## Louis Aledort

List of Publications by Year in descending order

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	567281	265206
1,926	15	42
citations	h-index	g-index
132	132	1645
docs citations	times ranked	citing authors
	citations 132	1,926 15 citations h-index  132 132

#	Article	IF	CITATIONS
1	In response to WFH guidelines for the management of haemophilia, 3 <sup>rd</sup> edition: Is there a difference between extendedâ€halfâ€life FVIII products or not?. Haemophilia, 2021, 27, e762-e764.	2.1	O
2	Deaths Associated with Emicizumab in Patients with Hemophilia A. New England Journal of Medicine, 2019, 381, 1878-1879.	27.0	20
3	Why plasmaâ€derived factor VIII?. Haemophilia, 2019, 25, e183-e185.	2.1	O
4	The ongoing imperative for immune tolerance induction in inhibitor management. Haemophilia, 2019, 25, 183-186.	2.1	9
5	Factor VIII replacement is still the standard of care in haemophilia A. Blood Transfusion, 2019, 17, 479-486.	0.4	53
6	Application of phospho-CyTOF to characterize immune activation in patients with sickle cell disease in an ex vivo model of thrombosis. Journal of Immunological Methods, 2018, 453, 11-19.	1.4	11
7	Evaluating the safety of emicizumab in patients with hemophilia A. Expert Opinion on Drug Safety, 2018, 17, 1233-1237.	2.4	15
8	Prophylaxis reâ€visited: The potential impact of novel factor and nonâ€factor therapies on prophylaxis. Haemophilia, 2018, 24, 845-848.	2.1	5
9	An International Prophylaxis Study Group (IPSG) survey of prophylaxis in inhibitor positive children/adults with severe haemophilia. Haemophilia, 2017, 23, e444-e447.	2.1	3
10	An International Prophylaxis Study Group (IPSG) survey of prophylaxis in adults with severe haemophilia. Haemophilia, 2017, 23, e447-e450.	2.1	4
11	Treatment of Congenital Thrombotic Thrombocytopenia Purpura: A New Paradigm. Journal of Pediatric Hematology/Oncology, 2017, 39, 524-527.	0.6	14
12	The Role of Platelet Activation and Inflammation in Early Brain Injury Following Subarachnoid Hemorrhage. Neurocritical Care, 2017, 26, 48-57.	2.4	112
13	Hemophilia Lend-Lease Program in Italy Successfully Meets Albania Factor Needs. Blood, 2016, 128, 5916-5916.	1.4	1
14	Efficacy and safety of the thrombopoietin receptor agonist romiplostim in patients aged ≥65Âyears with immune thrombocytopenia. Annals of Hematology, 2015, 94, 1973-1980.	1.8	25
15	Factor VIII therapy for hemophilia A: current and future issues. Expert Review of Hematology, 2014, 7, 373-385.	2.2	27
16	A randomized trial of avatrombopag, an investigational thrombopoietin-receptor agonist, in persistent and chronic immune thrombocytopenia. Blood, 2014, 123, 3887-3894.	1.4	112
17	Managing incidentally diagnosed isolated factor VII deficiency perioperatively: a brief expert consensus report. Expert Review of Hematology, 2012, 5, 47-50.	2.2	8
18	Why should we care about quality of life in persons with haemophilia?. Haemophilia, 2012, 18, e154-7.	2.1	34

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19	Tertiary prophylaxis in adults: is there a rationale?. Haemophilia, 2012, 18, 722-728.	2.1	26
20	The use of a single von Willebrand factor-containing, plasma-derived FVIII product in hemophilia A immune tolerance induction: the US experience. Journal of Thrombosis and Haemostasis, 2011, 9, 2229-2234.	3.8	42
21	Acquired hemophilia: We now see it with myeloproliferative neoplasms. American Journal of Hematology, 2011, 86, 329-330.	4.1	8
22	<i>Expert Review of Hematology:</i> ddressing the needs of a broad research field. Expert Review of Hematology, 2008, 1, 1-2.	2.2	1
23	Hemophilia lives: the impact of prophylaxis. Blood, 2008, 111, 1752-1753.	1.4	1
24	Hemophilia Treatment - A Model for Appropriate Cost Control Blood, 2007, 110, 3942-3942.	1.4	0
25	Do hemophilia carriers bleed? Yes Blood, 2006, 108, 6-6.	1.4	12
26	The Clinical and Direct Medical Cost Burden of Splenectomy among Managed Care Patients with Chronic Immune Thrombocytopenic Purpura (ITP) Blood, 2006, 108, 5536-5536.	1.4	2
27	Retrospective Matched Cohort Study of Immune Thrombocytopenic Purpura (ITP): Complications Related to Corticosteroid (CS) Use Blood, 2006, 108, 3295-3295.	1.4	24
28	Plug the hole. Blood, 2005, 105, 2621-2621.	1.4	1
29	United States' factor XI-deficiency patients need a safer treatment. American Journal of Hematology, 2005, 80, 301-302.	4.1	12
30	Adverse Events (AEs): Are They Factored into Costs of Hemophilia Care? Blood, 2005, 106, 5550-5550.	1.4	0
31	Can Health Care Plans Afford Hemophilia Costs? Yes Blood, 2005, 106, 5551-5551.	1.4	O
32	Platelet bleeding disorders and approach to their management. Psychophysiology, 2005, 4, 85-7.	1.1	0
33	An Open Label, Multi Center Study on the Efficacy and Safety of a Liquid, Ready-to-Use Intravenously Administered Anti-D Immunoglobulin in Patients with Chronic Immune Thrombocytopenic Purpura Blood, 2004, 104, 3943-3943.	1.4	12
34	Do we need a better test? Can thrombin generation be useful?. Psychophysiology, 2004, 3, 75-6.	1.1	0
35	Orthopedic Outcome Studies and Cost Issues. Seminars in Thrombosis and Hemostasis, 2003, 29, 055-060.	2.7	4
36	Why Thrombin Generation? From Bench to Bedside. Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research, 2003, 33, 2-3.	0.3	12

