## Hubert Schrezenmeier

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	The Complement Inhibitor Eculizumab in Paroxysmal Nocturnal Hemoglobinuria. New England Journal of Medicine, 2006, 355, 1233-1243.	27.0	1,060
2	SARS-CoV-2 variants B.1.351 and P.1 escape from neutralizing antibodies. Cell, 2021, 184, 2384-2393.e12.	28.9	848
3	Multicenter phase 3 study of the complement inhibitor eculizumab for the treatment of patients with paroxysmal nocturnal hemoglobinuria. Blood, 2008, 111, 1840-1847.	1.4	534
4	Effect of the complement inhibitor eculizumab on thromboembolism in patients with paroxysmal nocturnal hemoglobinuria. Blood, 2007, 110, 4123-4128.	1.4	481
5	Cdc42 Activity Regulates Hematopoietic Stem Cell Aging and Rejuvenation. Cell Stem Cell, 2012, 10, 520-530.	11.1	438
6	Treatment of Aplastic Anemia with Antilymphocyte Globulin and Methylprednisolone with or without Cyclosporine. New England Journal of Medicine, 1991, 324, 1297-1304.	27.0	406
7	Longâ€ŧerm safety and efficacy of sustained eculizumab treatment in patients with paroxysmal nocturnal haemoglobinuria. British Journal of Haematology, 2013, 162, 62-73.	2.5	320
8	Outcome of patients with acquired aplastic anemia given first line bone marrow transplantation or immunosuppressive treatment in the last decade: a report from the European Group for Blood and Marrow Transplantation. Haematologica, 2007, 92, 11-18.	3.5	318
9	Antithymocyte globulin with or without cyclosporin A: 11-year follow-up of a randomized trial comparing treatments of aplastic anemia. Blood, 2003, 101, 1236-1242.	1.4	298
10	A Differentiation Checkpoint Limits Hematopoietic Stem Cell Self-Renewal in Response to DNA Damage. Cell, 2012, 148, 1001-1014.	28.9	296
11	Uptake of functionalized, fluorescent-labeled polymeric particles in different cell lines and stem cells. Biomaterials, 2006, 27, 2820-2828.	11.4	279
12	Interleukin 21–Induced Granzyme B–Expressing B Cells Infiltrate Tumors and Regulate T Cells. Cancer Research, 2013, 73, 2468-2479.	0.9	277
13	Worse outcome and more chronic GVHD with peripheral blood progenitor cells than bone marrow in HLA-matched sibling donor transplants for young patients with severe acquired aplastic anemia. Blood, 2007, 110, 1397-1400.	1.4	260
14	Uptake Mechanism of Oppositely Charged Fluorescent Nanoparticles in HeLa Cells. Macromolecular Bioscience, 2008, 8, 1135-1143.	4.1	256
15	Platelet lysate from whole blood-derived pooled platelet concentrates and apheresis-derived platelet concentrates for the isolation and expansion of human bone marrow mesenchymal stromal cells: production process, content and identification of active components. Cytotherapy, 2012, 14, 540-554.	0.7	246
16	Ravulizumab (ALXN1210) vs eculizumab in adult patients with PNH naive to complement inhibitors: the 301 study. Blood, 2019, 133, 530-539.	1.4	227
17	Treatment of acquired severe aplastic anemia: Bone marrow transplantation compared with immunosuppressive therapy-the European group for blood and marrow transplantation experience. Seminars in Hematology, 2000, 37, 69-80.	3.4	223
18	Impaired humoral immunity to SARS-CoV-2 BNT162b2 vaccine in kidney transplant recipients and dialysis patients. Science Immunology, 2021, 6, eabj1031.	11.9	223

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19	Impaired humoral and cellular immunity after SARS-CoV-2 BNT162b2 (tozinameran) prime-boost vaccination in kidney transplant recipients. Journal of Clinical Investigation, 2021, 131, .	8.2	212
20	Eculizumab in Pregnant Patients with Paroxysmal Nocturnal Hemoglobinuria. New England Journal of Medicine, 2015, 373, 1032-1039.	27.0	201
