Alok Srivastava

List of Publications by Year
in descending order

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Version: 2024-02-01

322 papers	9,983 citations	70961 41 h-index	42291 92 g-index
332 all docs	332 docs citations	332 times ranked	7577 citing authors

#	Article	IF	Citations
1	Guidelines for the management of hemophilia. Haemophilia, 2013, 19, e1-47.	1.0	1,538
2	Update on the pathophysiology and classification of von Willebrand disease: a report of the Subcommittee on von Willebrand Factor. Journal of Thrombosis and Haemostasis, 2006, 4, 2103-2114.	1.9	1,061
3	WFH Guidelines for the Management of Hemophilia, 3rd edition. Haemophilia, 2020, 26, 1-158.	1.0	915
4	Definitions in hemophilia: communication from the SSC of the ISTH. Journal of Thrombosis and Haemostasis, 2014, 12, 1935-1939.	1.9	530
5	Impact, Diagnosis and Treatment of von Willebrand Disease. Thrombosis and Haemostasis, 2000, 84, 160-174.	1.8	478
6	The discriminant power of bleeding history for the diagnosis of type 1 von Willebrand disease: an international, multicenter study. Journal of Thrombosis and Haemostasis, 2005, 3, 2619-2626.	1.9	317
7	New ISSCR Guidelines Underscore Major Principles for Responsible Translational Stem Cell Research. Cell Stem Cell, 2008, 3, 607-609.	5.2	218
8	Glutathione S-transferase M1 polymorphism: a risk factor for hepatic venoocclusive disease in bone marrow transplantation. Blood, 2004, 104, 1574-1577.	0.6	155
9	Quality control guidelines for clinical-grade human induced pluripotent stem cell lines. Regenerative Medicine, 2018, 13, 859-866.	0.8	147
10	Genetic diagnosis of haemophilia and other inherited bleeding disorders. Haemophilia, 2006, 12, 82-89.	1.0	123
11	Diagnostic Accuracy of Ultrasound for Assessment of Hemophilic Arthropathy: MRI Correlation. American Journal of Roentgenology, 2015, 204, W336-W347.	1.0	98
12	Acute myeloid leukaemia: challenges and real world data from India. British Journal of Haematology, 2015, 170, 110-117.	1.2	96
13	Molecular defects in type 3 von Willebrand disease: updated results from 40 multiethnic patients. Blood Cells, Molecules, and Diseases, 2003, 30, 264-270.	0.6	95
14	Functional Independence Score in Haemophilia: a new performance-based instrument to measure disability. Haemophilia, 2005, 11, 598-602.	1.0	92
15	One and a half million hematopoietic stem cell transplants: continuous and differential improvement in worldwide access with the use of non-identical family donors. Haematologica, 2022, 107, 1045-1053.	1.7	87
16	SARS-2 Coronavirusâ€“Associated Hemostatic Lung Abnormality in COVID-19: Is It Pulmonary Thrombosis or Pulmonary Embolism?. Seminars in Thrombosis and Hemostasis, 2020, 46, 777-780.	1.5	85
17	Comprehensive care for haemophilia around the world. Haemophilia, 2004, 10, 9-13.	1.0	83
18	Improved Clinical Outcomes of High Risk Î² Thalassemia Major Patients Undergoing a HLA Matched Related Allogeneic Stem Cell Transplant with a Treosulfan Based Conditioning Regimen and Peripheral Blood Stem Cell Grafts. PLoS ONE, 2013, 8, e61637.	1.1	78

