## Alain Stépanian

List of Publications by Year in descending order

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49 papers

2,254 citations

331670 21 h-index 223800 46 g-index

55 all docs 55 docs citations

55 times ranked 3736 citing authors

#	Article	IF	Citations
1	Case series of massive direct oral anticoagulant ingestionâ€"Treatment and pharmacokinetics data. European Journal of Clinical Investigation, 2022, , e13746.	3.4	5
2	No VTE Recurrence After 1-Year Follow-Up of Hospitalized Patients With COVID-19 and a VTE Event. Chest, 2022, 162, 226-229.	0.8	6
3	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, e2200413119.	7.1	110
4	Efficacy of subcutaneous preemptive rituximab in immuneâ€mediated thrombotic thrombocytopenic purpura: Experience from the first 12 cases. American Journal of Hematology, 2021, 96, E26-E29.	4.1	5
5	The clinical picture of thrombotic microangiopathy in patients older than 60Âyears of age. British Journal of Haematology, 2021, 192, e25-e28.	2.5	3
6	Performance of Diagnostic Scores in Thrombotic Microangiopathy Patients in the Intensive Care Unit: A Monocentric Study. Thrombosis and Haemostasis, 2021, 121, 1427-1434.	3.4	2
7	D-Dimer Level and Neutrophils Count as Predictive and Prognostic Factors of Pulmonary Embolism in Severe Non-ICU COVID-19 Patients. Viruses, 2021, 13, 758.	3.3	19
8	Viscoelastometric Testing to Assess Hemostasis of COVID-19: A Systematic Review. Journal of Clinical Medicine, 2021, 10, 1740.	2.4	43
9	White blood count, Dâ€dimers, and ferritin levels as predictive factors of pulmonary embolism suspected upon admission in noncritically ill COVIDâ€19 patients: The French multicenter CLOTVID retrospective study. European Journal of Haematology, 2021, 107, 190-201.	2.2	12
10	Is platelet function testing at the acute phase under P2Y12 inhibitors helpful in predicting bleeding in real-life patients with acute coronary syndrome? The AVALANCHE study. Archives of Cardiovascular Diseases, 2021, 114, 612-623.	1.6	0
11	Coagulation disorders in patients with severe hemophagocytic lymphohistiocytosis. PLoS ONE, 2021, 16, e0251216.	2.5	12
12	Autoantibodies neutralizing type I IFNs are present in $\sim$ 4% of uninfected individuals over 70 years old and account for $\sim$ 20% of COVID-19 deaths. Science Immunology, 2021, 6, .	11.9	357
13	Diagnosis and followâ€up of thrombotic thrombocytopenic purpura with an automated chemiluminescent ADAMTS13 activity immunoassay. Research and Practice in Thrombosis and Haemostasis, 2021, 5, 81-93.	2.3	12
14	Contrast between Prevalence of HIT Antibodies and Confirmed HIT in Hospitalized COVID-19 Patients: A Prospective Study with Clinical Implications. Thrombosis and Haemostasis, 2021, 121, 971-975.	3.4	12
15	von Willebrand factor/ADAMTS13 axis and venous thromboembolism in moderateâ€toâ€severe COVIDâ€19 patients. British Journal of Haematology, 2021, 192, 1097-1100.	2.5	28
16	Are antiphospholipid antibodies associated with thrombotic complications in critically ill COVID-19 patients?. Thrombosis Research, 2020, 195, 74-76.	1.7	64
17	Plasma Serotonin is Elevated in Adult Patients with Sudden Sensorineural Hearing Loss. Thrombosis and Haemostasis, 2020, 120, 1291-1299.	3.4	12
18	High Prevalence of Deep Vein Thrombosis in Mechanically Ventilated COVID-19 Patients. Journal of the American College of Cardiology, 2020, 76, 480-482.	2.8	82

