Joan D Beckman

List of Publications by Year in descending order

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840776 713466 32 685 11 21 citations h-index g-index papers 32 32 32 1193 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Heme Degradation and Vascular Injury. Antioxidants and Redox Signaling, 2010, 12, 233-248.	5.4	196
2	Elevated hematocrit enhances platelet accumulation following vascular injury. Blood, 2017, 129, 2537-2546.	1.4	90
3	Heme oxygenase-1 gene delivery by Sleeping Beauty inhibits vascular stasis in a murine model of sickle cell disease. Journal of Molecular Medicine, 2010, 88, 665-675.	3.9	77
4	Regulation of Heme Oxygenase-1 Protein Expression by miR-377 in Combination with miR-217. Journal of Biological Chemistry, 2011, 286, 3194-3202.	3.4	76
5	Red blood cells modulate structure and dynamics of venous clot formation in sickle cell disease. Blood, 2019, 133, 2529-2541.	1.4	51
6	Inhaled carbon monoxide reduces leukocytosis in a murine model of sickle cell disease. American Journal of Physiology - Heart and Circulatory Physiology, 2009, 297, H1243-H1253.	3.2	47
7	Isolation and Characterization of Ovine Luteal Pericytes and Effects of Nitric Oxide on Pericyte Expression of Angiogenic Factors. Endocrine, 2006, 29, 467-476.	2.2	28
8	Endothelial TLR4 Expression Mediates Vaso-Occlusive Crisis in Sickle Cell Disease. Frontiers in Immunology, 2020, 11, 613278.	4.8	20
9	Bortezomib for Refractory Immune-Mediated Thrombocytopenia Purpura. American Journal of Therapeutics, 2018, 25, e270-e272.	0.9	19
10	Novel Pathophysiological Mechanisms of Thrombosis in Myeloproliferative Neoplasms. Current Hematologic Malignancy Reports, 2021, 16, 304-313.	2.3	14
11	Hematocrit and incidence of venous thromboembolism. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 422-428.	2.3	13
12	Integration of clinical parameters, genotype and epistaxis severity score to guide treatment for hereditary hemorrhagic telangiectasia associated bleeding. Orphanet Journal of Rare Diseases, 2020, 15, 185.	2.7	11
13	Challenges in diagnosis and management of acquired factor XIII (FXIII) inhibitors. Haemophilia, 2018, 24, e417-e420.	2.1	10
14	Soluble MD-2 and Heme in Sickle Cell Disease Plasma Promote Pro-Inflammatory Signaling in Endothelial Cells. Frontiers in Immunology, 2021, 12, 632709.	4.8	8
15	Factor XIII cotreatment with hemostatic agents in hemophilia A increases fibrin αâ€chain crosslinking. Journal of Thrombosis and Haemostasis, 2018, 16, 131-141.	3.8	6
16	Quantitative real-time polymerase chain reaction (qRT-PCR) restriction fragment length polymorphism (RFLP) method for monitoring highly conserved transgene expression during gene therapy. Translational Research, 2008, 152, 290-297.	5.0	4
17	High incidence of thromboembolism in patients with chronic GVHD: association with severity of GVHD and donor-recipient ABO blood group. Blood Cancer Journal, 2021, 11, 96.	6.2	4
18	Mechanistic rationale for factor XIII cotreatment in haemophilia. Haemophilia, 2019, 25, e377-e378.	2.1	3

#	Article	IF	CITATIONS
19	Ruxolitinib Reduces Endothelial Pro-Adhesive Interactions: Implications for JAK2V617+ MPN Thrombosis. Blood, 2020, 136, 1-1.	1.4	3
20	Gastrointestinal metastatic breast cancer unmasked by anticoagulation. Current Problems in Cancer Case Reports, 2020, 1, 100002.	0.1	2
21	Targeted sequencing of candidate gene regions for myelofibrosis in dogs. Journal of Veterinary Internal Medicine, 0, , .	1.6	2
22	Adult Langerhans histiocytosis with rare BRAF mutation complicated by massive pulmonary embolism. Thrombosis Research, 2020, 193, 207-210.	1.7	1
23	Transition of Human Embryonic Stem Cell-Derived Endothelial Cells to Smooth Muscle Cells in Culture as a Model for Vascular Development Blood, 2005, 106, 3683-3683.	1.4	0
24	Potential Role of Heme Oxygenase-1 in Arsenic Trioxide and Hemin-Mediated Differentiation of Human Promyelocytic Leukemia Cells. Blood, 2008, 112, 3981-3981.	1.4	0
25	Carbon Monoxide Decreases Leukocytosis in Murine Sickle Cell Disease Models Via Decreased Granulopoiesis Blood, 2008, 112, 1433-1433.	1.4	0
26	Heme Oxygenase-1 Gene Therapy in a Murine Model of Sickle Cell Disease Blood, 2009, 114, 1527-1527.	1.4	0
27	Carbon Monoxide Therapy Reduces Reactive Oxygen Species Production and the Short-Term Hematopoietic Stem Cell Population In Heme-Oxygenase-1 Knockout Mice. Blood, 2010, 116, 4767-4767.	1.4	O
28	Carbon Monoxide Therapy Modulates Hematopoietic Stem Cell Development in Heme-Oxygenase-1 Knockout Mice. Blood, 2011, 118, 1318-1318.	1.4	0
29	Effect of Genotype and Antifibrinolytic Therapy on the Severity of Epistaxis in Hereditary Hemorrhagic Telangiectasia. Blood, 2014, 124, 1515-1515.	1.4	O
30	Co-Administration of Factor XIII with Hemostatic Agents in Hemophilia Promotes Clot Stability and Composition. Blood, 2016, 128, 2589-2589.	1.4	0
31	Mechanism Underlying a Role for Factor XIII (FXIII) Polymorphism in Sickle Cell Disease-Associated Priapism. Blood, 2018, 132, 2361-2361.	1.4	0
32	Microfluidic Methods to Advance Mechanistic Understanding and Translational Research in Sickle Cell Disease. Translational Research, 2022, , .	5.0	0