Sara Gamba

List of Publications by Year
in descending order

13 papers	2,826 citations	840776 11 h-index	1125743 13 g-index
13 all docs	13 docs citations	13 times ranked	2304 citing authors

#	Article	IF	Citations
1	C5a and C5aR1 are key drivers of microvascular platelet aggregation in clinical entities spanning from aHUS to COVID-19. Blood Advances, 2022, 6, 866-881.	5.2	31
2	C5 Convertase Blockade in Membranoproliferative Glomerulonephritis: A Single-Arm Clinical Trial. American Journal of Kidney Diseases, 2019, 74, 224-238.	1.9	45
3	Octreotide-LAR in later-stage autosomal dominant polycystic kidney disease (ALADIN 2): A randomized, double-blind, placebo-controlled, multicenter trial. PLoS Medicine, 2019, 16, e1002777.	8.4	42
4	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
5	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
6	A Multidrug, Antiproteinuric Approach to Alport Syndrome: A Ten-Year Cohort Study. Nephron, 2015, 130, 13-20.	1.8	9
7	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
8	Genetics of HUS: the impact of MCP, CFH, and IF mutations on clinical presentation, response to treatment, and outcome. Blood, 2006, 108, 1267-1279.	1.4	652
9	Direct medical costs of monitoring and treating patients with Takayasu arteritis in Italy. European Journal of Health Economics, 2004, 5, 330-334.	2.8	4
10	Familial haemolytic uraemic syndrome and an MCP mutation. Lancet, The, 2003, 362, 1542-1547.	13.7	303
11	Complement factor H mutations and gene polymorphisms in haemolytic uraemic syndrome: the C-257T, the A2089G and the G2881T polymorphisms are strongly associated with the disease. Human Molecular Genetics, 2003, 12, 3385-3395.	2.9	291
12	von Willebrand factor cleaving protease (ADAMTS13) is deficient in recurrent and familial thrombotic thrombocytopenic purpura and hemolytic uremic syndrome. Blood, 2002, 100, 778-785.	1.4	200
13	Interleukin-6 and RANTES in Takayasu Arteritis. Circulation, 1999, 100, 55-60.	1.6	216