

Christelle Orlando

List of Publications by Year in descending order

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Version: 2024-02-01

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	Recommendations for clinical laboratory testing for antithrombin deficiency; Communication from the SSC of the ISTH. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 17-22.	3.8	47
2	Antithrombin heparin binding site deficiency: A challenging diagnosis of a not so benign thrombophilia. <i>Thrombosis Research</i> , 2015, 135, 1179-1185.	1.7	28
3	Incidence and features of thrombosis in children with inherited antithrombin deficiency. <i>Haematologica</i> , 2019, 104, 2512-2518.	3.5	21
4	A pilot study on the impact of congenital thrombophilia in COVID-19. <i>European Journal of Clinical Investigation</i> , 2021, 51, e13546.	3.4	16
5	Belgian clinical guidance on anticoagulation management in hospitalised and ambulatory patients with COVID-19. <i>Acta Clinica Belgica</i> , 2020, , 1-6.	1.2	12
6	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. <i>American Journal of Kidney Diseases</i> , 2022, 79, 79-87.e1.	1.9	11
7	Hemodialysis Does Not Induce Detectable Activation of the Contact System of Coagulation. <i>Kidney International Reports</i> , 2020, 5, 831-838.	0.8	10
8	Homozygous Antithrombin Deficiency in Adolescents Presenting With Lower Extremity Thrombosis and Renal Complications. <i>Journal of Pediatric Hematology/Oncology</i> , 2014, 36, e190-e192.	0.6	8
9	Molecular Dissection of Structural Variations Involved in Antithrombin Deficiency. <i>Journal of Molecular Diagnostics</i> , 2022, 24, 462-475.	2.8	7
10	Antithrombin p.Thr147Ala: The First Founder Mutation in People of African Origin Responsible for Inherited Antithrombin Deficiency. <i>Thrombosis and Haemostasis</i> , 2021, 121, 182-191.	3.4	6
11	Identification of two de novo mutations responsible for type I antithrombin deficiency. <i>Thrombosis and Haemostasis</i> , 2012, 107, 187-189.	3.4	4
12	Heterozygous FGA p.Asp473Ter (fibrinogen Nieuwegein) presenting as antepartum cerebral thrombosis. <i>Thrombosis Research</i> , 2018, 163, 185-189.	1.7	4
13	Child-onset thrombotic thrombocytopenic purpura caused by p.R498C and p.G259PfsX133 mutations in ADAMTS13. <i>European Journal of Haematology</i> , 2018, 101, 191-199.	2.2	4
14	Routine haematology parameters in COVID-19 patients and clinical outcome: A Belgian single-centre study. <i>International Journal of Laboratory Hematology</i> , 2020, 42, e252-e255.	1.3	4
15	Thrombotic storm under DOAC treatment in a patient with homozygous antithrombin Budapest III mutation. <i>Thrombosis Research</i> , 2021, 201, 161-163.	1.7	1