21	Baseline characteristics and disease burden in patients in the International Paroxysmal Nocturnal Hemoglobinuria Registry. Haematologica, 2014, 99, 922-929.	3.5	195
22	TSG-6 Released from Intradermally Injected Mesenchymal Stem Cells Accelerates Wound Healing and Reduces Tissue Fibrosis in Murine Full-Thickness Skin Wounds. Journal of Investigative Dermatology, 2014, 134, 526-537.	0.7	195
23	A randomized controlled study in patients with newly diagnosed severe aplastic anemia receiving antithymocyte globulin (ATG), cyclosporine, with or without G-CSF: a study of the SAA Working Party of the European Group for Blood and Marrow Transplantation. Blood, 2011, 117, 4434-4441.	1.4	187
24	Clinical-Grade Mesenchymal Stromal Cells Produced Under Various Good Manufacturing Practice Processes Differ in Their Immunomodulatory Properties: Standardization of Immune Quality Controls. Stem Cells and Development, 2013, 22, 1789-1801.	2.1	186
25	Bone marrow versus peripheral blood as the stem cell source for sibling transplants in acquired aplastic anemia: survival advantage for bone marrow in all age groups. Haematologica, 2012, 97, 1142-1148.	3.5	167
26	Deficiency of Innate and Acquired Immunity Caused by an <i>IKBKB</i> Mutation. New England Journal of Medicine, 2013, 369, 2504-2514.	27.0	161
27	Treatment of acquired aplastic anemia: Bone marrow transplantation compared with immunosuppressive therapy[mdash ]The European Group for Blood and Marrow Transplantation Experience. Seminars in Hematology, 2000, 37, 69-80.	3.4	160
28	Prospective study of rabbit antithymocyte globulin and cyclosporine for aplastic anemia from the EBMT Severe Aplastic Anaemia Working Party. Blood, 2012, 119, 5391-5396.	1.4	156
29	GMP-Compliant Isolation and Large-Scale Expansion of Bone Marrow-Derived MSC. PLoS ONE, 2012, 7, e43255.	2.5	156
30	Bacterial contamination of platelet concentrates: results of a prospective multicenter study comparing pooled whole blood?derived platelets and apheresis platelets. Transfusion, 2007, 47, 644-652.	1.6	155
31	Similar outcome of upfrontâ€unrelated and matched sibling stem cell transplantation in idiopathic paediatric aplastic anaemia. A study on behalf of the <scp>UK</scp> Paediatric <scp>BMT</scp> Working Party, Paediatric Diseases Working Party and Severe Aplastic Anaemia Working Party of <scp>EBMT</scp> BMT	2.5	146
32	Telomerase gene mutations are associated with cirrhosis formation. Hepatology, 2011, 53, 1608-1617.	7.3	143
33	Impact of age on outcomes after bone marrow transplantation for acquired aplastic anemia using HLA-matched sibling donors. Haematologica, 2010, 95, 2119-2125.	3.5	137
34	Second Allograft for Hematologic Relapse of Acute Leukemia After First Allogeneic Stem-Cell Transplantation From Related and Unrelated Donors: The Role of Donor Change. Journal of Clinical Oncology, 2013, 31, 3259-3271.	1.6	137
35	Cell therapy induced regeneration of severely atrophied mandibular bone in a clinical trial. Stem Cell Research and Therapy, 2018, 9, 213.	5.5	132
36	Peptide nanofibrils boost retroviral gene transfer and provide a rapid means for concentrating viruses. Nature Nanotechnology, 2013, 8, 130-136.	31.5	125

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37	Incomplete inhibition by eculizumab: mechanistic evidence for residual C5 activity during strong complement activation. Blood, 2017, 129, 970-980.	1.4	119
38	Standardization of Good Manufacturing Practice–compliant production of bone marrow–derived human mesenchymal stromal cells for immunotherapeutic applications. Cytotherapy, 2015, 17, 128-139.	0.7	118
