

Louis Aledort

List of Publications by Year in descending order

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116
papers

1,926
citations

567281

15
h-index

265206

42
g-index

132
all docs

132
docs citations

132
times ranked

1645
citing authors

#	ARTICLE	IF	CITATIONS
1	Definitions in Hemophilia. <i>Thrombosis and Haemostasis</i> , 2001, 85, 560-560.	3.4	679
2	Human Recombinant DNA-Derived Antihemophilic Factor (Factor VIII) in the Treatment of Hemophilia A. <i>New England Journal of Medicine</i> , 1990, 323, 1800-1805.	27.0	261
3	A randomized trial of avatrombopag, an investigational thrombopoietin-receptor agonist, in persistent and chronic immune thrombocytopenia. <i>Blood</i> , 2014, 123, 3887-3894.	1.4	112
4	The Role of Platelet Activation and Inflammation in Early Brain Injury Following Subarachnoid Hemorrhage. <i>Neurocritical Care</i> , 2017, 26, 48-57.	2.4	112
5	Prophylactic Use of Factor VIII: an Economic Evaluation. <i>Thrombosis and Haemostasis</i> , 1998, 79, 932-937.	3.4	88
6	Factor VIII replacement is still the standard of care in haemophilia A. <i>Blood Transfusion</i> , 2019, 17, 479-486.	0.4	53
7	The use of a single von Willebrand factor-containing, plasma-derived FVIII product in hemophilia A immune tolerance induction: the US experience. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 2229-2234.	3.8	42
8	Histocompatibility antigen patterns in haemophilic patients with factor VIII antibodies. <i>British Journal of Haematology</i> , 1990, 76, 238-241.	2.5	40
9	Why should we care about quality of life in persons with haemophilia?. <i>Haemophilia</i> , 2012, 18, e154-7.	2.1	34
10	Factor VIII therapy for hemophilia A: current and future issues. <i>Expert Review of Hematology</i> , 2014, 7, 373-385.	2.2	27
11	Tertiary prophylaxis in adults: is there a rationale?. <i>Haemophilia</i> , 2012, 18, 722-728.	2.1	26
12	Efficacy and safety of the thrombopoietin receptor agonist romiplostim in patients aged ≥65 years with immune thrombocytopenia. <i>Annals of Hematology</i> , 2015, 94, 1973-1980.	1.8	25
13	Retrospective Matched Cohort Study of Immune Thrombocytopenic Purpura (ITP): Complications Related to Corticosteroid (CS) Use.. <i>Blood</i> , 2006, 108, 3295-3295.	1.4	24
14	Reproductive choices in hemophilic men and carriers. <i>American Journal of Medical Genetics Part A</i> , 1987, 26, 591-598.	2.4	23
15	Deaths Associated with Emicizumab in Patients with Hemophilia A. <i>New England Journal of Medicine</i> , 2019, 381, 1878-1879.	27.0	20
16	Inhibitors in Hemophilia B. , 0, , 97-100.		16
17	Inhibitors to Factor VIII”Epidemiology and Treatment. , 0, , 64-70.		15
18	Evaluating the safety of emicizumab in patients with hemophilia A. <i>Expert Opinion on Drug Safety</i> , 2018, 17, 1233-1237.	2.4	15

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19	Current Concepts in Diagnosis and Management of Hemophilia. Hospital Practice (1995), 1982, 17, 77-92.	1.0	14
20	Treatment of Congenital Thrombotic Thrombocytopenia Purpura: A New Paradigm. Journal of Pediatric Hematology/Oncology, 2017, 39, 524-527.	0.6	14
21	Concurrence of von Willebrand's disease and hemophilia A: Implications for carrier detection and prevalence. American Journal of Medical Genetics Part A, 1986, 24, 83-94.	2.4	13
22	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
23	United States' factor XI-deficiency patients need a safer treatment. American Journal of Hematology, 2005, 80, 301-302.	4.1	12
24	Do hemophilia carriers bleed? Yes.. Blood, 2006, 108, 6-6.	1.4	12
25	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
26	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
27	AIDS: An Update. Hospital Practice (1995), 1983, 18, 159-171.	1.0	10
28	Hepatitis A Virus Transmission by Blood Products in the United States. Vox Sanguinis, 1994, 67, 24-28.	1.5	9
29	The ongoing imperative for immune tolerance induction in inhibitor management. Haemophilia, 2019, 25, 183-186.	2.1	9
30	New Approaches to Management of Bleeding Disorders. Hospital Practice (1995), 1989, 24, 207-226.	1.0	8
31	Causes of death in haemophilia. Nature, 1995, 378, 124-124.	27.8	8
32	Inhibitors to Factor VIII/IX: Treatment of Inhibitorsâ€™ Immune Tolerance Induction. , 0, , 74-79.		8
33	Inhibitors of Factor VIII: Treatment of Acute Bleeds. , 0, , 80-85.		8
34	Acquired hemophilia: We now see it with myeloproliferative neoplasms. American Journal of Hematology, 2011, 86, 329-330.	4.1	8
35	Managing incidentally diagnosed isolated factor VII deficiency perioperatively: a brief expert consensus report. Expert Review of Hematology, 2012, 5, 47-50.	2.2	8
36	Laboratory Assays in Hemophilia. , 0, , 235-241.		6

