Olivier D Christophe

List of Publications by Year
in descending order

70	2,850	186265 28	175258 52
papers	citations	h-index	g-index

72	72	72	3238
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	von Willebrand factor biosynthesis, secretion, and clearance: connecting the far ends. Blood, 2015, 125, 2019-2028.	1.4	296
2	Therapeutic levels of FVIII following a single peripheral vein administration of rAAV vector encoding a novel human factor VIII variant. Blood, 2013, 121, 3335-3344.	1.4	236
3	Emicizumab, a bispecific antibody recognizing coagulation factors IX and X: how does it actually compare to factor VIII?. Blood, 2017, 130, 2463-2468.	1.4	197
4	Antigen-selective modulation of AAV immunogenicity with tolerogenic rapamycin nanoparticles enables successful vector re-administration. Nature Communications, 2018, 9, 4098.	12.8	184
5	P-selectin glycoprotein ligand 1 and Î²2-integrins cooperate in the adhesion of leukocytes to von Willebrand factor. Blood, 2006, 108, 3746-3752.	1.4	152
6	A role for exposed mannosylations in presentation of human therapeutic self-proteins to CD4+ T lymphocytes. Proceedings of the National Academy of Sciences of the United States of America, 2007, 104, 8965-8970.	7.1	110
7	Macrophage LRP1 contributes to the clearance of von Willebrand factor. Blood, 2012, 119, 2126-2134.	1.4	99
8	Enhanced liver gene transfer and evasion of preexisting humoral immunity with exosome-enveloped AAV vectors. Blood Advances, 2017, 1, 2019-2031.	5.2	90
9	Liver-Specific Transcriptional Modules Identified by Genome-Wide In Silico Analysis Enable Efficient Gene Therapy in Mice and Non-Human Primates. Molecular Therapy, 2014, 22, 1605-1613.	8.2	71
10	Factor VIII and von Willebrand factor are ligands for the carbohydrate-receptor Siglec-5. Haematologica, 2012, 97, 1855-1863.	3.5	70
11	Accelerated uptake of VWF/platelet complexes in macrophages contributes to VWD type 2Bâ€“associated thrombocytopenia. Blood, 2013, 122, 2893-2902.	1.4	68
12	Altered thrombus formation in von Willebrand factorâ€“deficient mice expressing von Willebrand factor variants with defective binding to collagen or GPIbIIIa. Blood, 2008, 112, 603-609.	1.4	67
13	Ca2+ Binding to the First Epidermal Growth Factor-like Domain of Human Blood Coagulation Factor IX Promotes Enzyme Activity and Factor VIII Light Chain Binding. Journal of Biological Chemistry, 1996, 271, 25332-25337.	3.4	65
14	The interaction between factor H and VWF increases factor H cofactor activity and regulates VWF prothrombotic status. Blood, 2014, 123, 121-125.	1.4	63
15	Mutation and ADAMTS13-dependent modulation of disease severity in a mouse model for von Willebrand disease type 2B. Blood, 2010, 115, 4870-4877.	1.4	60
16	Identification of Galectin-1 and Galectin-3 as Novel Partners for Von Willebrand Factor. Arteriosclerosis, Thrombosis, and Vascular Biology, 2012, 32, 894-901.	2.4	59
17	Catalytic IgG from Patients with Hemophilia A Inactivate Therapeutic Factor VIII. Journal of Immunology, 2006, 177, 1355-1363.	0.8	45
18	Clearance of von Willebrand factor. Thrombosis and Haemostasis, 2008, 99, 271-278.	3.4	45