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37	Can costs of hemophilia products be curtailed? Not as we do business today!. Thrombosis and Haemostasis, 2002, 88, 541.	3.4	1
38	Definitions in Hemophilia. Thrombosis and Haemostasis, 2001, 85, 560-560.	3.4	679
39	Adverse reactions related to rVIIa?. American Journal of Hematology, 2001, 67, 213-213.	4.1	4
40	The above letter was sent to Dr. Aledort, who offered the following reply Transfusion, 2000, 40, 495-496.	1.6	0
41	Prophylactic Use of Factor VIII: an Economic Evaluation. Thrombosis and Haemostasis, 1998, 79, 932-937.	3.4	88
42	von Willebrand Disease: From the Bedside to Therapy. Thrombosis and Haemostasis, 1997, 78, 562-565.	3.4	3
43	Inhibitors to Coagulation: Can We Afford Immune Tolerance Induction Regimens?. Vox Sanguinis, 1996, 70, 77-78.	1.5	3
44	Medical necessity-a threat to treatment of chronic disease. Transfusion, 1995, 35, 712-712.	1.6	0
45	Causes of death in haemophilia. Nature, 1995, 378, 124-124.	27.8	8
46	Some Aspects on the Management of Hemophilia. Thrombosis and Haemostasis, 1995, 74, 440-443.	3.4	6
46	Some Aspects on the Management of Hemophilia. Thrombosis and Haemostasis, 1995, 74, 440-443.  Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.	3.4	6
47	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.  Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. British Journal	1.5	9
47	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.  Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. British Journal of Haematology, 1990, 76, 238-241.  Human Recombinant DNA–Derived Antihemophilic Factor (Factor VIII) in the Treatment of Hemophilia A.	1.5 2.5	9 40
47 48 49	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.  Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. British Journal of Haematology, 1990, 76, 238-241.  Human Recombinant DNA–Derived Antihemophilic Factor (Factor VIII) in the Treatment of Hemophilia A. New England Journal of Medicine, 1990, 323, 1800-1805.	1.5 2.5 27.0	9 40 261
47 48 49 50	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.  Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. British Journal of Haematology, 1990, 76, 238-241.  Human Recombinant DNA–Derived Antihemophilic Factor (Factor VIII) in the Treatment of Hemophilia A. New England Journal of Medicine, 1990, 323, 1800-1805.  New Approaches to Management of Bleeding Disorders. Hospital Practice (1995), 1989, 24, 207-226.  Reproductive choices in hemophilic men and carriers. American Journal of Medical Genetics Part A,	1.5 2.5 27.0 1.0	9 40 261 8
47 48 49 50	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.  Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. British Journal of Haematology, 1990, 76, 238-241.  Human Recombinant DNA–Derived Antihemophilic Factor (Factor VIII) in the Treatment of Hemophilia A. New England Journal of Medicine, 1990, 323, 1800-1805.  New Approaches to Management of Bleeding Disorders. Hospital Practice (1995), 1989, 24, 207-226.  Reproductive choices in hemophilic men and carriers. American Journal of Medical Genetics Part A, 1987, 26, 591-598.  Concurrence of von Willebrand's disease and hemophilia A: Implications for carrier detection and	1.5 2.5 27.0 1.0	9 40 261 8