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37	Some Aspects on the Management of Hemophilia. <i>Thrombosis and Haemostasis</i> , 1995, 74, 440-443.	3.4	6
38	Inhibitors to Factor VIII—Molecular Basis. , 0, , 59-63.		5
39	Products used to Treat Hemophilia: Evolution of Treatment for Hemophilia A and B. , 0, , 131-135.		5
40	Von Willebrand Disease: Epidemiology. , 0, , 265-271.		5
41	Prophylaxis re-visited: The potential impact of novel factor and non-factor therapies on prophylaxis. <i>Haemophilia</i> , 2018, 24, 845-848.	2.1	5
42	Adverse reactions related to rVlla?. <i>American Journal of Hematology</i> , 2001, 67, 213-213.	4.1	4
43	Orthopedic Outcome Studies and Cost Issues. <i>Seminars in Thrombosis and Hemostasis</i> , 2003, 29, 055-060.	2.7	4
44	Inhibitors to Factor VIII—Mild and Moderate Hemophilia. , 0, , 71-73.		4
45	Acquired Inhibitors to Factor VIII. , 0, , 86-90.		4
46	The Neonate with Hemophilia. , 0, , 125-130.		4
47	Synoviorthesis in Hemophilia. , 0, , 169-173.		4
48	Pseudotumors in Patients with Hemophilia. , 0, , 174-176.		4
49	Congenital Factor VII Deficiency. , 0, , 311-314.		4
50	Quality of Life in Hemophilia. , 0, , 345-350.		4
51	An International Prophylaxis Study Group (IPSC) survey of prophylaxis in adults with severe haemophilia. <i>Haemophilia</i> , 2017, 23, e447-e450.	2.1	4
52	Inhibitors to Coagulation: Can We Afford Immune Tolerance Induction Regimens?. <i>Vox Sanguinis</i> , 1996, 70, 77-78.	1.5	3
53	Treatment of Inhibitors in Hemophilia B. , 0, , 101-105.		3
54	Products used to Treat Hemophilia: Recombinant Factor VIIa. , 0, , 147-152.		3

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55	Transfusion Transmitted Disease: History of Epidemics (Focus on HIV). , 0, , 200-206.		3
56	Gene Therapy for Hemophilia A. , 0, , 226-228.		3
57	Standardization of Assays. , 0, , 242-248.		3
58	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
59	von Willebrand Disease: From the Bedside to Therapy. Thrombosis and Haemostasis, 1997, 78, 562-565.	3.4	3
60	Factor XI Deficiency. , 0, , 321-327.		2
61	Hemophilia Bâ€™Molecular Basis. , 0, , 91-95.		2
62	Joint Replacement. , 0, , 164-168.		2
63	Magnetic Resonance Imaging/Joint Outcome Assessment. , 0, , 182-192.		2
64	Obstetrics and Gynecology: Hemophilia. , 0, , 249-256.		2
65	Treatment of Von Willebrand Disease: Desmopressin. , 0, , 285-288.		2
66	Natural History of Inhibitor Development in Children with Severe Hemophilia a Treated with Factor VIII Products. , 0, , 34-38.		2
67	The Economics of Hemophilia Treatments. , 0, , 351-358.		2
68	Comprehensive Care and Delivery of Care: The Developed World. , 0, , 359-365.		2
69	Comprehensive Care and Delivery of Care: The Developing World. , 0, , 366-370.		2
70	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
71	Plug the hole. Blood, 2005, 105, 2621-2621.	1.4	1
72	Factor X and Factor X Deficiency. , 0, , 315-320.		1