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19	von Willebrand factor mutation promotes thrombocytopathy by inhibiting integrin Î±IIbÎ²3. Journal of Clinical Investigation, 2013, 123, 5071-5081.	8.2	42
20	Blood Coagulation Factor IX Residues Glu78 and Arg94 Provide a Link between Both Epidermal Growth Factor-like Domains That Is Crucial in the Interaction with Factor VIII Light Chain. Journal of Biological Chemistry, 1998, 273, 222-227.	3.4	41
21	Proteolytic antibodies activate factor IX in patients with acquired hemophilia. Blood, 2011, 117, 2257-2264.	1.4	38
22	Antibody-based prevention of von Willebrand factor degradation mediated by circulatory assist devices. Thrombosis and Haemostasis, 2014, 112, 1014-1023.	3.4	37
23	ON THE VERSATILITY OF VON WILLEBRAND FACTOR. Mediterranean Journal of Hematology and Infectious Diseases, 2013, 5, e2013046.	1.3	36
24	Correction of Bleeding Symptoms in von Willebrand Factorâ€“Deficient Mice by Liver-Expressed von Willebrand Factor Mutants. Arteriosclerosis, Thrombosis, and Vascular Biology, 2008, 28, 419-424.	2.4	35
25	A Novel Single-Domain Antibody Against von Willebrand Factor A1 Domain Resolves Leukocyte Recruitment and Vascular Leakage During Inflammationâ€”Brief Report. Arteriosclerosis, Thrombosis, and Vascular Biology, 2017, 37, 1736-1740.	2.4	33
26	In Vivo Analysis of the Role of O-Glycosylations of Von Willebrand Factor. PLoS ONE, 2012, 7, e37508.	2.5	33
27	Macrophage scavenger receptor SR-AI contributes to the clearance of von Willebrand factor. Haematologica, 2018, 103, 728-737.	3.5	32
28	A hemophilia A mouse model for the in vivo assessment of emicizumab function. Blood, 2020, 136, 740-748.	1.4	32
29	Network-based analysis of omics data: the LEAN method. Bioinformatics, 2017, 33, 701-709.	4.1	29
30	Soluble Siglec-5 associates to PSGL-1 and displays anti-inflammatory activity. Scientific Reports, 2016, 6, 37953.	3.3	26
31	Role of the Gla and First Epidermal Growth Factor-like Domains of Factor X in the Prothrombinase and Tissue Factor-Factor VIIa Complexes. Journal of Biological Chemistry, 2003, 278, 10393-10399.	3.4	25
32	Mutations in the A3 domain of Von Willebrand factor inducing combined qualitative and quantitative defects in the protein. Blood, 2013, 121, 2135-2143.	1.4	25
33	Role of the Î±-Helix 163-170 in Factor Xa Catalytic Activity. Journal of Biological Chemistry, 2007, 282, 31569-31579.	3.4	23
34	LIM kinase/cofilin dysregulation promotes macrothrombocytopenia in severe von Willebrand disease-type 2B. JCI Insight, 2016, 1, e88643.	5.0	23
35	Surface Loop 199â€“204 in Blood Coagulation Factor IX Is a Cofactor-dependent Site Involved in Macromolecular Substrate Interaction. Journal of Biological Chemistry, 1999, 274, 29087-29093.	3.4	22
36	Hydrolysis of Coagulation Factors by Circulating IgG Is Associated with a Reduced Risk for Chronic Allograft Nephropathy in Renal Transplanted Patients. Journal of Immunology, 2008, 180, 8455-8460.	0.8	22

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37	Emerging Therapeutic Strategies in the Treatment of Hemophilia A. Seminars in Thrombosis and Hemostasis, 2017, 43, 581-590.	2.7	22
38	The VWF/LRP4/Î±VÎ²3-axis represents a novel pathway regulating proliferation of human vascular smooth muscle cells. Cardiovascular Research, 2022, 118, 622-637.	3.8	22
39	Singleâ€domain antibodies targeting antithrombin reduce bleeding in hemophilic mice with or without inhibitors. EMBO Molecular Medicine, 2020, 12, e11298.	6.9	20
40	Characterization of Recombinant von Willebrand Factors Mutated on Cysteine 509 or 695. Thrombosis and Haemostasis, 1996, 76, 453-459.	3.4	20
41	A genetically-engineered von Willebrand disease type 2B mouse model displays defects in hemostasis and inflammation. Scientific Reports, 2016, 6, 26306.	3.3	19
42	A factor VIIIâ€“nanobody fusion protein forming an ultrastable complex with VWF: effect on clearance and antibody formation. Blood, 2018, 132, 1193-1197.	1.4	19
43	Thermodynamic Analysis of the Interaction of Factor VIII with von Willebrand Factor. Biochemistry, 2012, 51, 4108-4116.	2.5	17
44	Kinesin-1 Is a New Actor Involved in Platelet Secretion and Thrombus Stability. Arteriosclerosis, Thrombosis, and Vascular Biology, 2018, 38, 1037-1051.	2.4	17
45	A murine model to characterize the antithrombotic effect of molecules targeting human von Willebrand factor. Blood, 2012, 120, 2723-2732.	1.4	16
46	Correction of bleeding in experimental severe hemophilia A by systemic delivery of factor VIII-encoding mRNA. Haematologica, 2020, 105, 1129-1137.	3.5	15
47	Targeting protease nexin-1, a natural anticoagulant serpin, to control bleeding and improve hemostasis in hemophilia. Blood, 2019, 134, 1632-1644.	1.4	14
48	In vitro recovery of FIX clotting activity as a marker of highly functional hepatocytes in a hemophilia B iPSC model. Hepatology, 2022, 75, 866-880.	7.3	12
49	Complex formation with pentraxin-2 regulates factor X plasma levels and macrophage interactions. Blood, 2017, 129, 2443-2454.	1.4	11
50	Development of a dual hybrid AAV vector for endothelial-targeted expression of von Willebrand factor. Gene Therapy, 2023, 30, 245-254.	4.5	11
51	Coagulation Factor X Interaction with Macrophages through Its N-Glycans Protects It from a Rapid Clearance. PLoS ONE, 2012, 7, e45111.	2.5	10
52	Protein kinase C signaling dysfunction in von Willebrand disease (p.V1316M) type 2B platelets. Blood Advances, 2018, 2, 1417-1428.	5.2	9
53	Macrophage receptor SR-AI is crucial to maintain normal plasma levels of coagulation factor X. Blood, 2016, 127, 778-786.	1.4	8
54	Camelidâ€derived singleâ€chain antibodies in hemostasis: Mechanistic, diagnostic, and therapeutic applications. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 1087-1110.	2.3	8