39	GMP-Compliant Isolation and Expansion of Bone Marrow-Derived MSCs in the Closed, Automated Device Quantum Cell Expansion System. Cell Transplantation, 2013, 22, 1981-2000.	2.5	115
40	Granulocyte-stimulating factor and severe aplastic anemia: a survey by the European Group for Blood and Marrow Transplantation (EBMT). Blood, 2007, 109, 2794-2796.	1.4	111
41	Allogeneic stem cell transplantation in paroxysmal nocturnal hemoglobinuria. Haematologica, 2012, 97, 1666-1673.	3.5	110
42	Outcome of aplastic anaemia in children. A study by the severe aplastic anaemia and paediatric disease working parties of the European group blood and bone marrow transplant. British Journal of Haematology, 2015, 169, 565-573.	2.5	104
43	Heterologous ChAdOx1 nCoV-19 and BNT162b2 prime-boost vaccination elicits potent neutralizing antibody responses and T cell reactivity against prevalent SARS-CoV-2 variants. EBioMedicine, 2022, 75, 103761.	6.1	104
44	Results of Intracoronary Stem Cell Therapy After Acute Myocardial Infarction. American Journal of Cardiology, 2010, 105, 804-812.	1.6	102
45	Pre-clinical studies of bone regeneration with human bone marrow stromal cells and biphasic calcium phosphate. Stem Cell Research and Therapy, 2014, 5, 114.	5.5	100
46	T lymphocyte activation by staphylococcal enterotoxins: Role of class II molecules and T cell surface structures. Cellular Immunology, 1989, 120, 92-101.	3.0	96
47	Eosinophils Oxidize Damage-Associated Molecular Pattern Molecules Derived from Stressed Cells. Journal of Immunology, 2009, 183, 5023-5031.	0.8	96
48	Phenotypic Characterization of Mesenchymal Stem Cells from Various Tissues. Transfusion Medicine and Hemotherapy, 2008, 35, 168-184.	1.6	94
49	Synthesis and biomedical applications of functionalized fluorescent and magnetic dual reporter nanoparticles as obtained in the miniemulsion process. Journal of Physics Condensed Matter, 2006, 18, S2581-S2594.	1.8	89
50	Feasibility and safety of treating non-unions in tibia, femur and humerus with autologous, expanded, bone marrow-derived mesenchymal stromal cells associated with biphasic calcium phosphate biomaterials in a multicentric, non-comparative trial. Biomaterials, 2019, 196, 100-108.	11.4	87
51	Human B cells differentiate into granzyme Bâ€secreting cytotoxic B lymphocytes upon incomplete Tâ€cell help. Immunology and Cell Biology, 2012, 90, 457-467.	2.3	82
52	B and T Cell Responses after a Third Dose of SARS-CoV-2 Vaccine in Kidney Transplant Recipients. Journal of the American Society of Nephrology: JASN, 2021, 32, 3027-3033.	6.1	82
53	Serial chimerism analyses indicate that mixed haemopoietic chimerism influences the probability of graft rejection and disease recurrence following allogeneic stem cell transplantation (SCT) for severe aplastic anaemia (SAA): indication for routine assessment of chimerism post SCT for SAA. British Journal of Haematology, 2009, 144, 933-945	2.5	80
54	Characterization of breakthrough hemolysis events observed in the phase 3 randomized studies of ravulizumab versus eculizumab in adults with paroxysmal nocturnal hemoglobinuria. Haematologica, 2020, 106, 230-237.	3.5	77

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55	Anticoagulant-induced Pseudothrombocytopenia and Pseudoleucocytosis. Thrombosis and Haemostasis, 1995, 73, 506-513.	3.4	75
56	Drug induced immune haemolytic anaemia in the Berlin Case ontrol Surveillance Study. British Journal of Haematology, 2011, 154, 644-653.	2.5	75
57	Outcome of Pregnancy and Disease Course among Women with Aplastic Anemia Treated with Immunosuppression. Annals of Internal Medicine, 2002, 137, 164.	3.9	74
