Elena Verrecchia

List of Publications by Year
in descending order

36 papers	722 citations	758635 12 h-index	580395 25 g-index
36 all docs	36 docs citations	36 times ranked	1022 citing authors

#	Article	IF	Citations
1	Familial Mediterranean Fever: A review for clinical management. Joint Bone Spine, 2009, 76, 227-233.	0.8	99
2	Caveats and truths in genetic, clinical, autoimmune and autoinflammatory issues in Blau syndrome and early onset sarcoidosis. Autoimmunity Reviews, 2014, 13, 1220-1229.	2.5	86
3	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.	1.6	72
4	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.	1.0	64
5	Giant cell arteritis and polymyalgia rheumatica after influenza vaccination: report of 10 cases and review of the literature. Lupus, 2012, 21, 153-157.	0.8	61
6	Diagnostic Criteria for Adult-Onset Periodic Fever, Aphthous Stomatitis, Pharyngitis, and Cervical Adenitis (PFAPA) Syndrome. Frontiers in Immunology, 2017, 8, 1018.	2.2	37
7	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.	1.2	31
8	Efficacy of etanercept in the treatment of a patient with BehÃ§etâ€™s disease. Clinical Rheumatology, 2008, 27, 933-936.	1.0	27
9	Clinical hints to diagnosis of attenuated forms of Mucopolysaccharidoses. Italian Journal of Pediatrics, 2018, 44, 132.	1.0	25
10	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.	1.4	24
11	Macular Impairment in Fabry Disease: A Morpho-functional Assessment by Swept-Source OCT Angiography and Focal Electroretinography. , 2019, 60, 2667.		23
12	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.	1.4	21
13	Children and Adults with PFAPA Syndrome: Similarities and Divergences in a Real-Life Clinical Setting. Advances in Therapy, 2021, 38, 1078-1093.	1.3	15
14	Small Intestinal Bacterial Overgrowth Affects the Responsiveness to Colchicine in Familial Mediterranean Fever. Mediators of Inflammation, 2017, 2017, 1-6.	1.4	14
15	Prognostic significance of right ventricular hypertrophy and systolic function in Andersonâ€“Fabry disease. ESC Heart Failure, 2020, 7, 1605-1614.	1.4	14
16	Functional and pharmacological evaluation of novel GLA variants in Fabry disease identifies six (two) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 2018, 481, 25-33.	0.5	13
17	New insights from the application of the FAbry STabilization indEX in a large population of Fabry cases. CKJ: Clinical Kidney Journal, 2019, 12, 65-70.	1.4	10
18	Right ventricular strain in Anderson-Fabry disease. International Journal of Cardiology, 2021, 330, 84-90.	0.8	10

#	Article	IF	Citations
19	The impact of fever/hyperthermia in the diagnosis of Fabry: A retrospective analysis. European Journal of Internal Medicine, 2016, 32, 26-30.	1.0	9
20	Mesothelioma in Familial Mediterranean Fever With Colchicine Intolerance: A Case Report and Literature Review. Frontiers in Immunology, 2020, 11, 889.	2.2	8
21	Effectiveness of Colchicine Therapy in 4 Cases of Retroperitoneal Fibrosis Associated with Autoinflammatory Diseases. Journal of Rheumatology, 2010, 37, 1971-1972.	1.0	7
22	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.	1.0	7
23	Resolution of femoral metaphyseal dysplasia in CINCA syndrome after long-term treatment with interleukin-1 blockade. Clinical Rheumatology, 2018, 37, 2007-2009.	1.0	7
24	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.	1.2	6
25	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.	1.0	6
26	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.	0.3	5
27	Caregiversâ€™ and Physiciansâ€™ Perspectives on Alpha-Mannosidosis: A Report from Italy. Advances in Therapy, 2021, 38, 1-10.	1.3	4
28	Drug survival of anakinra and canakinumab in monogenic autoinflammatory diseases: observational study from the International AIDA Registry. Rheumatology, 2021, 60, 5705-5712.	0.9	4
29	Wegenerâ€™s granulomatosis: an update on diagnosis and therapy. Expert Review of Clinical Immunology, 2008, 4, 481-495.	1.3	3
30	Switch from anakinra to canakinumab in a severe case of CINCA syndrome. International Journal of Rheumatic Diseases, 2016, 19, 1354-1356.	0.9	3
31	Folic acid supplementation during methotrexate treatment: nonsense?. British Journal of Rheumatology, 2005, 44, 563-564.	2.5	2
32	Colchicine trial in PFAPA Syndrome and MEFV-negative patients. Pediatric Rheumatology, 2015, 13, .	0.9	2
33	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.	0.5	2
34	La maladie pÃ©riodique. Revue Du Rhumatisme (Edition Francaise), 2009, 76, 382-389.	0.0	1
35	PFAPA syndrome as an hereditary autoinflamatory disorder. Pediatric Rheumatology, 2015, 13, .	0.9	0
36	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.	0.2	0