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19	A study of reported factor VIII use around the world. Haemophilia, 2010, 16, 33-46.	1.0	75
20	Tests of global haemostasis and their applications in bleeding disorders. Haemophilia, 2010, 16, 85-92.	1.0	75
21	Identification of factor VIII gene mutations in 101 patients with haemophilia A: mutation analysis by inversion screening and multiplex PCR and CSGE and molecular modelling of 10 novel missense substitutions. Haemophilia, 2005, 11, 481-491.	1.0	69
22	A New Stratification Strategy That Identifies a Subset of Class III Patients with an Adverse Prognosis among Children with Î² Thalassemia Major Undergoing a Matched Related Allogeneic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2007, 13, 889-894.	2.0	67
23	Psychometric analysis of the Functional Independence Score in Haemophilia (FISH). Haemophilia, 2007, 13, 620-626.	1.0	65
24	Choosing outcome assessment tools in haemophilia care and research: a multidisciplinary perspective. Haemophilia, 2017, 23, 11-24.	1.0	63
25	Molecular Characterization of a Multiethnic Group of 21 Patients with Type 3 von Willebrand Disease. Thrombosis and Haemostasis, 2000, 84, 536-540.	1.8	61
26	The Phenotypic Heterogeneity of Severe Hemophilia. Seminars in Thrombosis and Hemostasis, 2008, 34, 128-141.	1.5	61
27	Nuclear factor (NF)â€ÎºB and its associated pathways are major molecular regulators of bloodâ€induced joint damage in a murine model of hemophilia. Journal of Thrombosis and Haemostasis, 2013, 11, 293-306.	1.9	61
28	Randomized trial of two different conditioning regimens for bone marrow transplantation in thalassemia â€“ the role of busulfan pharmacokinetics in determining outcome. Bone Marrow Transplantation, 2005, 36, 839-845.	1.3	60
29	Cure for thalassemia major â€“ from allogeneic hematopoietic stem cell transplantation to gene therapy. Haematologica, 2017, 102, 214-223.	1.7	57
30	Molecular mechanisms underlying hemophilia A phenotype in seven females. Journal of Thrombosis and Haemostasis, 2009, 7, 976-982.	1.9	56
31	Long-term outcome following splenectomy for chronic and persistent immune thrombocytopenia (ITP) in adults and children. Annals of Hematology, 2016, 95, 1429-1434.	0.8	56
32	When and how to start prophylaxis in boys with severe hemophilia without inhibitors: communication from the SSC of the ISTH. Journal of Thrombosis and Haemostasis, 2016, 14, 1105-1109.	1.9	54
33	Increased ABCG2 Expression Could Be Responsible for Resistance to Imatinib Mesylate in Patients with Chronic Myeloid Leukemia Who Do Not Have Mutations in BCR-ABL Kinase Domain. Blood, 2008, 112, 5026-5026.	0.6	53
34	Laboratory issues in bleeding disorders. Haemophilia, 2006, 12, 68-75.	1.0	51
35	Real-World Issues and Potential Solutions in Hematopoietic Cell Transplantation during the COVID-19 Pandemic: Perspectives from the Worldwide Network for Blood and Marrow Transplantation and Center for International Blood and Marrow Transplant Research Health Services and International Studies Committee. Biology of Blood and Marrow Transplantation, 2020, 26, 2181-2189.	2.0	51
36	Long-Term Cultured Human Term Placenta-Derived Mesenchymal Stem Cells of Maternal Origin Displays Plasticity. Stem Cells International, 2012, 2012, 1-11.	1.2	50