58	Eculizumab in cold agglutinin disease (DECADE): an open-label, prospective, bicentric, nonrandomized phase 2 trial. Blood Advances, 2018, 2, 2543-2549.	5.2	74
59	Clonal analysis of human T cell activation by the Mycoplasma arthritidis mitogen (MAS). European Journal of Immunology, 1988, 18, 1733-1738.	2.9	73
60	Influence of Donor/Recipient Sex Matching on Outcome of Allogeneic Hematopoietic Stem Cell Transplantation for Aplastic Anemia. Transplantation, 2006, 82, 218-226.	1.0	73
61	Outcome of aplastic anemia in adolescence: a survey of the Severe Aplastic Anemia Working Party of the European Group for Blood and Marrow Transplantation. Haematologica, 2014, 99, 1574-1581.	3.5	73
62	The complement C5 inhibitor crovalimab in paroxysmal nocturnal hemoglobinuria. Blood, 2020, 135, 912-920.	1.4	73
63	Results of the CAPSID randomized trial for high-dose convalescent plasma in patients with severe COVID-19. Journal of Clinical Investigation, 2021, 131, .	8.2	72
64	The spectrum of PIG-A gene mutations in aplastic anemia/paroxysmal nocturnal hemoglobinuria (AA/PNH): a high incidence of multiple mutations and evidence of a mutational hot spot. Blood, 2003, 101, 2833-2841.	1.4	71
65	Ravulizumab (ALXN1210) in patients with paroxysmal nocturnal hemoglobinuria: results of 2 phase 1b/2 studies. Blood Advances, 2018, 2, 2176-2185.	5.2	65
66	Alemtuzumab is safe and effective as immunosuppressive treatment for aplastic anaemia and singleâ€lineage marrow failure: a pilot study and a survey from the EBMT WPSAA. British Journal of Haematology, 2010, 148, 791-796.	2.5	63
67	Baseline clinical characteristics and disease burden in patients with paroxysmal nocturnal hemoglobinuria (PNH): updated analysis from the International PNH Registry. Annals of Hematology, 2020, 99, 1505-1514.	1.8	63
68	Temporary antimetabolite treatment hold boosts SARS-CoV-2 vaccination–specific humoral and cellular immunity in kidney transplant recipients. JCI Insight, 2022, 7, .	5.0	62
69	Drug-induced immune thrombocytopaenia: results from the Berlin Case–Control Surveillance Study. European Journal of Clinical Pharmacology, 2012, 68, 821-832.	1.9	59
70	Cerebral Ischemic Infarction in Paroxysmal Nocturnal Hemoglobinuria. Journal of Neurology, 2005, 252, 1379-1386.	3.6	58
71	Screening of platelet concentrates for bacterial contamination: spectrum of bacteria detected, proportionof transfused units, and clinical follow-up. Annals of Hematology, 2010, 89, 83-91.	1.8	57
72	Human mesenchymal stem cells respond to native but not oxidized damage associated molecular pattern molecules from necrotic (tumor) material. European Journal of Immunology, 2011, 41, 2021-2028.	2.9	57

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73	CD4+ T Cell–Derived IL-21 and Deprivation of CD40 Signaling Favor the In Vivo Development of Granzyme B–Expressing Regulatory B Cells in HIV Patients. Journal of Immunology, 2015, 194, 3768-3777.	0.8	57
74	Caspase-8L expression protects CD34+ hematopoietic progenitor cells and leukemic cells from CD95-mediated apoptosis. Oncogene, 2005, 24, 2421-2429.	5.9	56
75	Targeted Therapy with Eculizumab for Inherited CD59 Deficiency. New England Journal of Medicine, 2014, 370, 90-92.	27.0	55
76	Complement inhibition at the level of C3 or C5: mechanistic reasons for ongoing terminal pathway activity. Blood, 2021, 137, 443-455.	1.4	55
77	B Cell Numbers Predict Humoral and Cellular Response Upon <scp>SARS</scp> – <scp>CoV</scp> â€2 Vaccination Among Patients Treated With Rituximab. Arthritis and Rheumatology, 2022, 74, 934-947.	5.6	55
