Rossella Piras

List of Publications by Year
in descending order

19	1,572	759233 12	794594 19
papers	citations	h-index	g-index

21	21	21	1929
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	CFH and CFHR Copy Number Variations in C3 Glomerulopathy and Immune Complex-Mediated Membranoproliferative Glomerulonephritis. Frontiers in Genetics, 2021, 12, 670727.	2.3	11
2	Eculizumab in patients with severe coronavirus disease 2019 (COVID-19) requiring continuous positive airway pressure ventilator support: Retrospective cohort study. PLoS ONE, 2021, 16, e0261113.	2.5	25
3	Molecular Studies and an ex vivo Complement Assay on Endothelium Highlight the Genetic Complexity of Atypical Hemolytic Uremic Syndrome: The Case of a Pedigree With a Null CD46 Variant. Frontiers in Medicine, 2020, 7, 579418.	2.6	8
4	Morphofunctional Effects of C5 Convertase Blockade in Immune Complex-Mediated Membranoproliferative Glomerulonephritis: Report of Two Cases with Evidence of Terminal Complement Activation. Nephron, 2020, 144, 195-203.	1.8	4
5	Rare Functional Variants in Complement Genes and Anti-FH Autoantibodies-Associated aHUS. Frontiers in Immunology, 2019, 10, 853.	4.8	31
6	An ExÂVivo Test of Complement Activation on Endothelium for Individualized Eculizumab Therapy in Hemolytic Uremic Syndrome. American Journal of Kidney Diseases, 2019, 74, 56-72.	1.9	71
7	Cluster Analysis Identifies Distinct Pathogenetic Patterns in C3 Glomerulopathies/Immune Complexâ€“Mediated Membranoproliferative GN. Journal of the American Society of Nephrology: JASN, 2018, 29, 283-294.	6.1	89
8	Unraveling the Molecular Mechanisms Underlying Complement Dysregulation by Nephritic Factors in C3G and IC-MPGN. Frontiers in Immunology, 2018, 9, 2329.	4.8	37
9	Fibronectin glomerulopathy - A sporadic case with unusual clinical manifestation. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2017, 28, 1416.	0.3	3
10	Thrombotic microangiopathy without renal involvement: two novel mutations in complementâ€regulator genes. Journal of Thrombosis and Haemostasis, 2016, 14, 340-345.	3.8	6
11	Nephrotic-Range Proteinuria and Peripheral Edema in a Child: Not Only Idiopathic Nephrotic Syndrome. Case Reports in Nephrology and Dialysis, 2016, 6, 120-127.	0.6	7
12	Complement gene variants determine the risk of immunoglobulin-associated MPGN and C3 glomerulopathy and predict long-term renal outcome. Molecular Immunology, 2016, 71, 131-142.	2.2	126
13	Characterization of a New DGKE Intronic Mutation in Genetically Unsolved Cases of Familial Atypical Hemolytic Uremic Syndrome. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 1011-1019.	4.5	47
14	Atypical hemolytic uremic syndrome with MCP mutations preceded by respiratory infection. CEN Case Reports, 2013, 2, 34-37.	0.9	1
15	A Case of Familial Glomerulopathy With Fibronectin Deposits Caused by the Y973C Mutation in Fibronectin. American Journal of Kidney Diseases, 2013, 61, 514-518.	1.9	22
16	Postâ€transplant recurrence of atypical hemolytic uremic syndrome in a patient with thrombomodulin mutation. Pediatric Transplantation, 2013, 17, E177-81.	1.0	23
17	Fibronectin Glomerulopathy: An Unusual Cause of Adult-Onset Nephrotic Syndrome. American Journal of Kidney Diseases, 2012, 60, 839-842.	1.9	22
18	MYO1EMutations and Childhood Familial Focal Segmental Glomerulosclerosis. New England Journal of Medicine, 2011, 365, 295-306.	27.0	221

#	Article	IF	Citations
19	Relative Role of Genetic Complement Abnormalities in Sporadic and Familial aHUS and Their Impact on Clinical Phenotype. Clinical Journal of the American Society of Nephrology: CJASN, 2010, 5, 1844-1859.	4.5	818