

# Elena Verrecchia

## List of Publications by Year in descending order

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Version: 2024-02-01

36  
papers

722  
citations

759233

12  
h-index

580821

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. <i>Joint Bone Spine</i> , 2009, 76, 227-233.	1.6	99
2	Caveats and truths in genetic, clinical, autoimmune and autoinflammatory issues in Blau syndrome and early onset sarcoidosis. <i>Autoimmunity Reviews</i> , 2014, 13, 1220-1229.	5.8	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. <i>Frontiers in Pharmacology</i> , 2016, 7, 380.	3.5	72
4	Safety profile of the interleukin-1 inhibitors anakinra and canakinumab in real-life clinical practice: a nationwide multicenter retrospective observational study. <i>Clinical Rheumatology</i> , 2018, 37, 2233-2240.	2.2	64
5	Giant cell arteritis and polymyalgia rheumatica after influenza vaccination: report of 10 cases and review of the literature. <i>Lupus</i> , 2012, 21, 153-157.	1.6	61
6	Diagnostic Criteria for Adult-Onset Periodic Fever, Aphthous Stomatitis, Pharyngitis, and Cervical Adenitis (PFAPA) Syndrome. <i>Frontiers in Immunology</i> , 2017, 8, 1018.	4.8	37
7	Right Ventricular Hypertrophy, Systolic Function, and Disease Severity in Anderson-Fabry Disease: An Echocardiographic Study. <i>Journal of the American Society of Echocardiography</i> , 2017, 30, 282-291.	2.8	31
8	Efficacy of etanercept in the treatment of a patient with Behçet's disease. <i>Clinical Rheumatology</i> , 2008, 27, 933-936.	2.2	27
9	Clinical hints to diagnosis of attenuated forms of Mucopolysaccharidoses. <i>Italian Journal of Pediatrics</i> , 2018, 44, 132.	2.6	25
10	Clinical Features at Onset and Genetic Characterization of Pediatric and Adult Patients with TNF- $\alpha$ Receptor-Associated Periodic Syndrome (TRAPS): A Series of 80 Cases from the AIDA Network. <i>Mediators of Inflammation</i> , 2020, 2020, 1-12.	3.0	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. <i>European Journal of Human Genetics</i> , 2019, 27, 1361-1368.	2.8	21
13	Children and Adults with PFAPA Syndrome: Similarities and Divergences in a Real-Life Clinical Setting. <i>Advances in Therapy</i> , 2021, 38, 1078-1093.	2.9	15
14	Small Intestinal Bacterial Overgrowth Affects the Responsiveness to Colchicine in Familial Mediterranean Fever. <i>Mediators of Inflammation</i> , 2017, 2017, 1-6.	3.0	14
15	Prognostic significance of right ventricular hypertrophy and systolic function in Anderson's Fabry disease. <i>ESC Heart Failure</i> , 2020, 7, 1605-1614.	3.1	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.	1.1	13
17	New insights from the application of the FABry STabilization indEX in a large population of Fabry cases. <i>CKJ: Clinical Kidney Journal</i> , 2019, 12, 65-70.	2.9	10
18	Right ventricular strain in Anderson-Fabry disease. <i>International Journal of Cardiology</i> , 2021, 330, 84-90.	1.7	10

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19	The impact of fever/hyperthermia in the diagnosis of Fabry: A retrospective analysis. <i>European Journal of Internal Medicine</i> , 2016, 32, 26-30.	2.2	9
20	Mesothelioma in Familial Mediterranean Fever With Colchicine Intolerance: A Case Report and Literature Review. <i>Frontiers in Immunology</i> , 2020, 11, 889.	4.8	8
21	Effectiveness of Colchicine Therapy in 4 Cases of Retroperitoneal Fibrosis Associated with Autoinflammatory Diseases. <i>Journal of Rheumatology</i> , 2010, 37, 1971-1972.	2.0	7
22	Recommendations for the inclusion of Fabry disease as a rare febrile condition in existing algorithms for fever of unknown origin. <i>Internal and Emergency Medicine</i> , 2017, 12, 1059-1067.	2.0	7
23	Resolution of femoral metaphyseal dysplasia in CINCA syndrome after long-term treatment with interleukin-1 blockade. <i>Clinical Rheumatology</i> , 2018, 37, 2007-2009.	2.2	7
24	Biotechnological Agents for Patients With Tumor Necrosis Factor Receptor Associated Periodic Syndrome—Therapeutic Outcome and Predictors of Response: Real-Life Data From the AIDA Network. <i>Frontiers in Medicine</i> , 2021, 8, 668173.	2.6	6
25	The Use of Chitotriosidase as a Marker of Active Sarcoidosis and in the Diagnosis of Fever of Unknown Origin (FUO). <i>Journal of Clinical Medicine</i> , 2021, 10, 5283.	2.4	6
26	Rapid resolution of severe pericardial effusion using anakinra in a patient with COVID-19 vaccine-related acute pericarditis relapse: a case report. <i>European Heart Journal - Case Reports</i> , 2022, 6, ytacl23.	0.6	5
27	Caregivers' and Physicians' Perspectives on Alpha-Mannosidosis: A Report from Italy. <i>Advances in Therapy</i> , 2021, 38, 1-10.	2.9	4
28	Drug survival of anakinra and canakinumab in monogenic autoinflammatory diseases: observational study from the International AIDA Registry. <i>Rheumatology</i> , 2021, 60, 5705-5712.	1.9	4
29	Wegener's granulomatosis: an update on diagnosis and therapy. <i>Expert Review of Clinical Immunology</i> , 2008, 4, 481-495.	3.0	3
30	Switch from anakinra to canakinumab in a severe case of CINCA syndrome. <i>International Journal of Rheumatic Diseases</i> , 2016, 19, 1354-1356.	1.9	3
31	Folic acid supplementation during methotrexate treatment: nonsense?. <i>British Journal of Rheumatology</i> , 2005, 44, 563-564.	2.3	2
32	Colchicine trial in PFAPA Syndrome and MEFV-negative patients. <i>Pediatric Rheumatology</i> , 2015, 13, .	2.1	2
33	Improvement of liver involvement in familial Mediterranean fever after introduction of canakinumab: a case report. <i>Mediterranean Journal of Hematology and Infectious Diseases</i> , 2020, 12, e20200059.	1.3	2
34	La maladie périodique. <i>Revue Du Rhumatisme (Edition Francaise)</i> , 2009, 76, 382-389.	0.0	1
35	PFAPA syndrome as an hereditary autoinflammatory disorder. <i>Pediatric Rheumatology</i> , 2015, 13, .	2.1	0
36	Anderson-Fabry's Disease: A Rare but Treatable Case of Fever of Unknown Origin. <i>European Journal of Case Reports in Internal Medicine</i> , 2017, 2, 000645.	0.4	0