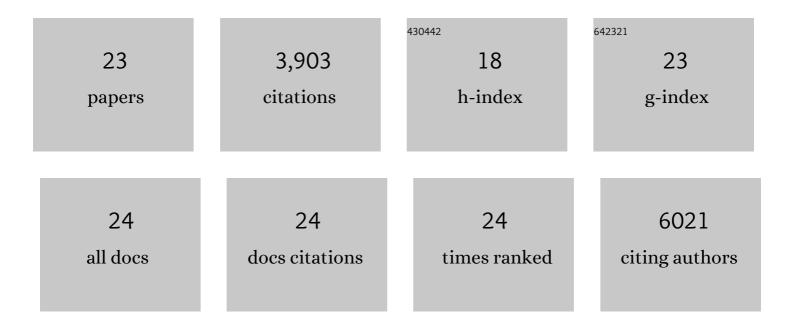
## Emilie Vénéreau

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/5549660/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Mutually exclusive redox forms of HMGB1 promote cell recruitment or proinflammatory cytokine release. Journal of Experimental Medicine, 2012, 209, 1519-1528.	4.2	590
2	HMGB1 promotes recruitment of inflammatory cells to damaged tissues by forming a complex with CXCL12 and signaling via CXCR4. Journal of Experimental Medicine, 2012, 209, 551-563.	4.2	539
3	DAMPs from Cell Death to New Life. Frontiers in Immunology, 2015, 6, 422.	2.2	500
4	Chronically Inflamed Human Tissues Are Infiltrated by Highly Differentiated Th17 Lymphocytes. Journal of Immunology, 2008, 180, 7423-7430.	0.4	470
5	Redox Modification of Cysteine Residues Regulates the Cytokine Activity of High Mobility Group Box-1 (HMGB1). Molecular Medicine, 2012, 18, 250-259.	1.9	378
6	Highâ€mobility group box 1 protein orchestrates responses to tissue damage via inflammation, innate and adaptive immunity, and tissue repair. Immunological Reviews, 2017, 280, 74-82.	2.8	281
7	HMGB1 as biomarker and drug target. Pharmacological Research, 2016, 111, 534-544.	3.1	214
8	HMGB1 and leukocyte migration during trauma and sterile inflammation. Molecular Immunology, 2013, 55, 76-82.	1.0	189
9	Oncostatin M Secreted by Skin Infiltrating T Lymphocytes Is a Potent Keratinocyte Activator Involved in Skin Inflammation. Journal of Immunology, 2007, 178, 4615-4622.	0.4	160
10	TLR4-mediated skin carcinogenesis is dependent on immune and radioresistant cells. EMBO Journal, 2010, 29, 2242-2252.	3.5	148
11	High mobility group box 1 orchestrates tissue regeneration via CXCR4. Journal of Experimental Medicine, 2018, 215, 303-318.	4.2	131
12	Aspirin's Active Metabolite Salicylic Acid Targets High Mobility Group Box 1 to Modulate Inflammatory Responses. Molecular Medicine, 2015, 21, 526-535.	1.9	97
13	HMGB1 is upregulated in the airways in asthma and potentiates airway smooth muscle contraction via TLR4. Journal of Allergy and Clinical Immunology, 2017, 140, 584-587.e8.	1.5	55
14	Molecular and Functional Characterization of a Soluble Form of Oncostatin M/Interleukin-31 Shared Receptor*. Journal of Biological Chemistry, 2006, 281, 36673-36682.	1.6	37
15	Rebalancing expression of HMGB1 redox isoforms to counteract muscular dystrophy. Science Translational Medicine, 2021, 13, .	5.8	26
16	Redox modifications of cysteine residues regulate the cytokine activity of HMGB1. Molecular Medicine, 2021, 27, 58.	1.9	25
17	Definition and Characterization of an Inhibitor for Interleukin-31. Journal of Biological Chemistry, 2010, 285, 14955-14963.	1.6	23
18	Oxidation of HMGB1 Is a Dynamically Regulated Process in Physiological and Pathological Conditions. Frontiers in Immunology, 2020, 11, 1122.	2.2	23

#	Article	IF	CITATIONS
19	Editorial: Seeing is not always believing: lessons from knockout mice. Journal of Leukocyte Biology, 2017, 101, 353-356.	1.5	4
20	Exploiting Live Imaging to Track Nuclei During Myoblast Differentiation and Fusion. Journal of Visualized Experiments, 2019, , .	0.2	4
21	Stress and Alarmins. Report from the 9th iD&EAs meeting. Cell Death and Disease, 2019, 10, 937.	2.7	3
22	Expression of Concern to: Redox modification of cysteine residues regulates the cytokine activity of high mobility group box-1 (HMGB1). Molecular Medicine, 2020, 26, 18.	1.9	3
23	Mutually exclusive redox forms of HMGB1 promote cell recruitment or proinflammatory cytokine release. Journal of General Physiology, 2012, 140, i3-i3.	0.9	0