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37	Management of haemophilia in the developing world. Haemophilia, 1998, 4, 474-480.	1.0	48
38	Dose and response in haemophilia - optimization of factor replacement therapy. British Journal of Haematology, 2004, 127, 12-25.	1.2	46
39	Positioning a Scientific Community on Unproven Cellular Therapies: The 2015 International Society for Cellular Therapy Perspective. Cytotherapy, 2015, 17, 1663-1666.	0.3	44
40	Correlating clinical and radiological assessment of joints in haemophilia: results of a cross sectional study. Haemophilia, 2016, 22, 925-933.	1.0	44
41	Fundamentals for a Systematic Approach to Mild and Moderate Inherited Bleeding Disorders: An EHA Consensus Report. HemaSphere, 2019, 3, e286.	1.2	43
42	Comparison of Newly Diagnosed and Relapsed Patients with Acute Promyelocytic Leukemia Treated with Arsenic Trioxide: Insight into Mechanisms of Resistance. PLoS ONE, 2015, 10, e0121912.	1.1	43
43	RNA expression of genes involved in cytarabine metabolism and transport predicts cytarabine response in acute myeloid leukemia. Pharmacogenomics, 2015, 16, 877-890.	0.6	41
44	Preclinical Development of a Hematopoietic Stem and Progenitor Cell Bioengineered Factor VIII Lentiviral Vector Gene Therapy for Hemophilia A. Human Gene Therapy, 2018, 29, 1183-1201.	1.4	39
45	Epidemiology of von Willebrand Disease in Developing Countries. Seminars in Thrombosis and Hemostasis, 2005, 31, 569-576.	1.5	38
46	Blood-induced bone loss in murine hemophilic arthropathy is prevented by blocking the iRhom2/ADAM17/TNF-Î± pathway. Blood, 2018, 132, 1064-1074.	0.6	38
47	Improved joint health in subjects with severe haemophilia A treated prophylactically with recombinant factor <scp>VIII</scp> Fc fusion protein. Haemophilia, 2018, 24, 77-84.	1.0	37
48	Informativeness of linkage analysis for genetic diagnosis of haemophilia A in India. Haemophilia, 2004, 10, 553-559.	1.0	36
49	Delivery of haemophilia care in the developing world. Haemophilia, 1998, 4, 33-40.	1.0	34
50	Neighborhood poverty and pediatric allogeneic hematopoietic cell transplantation outcomes: a CIBMTR analysis. Blood, 2021, 137, 556-568.	0.6	34
51	Low-dose intermittent factor replacement for post-operative haemostasis in haemophilia. Haemophilia, 1998, 4, 799-801.	1.0	33
52	High-performance liquid chromatographic method for quantification of busulfan in plasma after derivatization by tetrafluorothiophenol. Biomedical Applications, 1999, 721, 147-152.	1.7	33
53	Part 1: Defining unproven cellular therapies. Cytotherapy, 2016, 18, 117-119.	0.3	33
54	Pharmacokinetics, clot strength and safety of a new fibrinogen concentrate: randomized comparison with active control in congenital fibrinogen deficiency. Journal of Thrombosis and Haemostasis, 2018, 16, 253-261.	1.9	33