#	Article	IF	CITATIONS
55	Women and Von Willebrand Disease. , 0, , 296-301.		O
56	Products Used to Treat Hemophilia: Plasma-Derived Coagulation Factor Concentrates., 0,, 142-146.		O
57	Inhibitors to Factor VIII—Epidemiology and Treatment. , 0, , 64-70.		15
58	Von Willebrand Disease: Molecular Aspects. , 0, , 257-264.		0
59	Treatment of Inhibitors in Hemophilia B. , 0, , 101-105.		3
60	Inhibitors in Hemophilia B., 0,, 97-100.		16
61	Factor X and Factor X Deficiency. , 0, , 315-320.		1
62	Factor XI Deficiency., 0,, 321-327.		2
63	Inhibitors to Factor VIII—Molecular Basis. , 0, , 59-63.		5
64	Inhibitors to Factor VIII—Mild and Moderate Hemophilia. , 0, , 71-73.		4
65	Inhibitors to Factor VIII/IX: Treatment of Inhibitorsâ€"Immune Tolerance Induction. , 0, , 74-79.		8
66	Inhibitors of Factor VIII: Treatment of Acute Bleeds. , 0, , 80-85.		8
67	Acquired Inhibitors to Factor VIII., 0,, 86-90.		4
68	Hemophilia B—Molecular Basis. , 0, , 91-95.		2
69	Cellular Processing of Factors VIII and IX. , 0, , 5-12.		O
70	Work-Up of a Bleeding Child., 0,, 112-119.		0
71	Care of the Child with Hemophilia. , 0, , 120-124.		1
72	The Neonate with Hemophilia. , 0, , 125-130.		4

#	Article	IF	CITATIONS
73	Products used to Treat Hemophilia: Evolution of Treatment for Hemophilia A and B., 0,, 131-135.		5
74	Products used to Treat Hemophilia: Recombinant Products., 0,, 136-141.		O
75	Products used to Treat Hemophilia: Recombinant Factor VIIa. , 0, , 147-152.		3
76	Products used to Treat Hemophilia: Dosing. , 0, , 153-157.		1
77	Products Used to Treat Hemophilia: Regulation. , 0, , 158-163.		O
78	Joint Replacement., 0,, 164-168.		2
79	Work-Up of a Bleeding Adult., 0,, 13-18.		O
80	Synoviorthesis in Hemophilia. , 0, , 169-173.		4
81	Pseudotumors in Patients with Hemophilia. , 0, , 174-176.		4
82	Magnetic Resonance Imaging/Joint Outcome Assessment. , 0, , 182-192.		2
83	Physiotherapy in the Management of Hemophilia. , 0, , 193-199.		1
84	Transfusion Transmitted Disease: History of Epidemics (Focus on HIV)., 0,, 200-206.		3
85	Transfusion Transmitted Disease: Hepatitis C Virus Infection and Liver Transplantation., 0,, 207-213.		1
86	Gene Therapy: Introduction and Overview. , 0, , 214-219.		0
87	Gene Therapy for Hemophilia B. , 0, , 220-225.		O
88	Gene Therapy for Hemophilia A., 0,, 226-228.		3
89	Molecular Basis of Hemophilia A. , 0, , 19-26.		0
90	Gene Therapy: Molecular Engineering of Factor VIII and Factor IX., 0,, 229-234.		0

#	Article	IF	Citations
91	Laboratory Assays in Hemophilia. , 0, , 235-241.		6
92	Standardization of Assays. , 0, , 242-248.		3
93	Obstetrics and Gynecology: Hemophilia. , 0, , 249-256.		2
94	Von Willebrand Disease: Epidemiology. , 0, , 265-271.		5
95	Von Willebrand Disease: Biological Diagnosis. , 0, , 272-278.		1
96	Classification and Clinical Aspects of Von Willebrand Disease. , 0, , 279-284.		0
97	Treatment of Von Willebrand Disease: Desmopressin. , 0, , 285-288.		2
98	Treatment of Von Willebrand Disease: Therapeutic Concentrates., 0,, 289-295.		1
99	Hemophilia A: Role of Factor VIII in Coagulation. , 0, , 27-33.		O
100	Factor II., 0,, 302-305.		0
101	Factor V and Combined Factor V and VIII Deficiencies. , 0, , 306-310.		1
102	Congenital Factor VII Deficiency. , 0, , 311-314.		4
103	Factor XIII., 0, , 328-331.		0
104	Fibrinogen., 0,, 332-337.		0
105	Miscellaneous Rare Bleeding Disorders. , 0, , 338-344.		0
106	Quality of Life in Hemophilia. , 0, , 345-350.		4
107	Natural History of Inhibitor Development in Children with Severe Hemophilia a Treated with Factor VIII Products., 0,, 34-38.		2
108	The Economics of Hemophilia Treatments. , 0, , 351-358.		2

#	Article	IF	CITATIONS
109	Comprehensive Care and Delivery of Care: The Developed World. , 0, , 359-365.		2
110	Comprehensive Care and Delivery of Care: The Developing World., 0,, 366-370.		2
111	Comprehensive Care and Delivery of Care: The Global Perspective., 0,, 371-377.		1
112	Prophylaxis., 0,, 39-45.		1
113	Continuous Infusion of Coagulation Products in Hemophilia. , 0, , 46-52.		0
114	Inhibitors to Factor VIII—Immunology. , 0, , 53-58.		1
115	Colour Plate. , 0, , 409-412.		0
116	Overview of Hemostasis. , 0, , 1-4.		0