78	Robust and durable serological response following pediatric SARS-CoV-2 infection. Nature Communications, 2022, 13, 128.	12.8	54
79	Effect of functionalised fluorescence-labelled nanoparticles on mesenchymal stem cell differentiation. Biomaterials, 2010, 31, 2064-2071.	11.4	51
80	Telomere elongation and clinical response to androgen treatment in a patient with aplastic anemia and a heterozygous hTERT gene mutation. Annals of Hematology, 2012, 91, 1115-1120.	1.8	51
81	Comparative Analysis of Different Platelet Lysates and Platelet Rich Preparations to Stimulate Tendon Cell Biology: An In Vitro Study. International Journal of Molecular Sciences, 2018, 19, 212.	4.1	51
82	Independent Side-by-Side Validation and Comparison of 4 Serological Platforms for SARS-CoV-2 Antibody Testing. Journal of Infectious Diseases, 2021, 223, 796-801.	4.0	51
83	mRNAâ€Mediated Gene Delivery Into Human Progenitor Cells Promotes Highly Efficient Protein Expression. Journal of Cellular and Molecular Medicine, 2007, 11, 521-530.	3.6	48
84	Drug-induced agranulocytosis in the Berlin case–control surveillance study. European Journal of Clinical Pharmacology, 2014, 70, 339-345.	1.9	46
85	Arsenic trioxide therapy in acute promyelocytic leukemia and beyond: from bench to bedside. Leukemia and Lymphoma, 2004, 45, 2387-2401.	1.3	42
86	The K+ channel openers diazoxide and NS1619 induce depolarization of mitochondria and have differential effects on cell Ca2+ in CD34+ cell line KG-1a. Experimental Hematology, 2003, 31, 815-823.	0.4	41
87	Matching for the MICA-129 polymorphism is beneficial in unrelated hematopoietic stem cell transplantation. Blood, 2016, 128, 3169-3176.	1.4	41
88	Mitogenic activity of staphylococcal protein A is due to contaminating staphylococcal enterotoxins. Journal of Immunological Methods, 1987, 105, 133-137.	1.4	39
89	Crucial Role of IL1beta and C3a in the In Vitro-Response of Multipotent Mesenchymal Stromal Cells to Inflammatory Mediators of Polytrauma. PLoS ONE, 2015, 10, e0116772.	2.5	39
90	Thrombopoietin serum levels in patients with aplastic anaemia: correlation with platelet count and persistent elevation in remission. British Journal of Haematology, 1998, 100, 571-576.	2.5	37

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91	Paroxysmal nocturnal hemoglobinuria (PNH): higher sensitivity and validity in diagnosis and serial monitoring by flow cytometric analysis of reticulocytes. Annals of Hematology, 2011, 90, 887-899.	1.8	37
92	Comparative Analysis of Novel Complement-Targeted Inhibitors, MiniFH, and the Natural Regulators Factor H and Factor H–like Protein 1 Reveal Functional Determinants of Complement Regulation. Journal of Immunology, 2016, 196, 866-876.	0.8	37
93	Fracture Healing Is Delayed in Immunodeficient NOD/scidâ€ʿi»¿IL2Rγcnull Mice. PLoS ONE, 2016, 11, e014746	52.5	37
94	The power of DNA doubleâ€strand break (DSB) repair testing to predict breast cancer susceptibility. FASEB Journal, 2012, 26, 2094-2104.	0.5	36
95	Complement and inflammasome overactivation mediates paroxysmal nocturnal hemoglobinuria with autoinflammation. Journal of Clinical Investigation, 2019, 129, 5123-5136.	8.2	36
96	Serum erythropotietin and serum transferrin receptor levels in aplastic anaemia. British Journal of Haematology, 1994, 88, 286-294.	2.5	35
97	S100A4 and Uric Acid Promote Mesenchymal Stromal Cell Induction of IL-10+/IDO+ Lymphocytes. Journal of Immunology, 2014, 192, 6102-6110.	0.8	35
98	An enzyme-based immunodetection assay to quantify SARS-CoV-2 infection. Antiviral Research, 2020, 181, 104882.	4.1	34
