

Salam Salloum-Asfar

List of Publications by Year
in descending order

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Version: 2024-02-01

26
papers

412
citations

759233
12
h-index

752698
20
g-index

26
all docs

26
docs citations

26
times ranked

686
citing authors

#	ARTICLE	IF	CITATIONS
1	Rules all PIs should follow. Science, 2022, 376, 24-26.	12.6	0
2	Single Extracellular Vesicle Analysis Using Flow Cytometry for Neurological Disorder Biomarkers. Frontiers in Integrative Neuroscience, 2022, 16, .	2.1	5
3	Circulating Non-Coding RNAs as a Signature of Autism Spectrum Disorder Symptomatology. International Journal of Molecular Sciences, 2021, 22, 6549.	4.1	13
4	Human induced pluripotent stem cell line (QBRli013-A) derivation from a 6-year-old female diagnosed with Autism spectrum disorder (ASD) and intellectual disability (ID). Stem Cell Research, 2021, 56, 102500.	0.7	0
5	Hyperosmotic Stress Induces a Specific Pattern for Stress Granule Formation in Human-Induced Pluripotent Stem Cells. Stem Cells International, 2021, 2021, 1-19.	2.5	1
6	The Potential Role of COVID-19 in the Pathogenesis of Multiple Sclerosisâ€”A Preliminary Report. Viruses, 2021, 13, 2091.	3.3	14
7	Identification of potential transcription factors that enhance human iPSC generation. Scientific Reports, 2020, 10, 21950.	3.3	13
8	When genetic and surname analyses meet historical sources: The C56R mutation associated with factor XI deficiency as a marker of human migration during the Spanish Reconquista. Medical Hypotheses, 2020, 141, 109709.	1.5	0
9	Circulating miRNAs, Small but Promising Biomarkers for Autism Spectrum Disorder. Frontiers in Molecular Neuroscience, 2019, 12, 253.	2.9	40
10	Archeogenetics of F11 p.Cys38Arg: a 5400-year-old mutation identified in different southwestern European countries. Blood, 2019, 133, 2618-2622.	1.4	6
11	Mouse venous thrombosis upon silencing of anticoagulants depends on tissue factor and platelets, not FXII or neutrophils. Blood, 2019, 133, 2090-2099.	1.4	23
12	Gynaecological and obstetrical bleeding in Caucasian women with congenital factor XI deficiency: Results from a twenty-year, retrospective, observational study. Medicina Clínica (English Edition), 2019, 153, 373-379.	0.2	0
13	Gynaecological and obstetrical bleeding in Caucasian women with congenital factor XI deficiency: Results from a twenty-year, retrospective, observational study. Medicina Clínica, 2019, 153, 373-379.	0.6	6
14	Assessment of two contact activation reagents for the diagnosis of congenital factor XI deficiency. Thrombosis Research, 2018, 163, 64-70.	1.7	12
15	Heestermans M, Salloum-Asfar S, Salvatori D, Laghmani EH, Luken BM, Zeerleder SS, Spronk HMH, Korporaal SJ, Wagenaar GTM, Reitsma PH, van Vlijmen BJM. Role of platelets, neutrophils, and factor XII in spontaneous venous thrombosis in mice. Blood. 2016;127(21):2630-2637.. Blood, 2018, 131, 2996-2996.	1.4	4
16	Regulation of TFPI± expression by miR-27a/b-3p in human endothelial cells under normal conditions and in response to androgens. Scientific Reports, 2017, 7, 43500.	3.3	20
17	Identification of coagulation gene 3â€²UTR variants that are potentially regulated by microRNAs. British Journal of Haematology, 2017, 177, 782-790.	2.5	15
18	High incidence of <sc>FXI</sc> deficiency in a Spanish town caused by 11 different mutations and the first duplication of <i>F11</i>: Results from the Yecla study. Haemophilia, 2017, 23, e488-e496.	2.1	22

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19	MiRNA-Based Regulation of Hemostatic Factors through Hepatic Nuclear Factor-4 Alpha. PLoS ONE, 2016, 11, e0154751.	2.5	19
20	Role of platelets, neutrophils, and factor XII in spontaneous venous thrombosis in mice. Blood, 2016, 127, 2630-2637.	1.4	26
21	The Immediate and Late Effects of Thyroid Hormone (Triiodothyronine) on Murine Coagulation Gene Transcription. PLoS ONE, 2015, 10, e0127469.	2.5	6
22	Peritoneal fluid modifies the microRNA expression profile in endometrial and endometriotic cells from women with endometriosis. Human Reproduction, 2015, 30, 2292-2302.	0.9	51
23	Regulation of Coagulation Factor XI Expression by MicroRNAs in the Human Liver. PLoS ONE, 2014, 9, e111713.	2.5	34
24	Prognostic role of MIR146A polymorphisms for cardiovascular events in atrial fibrillation. Thrombosis and Haemostasis, 2014, 112, 781-788.	3.4	36
25	Control of post-translational modifications in antithrombin during murine post-natal development by miR-200a. Journal of Biomedical Science, 2013, 20, 29.	7.0	10
26	miR-133a Regulates Vitamin K 2,3-Epoxy Reductase Complex Subunit 1 (VKORC1), a Key Protein in the Vitamin K Cycle. Molecular Medicine, 2012, 18, 1466-1472.	4.4	36