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73	Care of the Child with Hemophilia. , 0 , 120-124.		1
74	Products used to Treat Hemophilia: Dosing. , 0 , 153-157.		1
75	Physiotherapy in the Management of Hemophilia. , 0 , 193-199.		1
76	Transfusion Transmitted Disease: Hepatitis C Virus Infection and Liver Transplantation. , 0 , 207-213.		1
77	Von Willebrand Disease: Biological Diagnosis. , 0 , 272-278.		1
78	Treatment of Von Willebrand Disease: Therapeutic Concentrates. , 0 , 289-295.		1
79	Factor V and Combined Factor V and VIII Deficiencies. , 0 , 306-310.		1
80	Comprehensive Care and Delivery of Care: The Global Perspective. , 0 , 371-377.		1
81	Prophylaxis. , 0 , 39-45.		1
82	Inhibitors to Factor VIIIâ€™ Immunology. , 0 , 53-58.		1
83	<i>Expert Review of Hematology:</i>addressing the needs of a broad research field. Expert Review of Hematology, 2008, 1, 1-2.	2.2	1
84	Hemophilia lives: the impact of prophylaxis. Blood, 2008, 111, 1752-1753.	1.4	1
85	Hemophilia Lend-Lease Program in Italy Successfully Meets Albania Factor Needs. Blood, 2016, 128, 5916-5916.	1.4	1
86	Can costs of hemophilia products be curtailed? Not as we do business today!. Thrombosis and Haemostasis, 2002, 88, 541.	3.4	1
87	Medical necessity-a threat to treatment of chronic disease. Transfusion, 1995, 35, 712-712.	1.6	0
88	The above letter was sent to Dr. Aledort, who offered the following reply.. Transfusion, 2000, 40, 495-496.	1.6	0
89	Women and Von Willebrand Disease. , 0 , 296-301.		0
90	Products Used to Treat Hemophilia: Plasma-Derived Coagulation Factor Concentrates. , 0 , 142-146.		0

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91	Von Willebrand Disease: Molecular Aspects. , 0 , 257-264.		0
92	Cellular Processing of Factors VIII and IX. , 0 , 5-12.		0
93	Work-Up of a Bleeding Child. , 0 , 112-119.		0
94	Products used to Treat Hemophilia: Recombinant Products. , 0 , 136-141.		0
95	Products Used to Treat Hemophilia: Regulation. , 0 , 158-163.		0
96	Work-Up of a Bleeding Adult. , 0 , 13-18.		0
97	Gene Therapy: Introduction and Overview. , 0 , 214-219.		0
98	Gene Therapy for Hemophilia B. , 0 , 220-225.		0
99	Molecular Basis of Hemophilia A. , 0 , 19-26.		0
100	Gene Therapy: Molecular Engineering of Factor VIII and Factor IX. , 0 , 229-234.		0
101	Classification and Clinical Aspects of Von Willebrand Disease. , 0 , 279-284.		0
102	Hemophilia A: Role of Factor VIII in Coagulation. , 0 , 27-33.		0
103	Factor II. , 0 , 302-305.		0
104	Factor XIII. , 0 , 328-331.		0
105	Fibrinogen. , 0 , 332-337.		0
106	Miscellaneous Rare Bleeding Disorders. , 0 , 338-344.		0
107	Continuous Infusion of Coagulation Products in Hemophilia. , 0 , 46-52.		0
108	Colour Plate. , 0 , 409-412.		0

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109	Overview of Hemostasis. , 0, , 1-4.		0
110	Why plasma-derived factor VIII?. Haemophilia, 2019, 25, e183-e185.	2.1	0
111	In response to WFH guidelines for the management of haemophilia, 3 rd edition: Is there a difference between extended-half-life FVIII products or not?. Haemophilia, 2021, 27, e762-e764.	2.1	0
112	Adverse Events (AEs): Are They Factored into Costs of Hemophilia Care?.. Blood, 2005, 106, 5550-5550.	1.4	0
113	Can Health Care Plans Afford Hemophilia Costs? Yes.. Blood, 2005, 106, 5551-5551.	1.4	0
114	Hemophilia Treatment - A Model for Appropriate Cost Control.. Blood, 2007, 110, 3942-3942.	1.4	0
115	Do we need a better test? Can thrombin generation be useful?. Psychophysiology, 2004, 3, 75-6.	1.1	0
116	Platelet bleeding disorders and approach to their management. Psychophysiology, 2005, 4, 85-7.	1.1	0