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19	Placental Overexpression of Soluble CORIN inÂPreeclampsia. American Journal of Pathology, 2020, 190, 970-976.	3.8	13
20	HLA-DRB1*11 is a strong risk factor for acquired thrombotic thrombocytopenic purpura in children. Haematologica, 2020, 105, e531.	3.5	5
21	Factor VIII and IX assays for postâ€infusion monitoring in hemophilia patients: Guidelines from the French BIMHO group (GFHT). European Journal of Haematology, 2020, 105, 103-115.	2.2	5
22	Pathophysiological Processes Underlying the High Prevalence of Deep Vein Thrombosis in Critically Ill COVID-19 Patients. Frontiers in Physiology, 2020, 11, 608788.	2.8	24
23	Loss of von Willebrand factor high-molecular-weight multimers at acute phase is associated with detectable anti-ADAMTS13 IgG and neurological symptoms in acquired thrombotic thrombocytopenic purpura. Thrombosis Research, 2019, 181, 29-35.	1.7	7
24	Multicentre evaluation of $\langle scp \rangle CK \langle scp \rangle Prest \langle sup \rangle \hat{A}^{\otimes} \langle sup \rangle$ for assaying plasma levels of factor IX fused with albumin (Idelvion $\langle sup \rangle \hat{A}^{\otimes} \langle sup \rangle$ ). Haemophilia, 2019, 25, e327-e330.	2.1	5
25	Potential usefulness of activated charcoal (DOAC remove $\hat{A}^{\circ}$ ) for dRVVT testing in patients receiving Direct Oral AntiCoagulants. Thrombosis Research, 2019, 184, 86-91.	1.7	30
26	Modified ROTEM for the detection of rivaroxaban and apixaban anticoagulant activity in whole blood. European Journal of Anaesthesiology, 2019, 36, 449-456.	1.7	22
27	Multifactorial hypercoagulable state associated with a thrombotic phenotype in phosphomannomutase-2 congenital disorder of glycosylation (PMM2-CDG): Case report and brief review of the literature. Thrombosis Research, 2019, 178, 75-78.	1.7	4
28	Platelet Functions are Decreased in Obesity and Restored after Weight Loss: Evidence for a Role of the SERCA3-Dependent ADP Secretion Pathway. Thrombosis and Haemostasis, 2019, 119, 384-396.	3.4	13
29	Are Screening Tests Reliable to Rule Out Direct Oral Anticoagulant Plasma Levels at Various Thresholds (30,Â50, or 100Âng/mL) in Emergency Situations?. Chest, 2018, 153, 288-290.	0.8	15
30	ADAMTS13 Gene Mutations Influence ADAMTS13 Conformation and Disease Age-Onset in the French Cohort of Upshawâ€"Schulman Syndrome. Thrombosis and Haemostasis, 2018, 118, 1902-1917.	3.4	40
31	Comparison of risk prediction scores for venous thromboembolism in cancer patients: a prospective cohort study. Haematologica, 2017, 102, 1494-1501.	3.5	164
32	Fatal case of chikungunya and concomitant thrombotic thrombocytopenic purpura in French Guiana during air flight medical evacuation. Journal of Travel Medicine, 2017, 24, .	3.0	19
33	Epidemiology and pathophysiology of adulthood-onset thrombotic microangiopathy with severe ADAMTS13 deficiency (thrombotic thrombocytopenic purpura): a cross-sectional analysis of the French national registry for thrombotic microangiopathy. Lancet Haematology, the, 2016, 3, e237-e245.	4.6	218
34	Child-onset and adolescent-onset acquired thrombotic thrombocytopenic purpura with severe ADAMTS13 deficiency: a cohort study of the French national registry for thrombotic microangiopathy. Lancet Haematology,the, 2016, 3, e537-e546.	4.6	53
35	Apelin: an antithrombotic factor that inhibits platelet function. Blood, 2016, 127, 908-920.	1.4	45
36	Coagulation Disorders and Bleedings in Critically Ill Patients With Hemophagocytic Lymphohistiocytosis. Medicine (United States), 2015, 94, e1692.	1.0	62

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37	Highly Significant Association between Two Common Single Nucleotide Polymorphisms in CORIN Gene and Preeclampsia in Caucasian Women. PLoS ONE, 2014, 9, e113176.	2.5	31
38	Evaluation of a chromogenic commercial assay using VWF-73 peptide for ADAMTS13 activity measurement. Thrombosis Research, 2014, 134, 1074-1080.	1.7	28
39	Microparticle increase in severe obesity: Not related to metabolic syndrome and unchanged after massive weight loss. Obesity, 2013, 21, 2236-2243.	3.0	114
40	Evaluation of a commercial assay for ADAMTS13 activity measurement. Thrombosis and Haemostasis, 2013, 110, 852-853.	3.4	18
41	Unexpected frequency of Upshaw-Schulman syndrome in pregnancy-onset thrombotic thrombocytopenic purpura. Blood, 2012, 119, 5888-5897.	1.4	206
42	Elevated soluble endothelial cell protein C receptor (sEPCR) levels in women with preeclampsia: A marker of endothelial activation/damage?. Thrombosis Research, 2012, 129, 152-157.	1.7	22
43	Influence of ethnicity on the clinical and biologic expression of preâ€eclampsia in the ECLAXIR study. International Journal of Gynecology and Obstetrics, 2011, 115, 153-156.	2.3	7
44	Reference values for rotation thromboelastometry (ROTEM $\hat{A}^{@}$ ) parameters following non-haemorrhagic deliveries. Correlations with standard haemostasis parameters. Thrombosis and Haemostasis, 2011, 106, 176-178.	3.4	46
45	Von Willebrand Factor and ADAMTS13. Arteriosclerosis, Thrombosis, and Vascular Biology, 2011, 31, 1703-1709.	2.4	72
46	Cardiovascular and Thromboembolic Risk Factors in Idiopathic Sudden Sensorineural Hearing Loss: A Case-Control Study. Audiology and Neuro-Otology, 2011, 16, 55-66.	1.3	134
47	Search for an Association between V249I and T280M CX3CR1 Genetic Polymorphisms, Endothelial Injury and Preeclampsia: The ECLAXIR Study. PLoS ONE, 2009, 4, e6192.	2.5	15
48	Impaired dimerization of von Willebrand factor subunit due to mutation A2801D in the CK domain results in a recessive type 2A subtype IID von Willebrand disease. Thrombosis and Haemostasis, 2006, 95, 776-781.	3.4	10
49	A new mutation, S1285F, within the A1 loop of von Willebrand factor induces a conformational change in A1 loop with abnormal binding to platelet GPIb and botrocetin causing type 2M von Willebrand disease. British Journal of Haematology, 2003, 120, 643-651.	2.5	14