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55	Paraoxonase-1 expression is up-regulated in Down syndrome fetal liver. Biochemical and Biophysical Research Communications, 2006, 346, 1303-1306.	2.1	6
56	Expression of a structurally constrained von Willebrand factor variant triggers acute thrombotic thrombocytopenic purpura in mice. Blood, 2014, 123, 3344-3353.	1.4	6
57	Development and characterization of singleâ€domain antibodies neutralizing protease nexinâ€1 as tools to increase thrombin generation. Journal of Thrombosis and Haemostasis, 2020, 18, 2155-2168.	3.8	6
58	Biological outcome and mapping of total factor cascades in response to HIF induction during regenerative angiogenesis. Oncotarget, 2016, 7, 12102-12120.	1.8	6
59	A Thrombin-Activatable Factor X Variant Corrects Hemostasis in a Mouse Model for Hemophilia A. Thrombosis and Haemostasis, 2019, 119, 1981-1993.	3.4	5
60	In vivo modulation of a dominantâ€negative variant in mouse models of von Willebrand disease type 2A. Journal of Thrombosis and Haemostasis, 2021, 19, 139-146.	3.8	5
61	Apoptotic Platelet Events Are Not Observed in Severe von Willebrand Disease-Type 2B Mutation p.V1316M. PLoS ONE, 2015, 10, e0143896.	2.5	4
62	VON WILLEBRAND FACTOR ABNORMALITIES STUDIED IN THE MOUSE MODEL: WHAT WE LEARNED ABOUT VWF FUNCTIONS. Mediterranean Journal of Hematology and Infectious Diseases, 2013, 5, e2013047.	1.3	3
63	Removal of Mannose-Ending Glycan at Asn2118 Abrogates FVIII Presentation by Human Monocyte-Derived Dendritic Cells. Frontiers in Immunology, 2020, 11, 393.	4.8	3
64	A singleâ€domain antibody that blocks factorVIIa activity in the absence but not presence of tissue factor. Journal of Thrombosis and Haemostasis, 2019, 17, 2035-2046.	3.8	1
65	Identification of von Willebrand factor D4 domain mutations in patients of Afroâ€Caribbean descent: In vitro characterization. Research and Practice in Thrombosis and Haemostasis, 2022, 6, e12737.	2.3	1
66	Autologous cell/gene therapy approach of hemophilia B using patient specific induced Pluripotent Stem Cells. Journal of Hepatology, 2018, 68, S81-S82.	3.7	0
67	Siglecs as Novel Cellular Partners for Von Willebrand Factor. Blood, 2010, 116, 4306-4306.	1.4	0
68	Determinants of von Willebrand Factor Function. Blood, 2012, 120, SCI-17-SCI-17.	1.4	0
69	Mechanisms and Therapeutic Modulation of the Bleeding Tendency in Genetically-Engineered Von Willebrand Disease Type 2B Mice. Blood, 2015, 126, 3516-3516.	1.4	0
70	Antithrombotic potential of a singleâ€domain antibody enhancing the activated protein Câ€cofactor activity of protein S. Journal of Thrombosis and Haemostasis, 2022, , .	3.8	0