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	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
2	Circulating miRNAs, Small but Promising Biomarkers for Autism Spectrum Disorder. Frontiers in Molecular Neuroscience, 2019, 12, 253.	2.9	40
3	miR-133a Regulates Vitamin K 2,3-Epoxide Reductase Complex Subunit 1 (VKORC1), a Key Protein in the Vitamin K Cycle. Molecular Medicine, 2012, 18, 1466-1472.	4.4	36
4	Prognostic role of MIR146A polymorphisms for cardiovascular events in atrial fibrillation. Thrombosis and Haemostasis, 2014, 112, 781-788.	3.4	36
5	Regulation of Coagulation Factor XI Expression by MicroRNAs in the Human Liver. PLoS ONE, 2014, 9, e111713.	2.5	34
6	Role of platelets, neutrophils, and factor XII in spontaneous venous thrombosis in mice. Blood, 2016, 127, 2630-2637.	1.4	26
7	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
8	High incidence of <scp>FXI</scp> deficiency in a Spanish town caused by 11 different mutations and the first duplication of F11: Results from the Yecla study. Haemophilia, 2017, 23, e488-e496.	2.1	22
9	Regulation of TFPIÎ± 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
10	MiRNA-Based Regulation of Hemostatic Factors through Hepatic Nuclear Factor-4 Alpha. PLoS ONE, 2016, 11, e0154751.	2.5	19
11	Identification of coagulation gene 3â€²UTR variants that are potentially regulated by microRNAs. British Journal of Haematology, 2017, 177, 782-790.	2.5	15
12	The Potential Role of COVID-19 in the Pathogenesis of Multiple Sclerosisâ€”A Preliminary Report. Viruses, 2021, 13, 2091.	3.3	14
13	Identification of potential transcription factors that enhance human iPSC generation. Scientific Reports, 2020, 10, 21950.	3.3	13
14	Circulating Non-Coding RNAs as a Signature of Autism Spectrum Disorder Symptomatology. International Journal of Molecular Sciences, 2021, 22, 6549.	4.1	13
15	Assessment of two contact activation reagents for the diagnosis of congenital factor XI deficiency. Thrombosis Research, 2018, 163, 64-70.	1.7	12
16	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
17	The Immediate and Late Effects of Thyroid Hormone (Triiodothyronine) on Murine Coagulation Gene Transcription. PLoS ONE, 2015, 10, e0127469.	2.5	6
18	Archeogenetics of F11 p.Cys38Arg: a 5400-year-old mutation identified in different southwestern European countries. Blood, 2019, 133, 2618-2622.	1.4	6

#	Article	IF	Citations
19	Gynaecological and obstetrical bleeding in Caucasian women with congenital factor XI deficiency: Results from a twenty-year, retrospective, observational study. Medicina ClÃnica, 2019, 153, 373-379.	0.6	6
20	Single Extracellular Vesicle Analysis Using Flow Cytometry for Neurological Disorder Biomarkers. Frontiers in Integrative Neuroscience, 2022, 16, .	2.1	5
21	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
22	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
23	Gynaecological and obstetrical bleeding in Caucasian women with congenital factor XI deficiency: Results from a twenty-year, retrospective, observational study. Medicina ClÃnica (English Edition), 2019, 153, 373-379.	0.2	0
24	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
25	Human induced pluripotent stem cell line (QBRIi013-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
26	Rules all PIs should follow. Science, 2022, 376, 24-26.	12.6	0