99	Long-term outcome of a randomized controlled study in patients with newly diagnosed severe aplastic anemia treated with antithymocyte globulin and cyclosporine, with or without granulocyte colony-stimulating factor: a Severe Aplastic Anemia Working Party Trial from the European Group of Blood and Marrow Transplantation. Haematologica. 2020. 105. 1223-1231.	3.5	34
100	The tyrosine kinase NPM-ALK, associated with anaplastic large cell lymphoma, binds the intracellular domain of the surface receptor CD30 but is not activated by CD30 stimulation. Experimental Hematology, 1999, 27, 1796-1805.	0.4	33
101	Clinical relevance of the TNF-alpha promoter/enhancer polymorphism in patients with aplastic anemia. Annals of Hematology, 2002, 81, 566-569.	1.8	33
102	Translation of a standardized manufacturing protocol for mesenchymal stromal cells: A systematic comparison of validation and manufacturing data. Cytotherapy, 2019, 21, 468-482.	0.7	33
103	Haploidentical hematopoietic stem cell transplantation in aplastic anemia: a systematic review and meta-analysis of clinical outcome on behalf of the severe aplastic anemia working party of the European group for blood and marrow transplantation (SAAWP of EBMT). Bone Marrow Transplantation, 2020, 55, 1906-1917.	2.4	33
104	Altered increase in STAT1 expression and phosphorylation in severe COVIDâ€19. European Journal of Immunology, 2022, 52, 138-148.	2.9	33
105	First line treatment of aplastic anemia with thymoglobuline in Europe and Asia: Outcome of 955 patients treated 2001â€2012. American Journal of Hematology, 2018, 93, 643-648.	4.1	32
106	Early efficacy evaluation of mesenchymal stromal cells (MSC) combined to biomaterials to treat long bone non-unions. Injury, 2020, 51, S63-S73.	1.7	32
107	A phase I/II trial of recombinant human interleukinâ€6 in patients with aplastic anaemia. British Journal of Haematology, 1995, 90, 283-292.	2.5	31
108	Experimental blunt chest trauma-induced myocardial inflammation and alteration of gap-junction protein connexin 43. PLoS ONE, 2017, 12, e0187270.	2.5	31

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109	Self versus Nonself Discrimination by the Soluble Complement Regulators Factor H and FHL-1. Journal of Immunology, 2019, 202, 2082-2094.	0.8	31
110	SARS-CoV-2 neutralising antibody testing in Europe: towards harmonisation of neutralising antibody titres for better use of convalescent plasma and comparability of trial data. Eurosurveillance, 2021, 26, .	7.0	31
111	Activation of human T lymphocytes III. Triggering of bystander cytotoxicity in cytotoxic T cell clones by antibodies against the T3 antigen or by a calcium ionophore. European Journal of Immunology, 1985, 15, 1019-1024.	2.9	30
112	Labeling of mesenchymal stromal cells with iron oxide–poly(l-lactide) nanoparticles for magnetic resonance imaging: uptake, persistence, effects on cellular function and magnetic resonance imaging properties. Cytotherapy, 2011, 13, 962-975.	0.7	30
113	Paroxysmal Nocturnal Haemoglobinuria: A Replacement of Haematopoietic Tissue?. Acta Haematologica, 2000, 103, 41-48.	1.4	29
114	Selectivity of C3-opsonin targeted complement inhibitors: A distinct advantage in the protection of erythrocytes from paroxysmal nocturnal hemoglobinuria patients. Immunobiology, 2016, 221, 503-511.	1.9	28
115	HLA Matching in Unrelated Stem Cell Transplantation up to Date. Transfusion Medicine and Hemotherapy, 2019, 46, 326-336.	1.6	28
116	Impact of Donor Activating KIR Genes on HSCT Outcome in C1-Ligand Negative Myeloid Disease Patients Transplanted with Unrelated Donors—A Retrospective Study. PLoS ONE, 2017, 12, e0169512.	2.5	28
