Christelle Orlando

List of Publications by Year
in descending order

15 papers	183 citations	1163117 8 h-index	1125743 13 g-index
15 all docs	15 docs citations	15 times ranked	239 citing authors

#	Article	IF	Citations
1	Dialyzer Performance During Hemodialysis Without Systemic Anticoagulation Using a Heparin-Grafted Dialyzer Combined With a Citrate-Enriched Dialysate: Results of the Randomized Crossover Noninferiority EvoCit Study. American Journal of Kidney Diseases, 2022, 79, 79-87.e1.	1.9	11
2	Molecular Dissection of Structural Variations Involved in Antithrombin Deficiency. Journal of Molecular Diagnostics, 2022, 24, 462-475.	2.8	7
3	Antithrombin p.Thr147Ala: The First Founder Mutation in People of African Origin Responsible for Inherited Antithrombin Deficiency. Thrombosis and Haemostasis, 2021, 121, 182-191.	3.4	6
4	A pilot study on the impact of congenital thrombophilia in COVIDâ€19. European Journal of Clinical Investigation, 2021, 51, e13546.	3.4	16
5	Thrombotic storm under DOAC treatment in a patient with homozygous antithrombin Budapest III mutation. Thrombosis Research, 2021, 201, 161-163.	1.7	1
6	Recommendations for clinical laboratory testing for antithrombin deficiency; Communication from the SSC of the ISTH. Journal of Thrombosis and Haemostasis, 2020, 18, 17-22.	3.8	47
7	Routine haematology parameters in COVIDâ€19 patients and clinical outcome: A Belgian singleâ€centre study. International Journal of Laboratory Hematology, 2020, 42, e252-e255.	1.3	4
8	Belgian clinical guidance on anticoagulation management in hospitalised and ambulatory patients with COVID-19. Acta Clinica Belgica, 2020, , 1-6.	1.2	12
9	Hemodialysis Does Not Induce Detectable Activation of the Contact System of Coagulation. Kidney International Reports, 2020, 5, 831-838.	0.8	10
10	Incidence and features of thrombosis in children with inherited antithrombin deficiency. Haematologica, 2019, 104, 2512-2518.	3.5	21
11	Heterozygous FGA p.Asp473Ter (fibrinogen Nieuwegein) presenting as antepartum cerebral thrombosis. Thrombosis Research, 2018, 163, 185-189.	1.7	4
12	Childâ€onset thrombotic thrombocytopenic purpura caused by p.R498C and p.G259PfsX133 mutations in ADAMTS13. European Journal of Haematology, 2018, 101, 191-199.	2.2	4
13	Antithrombin heparin binding site deficiency: A challenging diagnosis of a not so benign thrombophilia. Thrombosis Research, 2015, 135, 1179-1185.	1.7	28
14	Homozygous Antithrombin Deficiency in Adolescents Presenting With Lower Extremity Thrombosis and Renal Complications. Journal of Pediatric Hematology/Oncology, 2014, 36, e190-e192.	0.6	8
15	Identification of two de novo mutations responsible for type I antithrombin deficiency. Thrombosis and Haemostasis, 2012, 107, 187-189.	3.4	4