologia Pediatrica). <i>JIMD Reports</i> , 2016 , 26, 37-43	1.9	28
22	Sirolimus for the treatment of multi-resistant autoimmune haemolytic anaemia in children. <i>British Journal of Haematology</i> , 2014 , 167, 571-4	4.5	24
21	FAS-mediated apoptosis impairment in patients with ALPS/ALPS-like phenotype carrying variants on CASP10 gene. <i>British Journal of Haematology</i> , 2019 , 187, 502-508	4.5	21
20	Agreement between physicians and parents in rating functional ability of children with juvenile idiopathic arthritis. <i>Pediatric Rheumatology</i> , 2007 , 5, 23	3.5	14
19	Therapeutic approaches for the treatment of renal disease in juvenile systemic lupus erythematosus: an international multicentre PRINTO study. <i>Annals of the Rheumatic Diseases</i> , 2013 , 72, 1503-9	2.4	11
18	A Meta-Analysis to Estimate the Placebo Effect in Randomized Controlled Trials in Juvenile Idiopathic Arthritis. <i>Arthritis and Rheumatology</i> , 2016 , 68, 1540-50	9.5	10
17	Sirolimus as a rescue therapy in children with immune thrombocytopenia refractory to mycophenolate mofetil. <i>American Journal of Hematology</i> , 2018 , 93, E175-E177	7.1	9
16	RAG deficiency with ALPS features successfully treated with TCR/CD19 cell depleted haploidentical stem cell transplant. <i>Clinical Immunology</i> , 2018 , 187, 102-103	9	9
15	Clinical features and therapeutic challenges of cytopenias belonging to alps and alps-related (ARS) phenotype. <i>British Journal of Haematology</i> , 2019 , 184, 861-864	4.5	7
14	Need of voriconazole high dosages, with documented cerebrospinal fluid penetration, for treatment of cerebral aspergillosis in a 6-month-old leukaemic girl. <i>Journal of Chemotherapy</i> , 2017 , 29, 42-44	2.3	4

13	Unusual Late-onset Enteropathy in a Patient With Lipopolysaccharide-responsive Beige-like Anchor Protein Deficiency. <i>Journal of Pediatric Hematology/Oncology</i> , 2020 , 42, e768-e771	1.2	4
12	Hepatic veno-occlusive disease during isavuconazole administration. <i>Journal of Chemotherapy</i> , 2018 , 30, 63-64	2.3	3
11	The challenge of early diagnosis of autoimmune lymphoproliferative syndrome in children with suspected autoinflammatory/autoimmune disorders. <i>Rheumatology</i> , 2021 ,	3.9	3
10	Genetic screening of children with marrow failure. The role of primary Immunodeficiencies. <i>American Journal of Hematology</i> , 2021 , 96, 1077-1086	7.1	3
9	Thrombotic thrombocytopenic purpura and defective apoptosis due to CASP8/10 mutations: the role of mycophenolate mofetil. <i>Blood Advances</i> , 2019 , 3, 3432-3435	7.8	3
8	Case Report: Deficiency of Adenosine Deaminase 2 Presenting With Overlapping Features of Autoimmune Lymphoproliferative Syndrome and Bone Marrow Failure. <i>Frontiers in Immunology</i> , 2021 , 12, 754029	8.4	2
7	Comment on: Invasive fungal infections in children with acute lymphoblastic leukemia. <i>Pediatric Blood and Cancer</i> , 2020 , 67, e28035	3	2
6	Tocilizumab may slow radiographic progression in patients with systemic or polyarticular-course juvenile idiopathic arthritis: post hoc radiographic analysis from two randomized controlled trials. <i>Arthritis Research and Therapy</i> , 2020 , 22, 211	5.7	2
5	Underlying CTLA4 Deficiency in a Patient With Juvenile Idiopathic Arthritis and Autoimmune Lymphoproliferative Syndrome Features Successfully Treated With Abatacept-A Case Report. <i>Journal of Pediatric Hematology/Oncology</i> , 2021 , 43, e1168-e1172	1.2	2
4	Intravenous isavuconazole can be administered 5 days-a-week. A possibility suggested by a real-life observation. <i>Journal of Chemotherapy</i> , 2020 , 32, 217-218	2.3	1
3	Targeted NGS Yields Plentiful Ultra-Rare Variants in Inborn Errors of Immunity Patients. <i>Genes</i> , 2021 , 12,	4.2	1
2	Autoimmune Lymphoproliferative Syndrome (ALPS) and ALPS-Related Disorders. Different Bio-Clinical Profile and Similar Response to Treatment: A Single Centre Experience. <i>Blood</i> , 2015 , 126, 4618-4618	2.2	0
1	Genetic Screening of Patients with Evans Syndrome: A Single Centre Analysis. <i>Blood</i> , 2021 , 138, 4198-4198		