117	Acute myeloid leukemia with mutated nucleophosmin 1: an immunogenic acute myeloid leukemia subtype and potential candidate for immune checkpoint inhibition. Haematologica, 2017, 102, e499-e501.	3.5	26
118	The changing scene of allogeneic stem cell transplantation for chronic myeloid leukemia—a report from the German Registry covering the period from 1998 to 2004. Annals of Hematology, 2009, 88, 1237-1247.	1.8	25
119	Development of a disease-specific quality of life questionnaire for patients with aplastic anemia and/or paroxysmal nocturnal hemoglobinuria (QLQ-AA/PNH)—report on phases I and II. Annals of Hematology, 2017, 96, 171-181.	1.8	25
120	The Terminal Complement Inhibitor Eculizumab Reduces Thrombosis in Patients with Paroxysmal Nocturnal Hemoglobinuria Blood, 2006, 108, 123-123.	1.4	25
121	Healthy donor hematopoietic stem cell mobilization with biosimilar granulocyte olonyâ€stimulating factor: safety, efficacy, and graft performance. Transfusion, 2016, 56, 3055-3064.	1.6	24
122	One-year efficacy and safety of ravulizumab in adults with paroxysmal nocturnal hemoglobinuria naìve to complement inhibitor therapy: open-label extension of a randomized study. Therapeutic Advances in Hematology, 2020, 11, 204062072096613.	2.5	24
123	Depolarisation of the plasma membrane in the arsenic trioxide (As2O3)-and anti-CD95-induced apoptosis in myeloid cells. FEBS Letters, 2004, 578, 85-89.	2.8	23
124	Should irradiated blood products be given routinely to all patients with aplastic anaemia undergoing immunosuppressive therapy with antithymocyte globulin (ATG)? A survey from the European Group for Blood and Marrow Transplantation Severe Aplastic Anaemia Working Party. British Journal of Haematology, 2010, 150, 377-379.	2.5	23
125	Drugs that inhibit complement. Transfusion and Apheresis Science, 2012, 46, 87-92.	1.0	23
126	Characterization of the SARS-CoV-2 Neutralization Potential of COVID-19–Convalescent Donors. Journal of Immunology, 2021, 206, 2614-2622.	0.8	22

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127	Evaluation of platelet-rich plasma and hydrostatic pressure regarding cell differentiation in nucleus pulposus tissue engineering. Journal of Tissue Engineering and Regenerative Medicine, 2013, 7, 244-252.	2.7	21
128	Natural Killer Cells Generated From Human Induced Pluripotent Stem Cells Mature to CD56brightCD16+NKp80+/-In-Vitro and Express KIR2DL2/DL3 and KIR3DL1. Frontiers in Immunology, 2021, 12, 640672.	4.8	21
129	Donors for SARS-CoV-2 Convalescent Plasma for a Controlled Clinical Trial: Donor Characteristics, Content and Time Course of SARS-CoV-2 Neutralizing Antibodies. Transfusion Medicine and Hemotherapy, 2021, 48, 137-147.	1.6	21
130	Endocrine Effects of Recombinant Interleukin 6 in Man. Neuroendocrinology, 1996, 63, 237-243.	2.5	20
131	Antimony-trioxide- and arsenic-trioxide-induced apoptosis in myelogenic and lymphatic cell lines, recruitment of caspases, and loss of mitochondrial membrane potential are enhanced by modulators of the cellular glutathione redox system. Annals of Hematology, 2009, 88, 1047-1058.	1.8	20
132	Leukemic progenitor cells are susceptible to targeting by stimulated cytotoxic <scp>T</scp> cells against immunogenic leukemiaâ€associated antigens. International Journal of Cancer, 2015, 137, 2083-2092.	5.1	19
133	Clinical benefit of eculizumab in patients with no transfusion history in the International Paroxysmal Nocturnal Haemoglobinuria Registry. Internal Medicine Journal, 2017, 47, 1026-1034.	0.8	19
