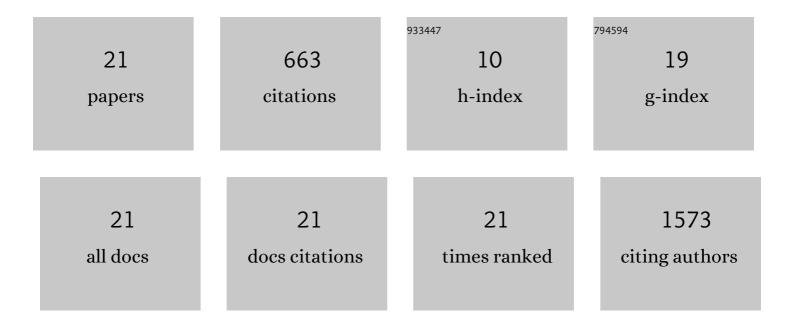
Erica Rurali

List of Publications by Year in descending order

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FRICA RUDALI

#	Article	IF	CITATIONS
1	Combined Complement Gene Mutations in Atypical Hemolytic Uremic Syndrome Influence Clinical Phenotype. Journal of the American Society of Nephrology: JASN, 2013, 24, 475-486.	6.1	308
2	A Genome-Wide Association Study of Diabetic Kidney Disease in Subjects With Type 2 Diabetes. Diabetes, 2018, 67, 1414-1427.	0.6	136
3	Vascular smooth muscle cells in Marfan syndrome aneurysm: the broken bricks in the aortic wall. Cellular and Molecular Life Sciences, 2017, 74, 267-277.	5.4	41
4	ADAMTS13 Predicts Renal and Cardiovascular Events in Type 2 Diabetic Patients and Response to Therapy. Diabetes, 2013, 62, 3599-3609.	0.6	25
5	Treatment of Congenital Thrombotic Thrombocytopenic Purpura With Eculizumab. American Journal of Kidney Diseases, 2015, 66, 1067-1070.	1.9	25
6	Cardiac fibrosis in regenerative medicine: destroy to rebuild. Journal of Thoracic Disease, 2018, 10, S2376-S2389.	1.4	15
7	Precise Therapy for Thoracic Aortic Aneurysm in Marfan Syndrome: A Puzzle Nearing Its Solution. Progress in Cardiovascular Diseases, 2018, 61, 328-335.	3.1	15
8	BM ageing: Implication for cell therapy with EPCs. Mechanisms of Ageing and Development, 2016, 159, 4-13.	4.6	14
9	Linking cell function with perfusion: insights from the transcatheter delivery of bone marrow-derived CD133+ cells in ischemic refractory cardiomyopathy trial (RECARDIO). Stem Cell Research and Therapy, 2018, 9, 235.	5.5	14
10	ADAMTS13 Secretion and Residual Activity among Patients with Congenital Thrombotic Thrombocytopenic Purpura with and without Renal Impairment. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 2002-2012.	4.5	12
11	Impact of a Complement Factor H Gene Variant on Renal Dysfunction, Cardiovascular Events, and Response to ACE Inhibitor Therapy in Type 2 Diabetes. Frontiers in Genetics, 2019, 10, 681.	2.3	11
12	Cyclophilin A/EMMPRIN Axis Is Involved in Pro-Fibrotic Processes Associated with Thoracic Aortic Aneurysm of Marfan Syndrome Patients. Cells, 2020, 9, 154.	4.1	11
13	Soluble EMMPRIN levels discriminate aortic ectasia in Marfan syndrome patients. Theranostics, 2019, 9, 2224-2234.	10.0	9
14	Congenital thrombotic thrombocytopenic purpura (cTTP) with two novel mutations. Pediatric Blood and Cancer, 2012, 59, 1296-1298.	1.5	6
15	New Strategies to Enhance Myocardial Regeneration: Expectations and Challenges from Preclinical Evidence. Current Stem Cell Research and Therapy, 2020, 15, 696-710.	1.3	6
16	Diabetes Induces a Transcriptional Signature in Bone Marrow–Derived CD34+ Hematopoietic Stem Cells Predictive of Their Progeny Dysfunction. International Journal of Molecular Sciences, 2021, 22, 1423.	4.1	5
17	Hereditary ADAMTS 13 deficiency presenting as recurrent acute kidney injury. Indian Journal of Nephrology, 2012, 22, 298.	0.5	4
18	Cyclophilin A in Arrhythmogenic Cardiomyopathy Cardiac Remodeling. International Journal of Molecular Sciences, 2019, 20, 2403.	4.1	4

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#	Article	IF	CITATIONS
19	Son of a Lesser God: The Case of Cell Therapy for Refractory Angina. Frontiers in Cardiovascular Medicine, 2021, 8, 709795.	2.4	2
20	Sensitive and quantitative method to evaluate DNA methylation of the positive regulatory domains (PRDI, PRDII) and cAMP response element (CRE) in human endothelial nitric oxide synthase promoter. Nitric Oxide - Biology and Chemistry, 2019, 92, 41-48.	2.7	0
21	Novel Targets for Old and Diseased Hearts. International Journal of Molecular Sciences, 2022, 23, 6627.	4.1	Ο