134	Osteonecrosis of the Femoral Head Safely Healed with Autologous, Expanded, Bone Marrow-Derived Mesenchymal Stromal Cells in a Multicentric Trial with Minimum 5 Years Follow-Up. Journal of Clinical Medicine, 2021, 10, 508.	2.4	19
135	Design and development of a disease-specific quality of life tool for patients with aplastic anaemia and/or paroxysmal nocturnal haemoglobinuria (QLQ-AA/PNH)—a report on phase III. Annals of Hematology, 2019, 98, 1547-1559.	1.8	18
136	Complement Activation and Organ Damage After Trauma—Differential Immune Response Based on Surgical Treatment Strategy. Frontiers in Immunology, 2020, 11, 64.	4.8	18
137	Tissue distribution of blood group membrane proteins beyond red cells: Evidence from cDNA libraries. Transfusion and Apheresis Science, 2006, 35, 71-82.	1.0	17
138	Direct and indirect effects of functionalised fluorescence-labelled nanoparticles on human osteoclast formation and activity. Biomaterials, 2011, 32, 1706-1714.	11.4	17
139	Human leukocyte antigen-E mismatch is associated with better hematopoietic stem cell transplantation outcome in acute leukemia patients. Haematologica, 2017, 102, 1947-1955.	3.5	17
140	Different Levels of Incomplete Terminal Pathway Inhibition by Eculizumab and the Clinical Response of PNH Patients. Frontiers in Immunology, 2019, 10, 1639.	4.8	17
141	Effectiveness of eculizumab in patients with paroxysmal nocturnal hemoglobinuria (PNH) with or without aplastic anemia in the International PNH Registry. American Journal of Hematology, 2019, 94, E37-E41.	4.1	17
142	Next-Generation Sequencing Technologies in Blood Group Typing. Transfusion Medicine and Hemotherapy, 2020, 47, 4-13.	1.6	17
143	Evaluation of SARSâ€CoVâ€2 antibody titers and potency for convalescent plasma donation: a brief commentary. Vox Sanguinis, 2021, 116, 493-496.	1.5	17
144	Complement Inhibition with Eculizumab in Patients with Cold Agglutinin Disease (CAD): Results from a Prospective Phase II Trial (DECADE Trial). Blood, 2015, 126, 274-274.	1.4	17

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145	Eculizumab opens a new era of treatment for paroxysmal nocturnal hemoglobinuria. Expert Review of Hematology, 2009, 2, 7-16.	2.2	16
146	Deregulated expression of circular RNAs in acute myeloid leukemia. Blood Advances, 2021, 5, 1490-1503.	5.2	16
147	Significant Disease Burden in Paroxysmal Nocturnal Hemoglobinuria Patients with Lower Levels of Hemolysis, Mild Anemia and Minimal Transfusion: Clinical Improvement with Eculizumab Therapy Blood, 2007, 110, 840-840.	1.4	16
148	Serum Erythropoietin and Thrombopoietin Levels in Patients with Essential Thrombocythaemia. Leukemia and Lymphoma, 2000, 36, 533-538.	1.3	15
149	Effects of a Ceramic Biomaterial on Immune Modulatory Properties and Differentiation Potential of Human Mesenchymal Stromal Cells of Different Origin. Tissue Engineering - Part A, 2015, 21, 767-781.	3.1	15
150	Different clinical characteristics of paroxysmal nocturnal hemoglobinuria in pediatric and adult patients. Haematologica, 2017, 102, e76-e79.	3.5	15
151	Autologous Mesenchymal Stroma Cells Are Superior to Allogeneic Ones in Bone Defect Regeneration. International Journal of Molecular Sciences, 2018, 19, 2526.	4.1	15
152	Inflammatory response of mesenchymal stromal cells after in vivo exposure with selected trauma-related factors and polytrauma serum. PLoS ONE, 2019, 14, e0216862.	2.5	15
153	Complement and the prothrombotic state. Blood, 2022, 139, 1954-1972.	1.4	15
154	DNA methylation in <i>PRDM8</i> is indicative for dyskeratosis congenita. Oncotarget, 2016, 7, 10765-10772.	1.8	15
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