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55	Factor replacement therapy in haemophilia - are there models for developing countries?. Haemophilia, 2003, 9, 391-396.	1.0	32
56	Blood-Induced Arthropathy in Hemophilia: Mechanisms and Heterogeneity. Seminars in Thrombosis and Hemostasis, 2015, 41, 832-837.	1.5	31
57	Measurement of joint health in persons with haemophilia: A systematic review of the measurement properties of haemophiliaâ€specific instruments. Haemophilia, 2019, 25, e1-e10.	1.0	31
58	Pharmacokinetics of oral busulphan in children with beta thalassaemia major undergoing allogeneic bone marrow transplantation. Bone Marrow Transplantation, 1999, 24, 5-11.	1.3	30
59	Definitions in Hemophilia: Resolved and Unresolved Issues. Seminars in Thrombosis and Hemostasis, 2015, 41, 819-825.	1.5	30
60	Frequency of rare <scp>BCR</scp>â€<scp>ABL</scp>1 fusion transcripts in chronic myeloid leukemia patients. International Journal of Laboratory Hematology, 2017, 39, 235-242.	0.7	30
61	Hemophilia Treatment in Developing Countries: Products and Protocols. Seminars in Thrombosis and Hemostasis, 2005, 31, 495-500.	1.5	29
62	Efficacy and safety of a <scp>VWF</scp>/<scp>FVIII</scp> concentrate (wilate^{Â®}) in inherited von Willebrand disease patients undergoing surgical procedures. Haemophilia, 2017, 23, 264-272.	1.0	29
63	Identification of novel HPFH-like mutations by CRISPR base editing that elevate the expression of fetal hemoglobin. ELife, 2022, 11, .	2.8	29
64	Role of Molecular Genetics in Hemophilia: From Diagnosis to Therapy. Seminars in Thrombosis and Hemostasis, 2012, 38, 64-78.	1.5	28
65	Rationale and efficacy of proteasome inhibitor combined with arsenic trioxide in the treatment of acute promyelocytic leukemia. Leukemia, 2016, 30, 2169-2178.	3.3	28
66	Developing an algorithm of informative markers for evaluation of chimerism after allogeneic bone marrow transplantation. Bone Marrow Transplantation, 2006, 37, 751-755.	1.3	27
67	Clinical Profile and Outcomes of Patients with Î² Thalassemia Major and Hepatitis C Virus Infection Undergoing an Allogeneic Stem Cell Transplant. Blood, 2012, 120, 4160-4160.	0.6	27
68	Invasive fungal infection following chemotherapy for acute myeloid leukaemiaâ€"Experience from a developing country. Mycoses, 2017, 60, 686-691.	1.8	26
69	Surgery for Hemophilia in Developing Countries. Seminars in Thrombosis and Hemostasis, 2005, 31, 538-543.	1.5	25
70	Inflammation is key to hemophilic arthropathy. Blood, 2015, 126, 2175-2176.	0.6	25
71	Assessments of outcome in haemophilia â€“ what is the added value of <scp>QoL</scp> tools?. Haemophilia, 2015, 21, 430-435.	1.0	25
72	Long-term outcome of mixed chimerism after stem cell transplantation for thalassemia major conditioned with busulfan and cyclophosphamide. Bone Marrow Transplantation, 2018, 53, 169-174.	1.3	25

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73	Factor concentrates for haemophilia in the developing world. Haemophilia, 1998, 4, 481-485.	1.0	24
74	ATP-binding casette transporter expression in acute myeloid leukemia: association within vitrocytotoxicity and prognostic markers. Pharmacogenomics, 2017, 18, 235-244.	0.6	24
75	Emerging therapies for haemophilia â€•Global perspective. Haemophilia, 2018, 24, 15-21.	1.0	24
76	Factor replacement for haemophilia - should cryoprecipitate be used?. Haemophilia, 1999, 5, 301-305.	1.0	23
77	A study of reported factor IX use around the world. Haemophilia, 2011, 17, 446-455.	1.0	23
78	Sensitivity and Robustness of Spatially Dependent Thrombin Generation and Fibrin Clot Propagation. Biophysical Journal, 2018, 115, 2461-2473.	0.2	23
79	Allogeneic Stem Cell Transplantation for Thalassemia Major. Hematology/Oncology Clinics of North America, 2014, 28, 1187-1200.	0.9	22
80	Systematic evaluation of markers used for the identification of human induced pluripotent stem cells. Biology Open, 2017, 6, 100-108.	0.6	22
81	Pharmacokinetics and Pharmacodynamics of Treosulfan in Patients With Thalassemia Major Undergoing Allogeneic Hematopoietic Stem Cell Transplantation. Clinical Pharmacology and Therapeutics, 2018, 104, 575-583.	2.3	22
82	Post-Transplant Cyclophosphamide as Sole Graft-versus-Host Disease Prophylaxis Is Feasible in Patients Undergoing Peripheral Blood Stem Cell Transplantation for Severe Aplastic Anemia Using Matched Sibling Donors. Biology of Blood and Marrow Transplantation, 2018, 24, 494-500.	2.0	22
83	Worldwide Network for Blood and Marrow Transplantation Recommendations for Establishing a Hematopoietic Stem Cell Transplantation Program in Countries with Limited Resources, Part II: Clinical, Technical, and Socioeconomic Considerations. Biology of Blood and Marrow Transplantation, 2019, 25, 2330-2337.	2.0	22
84	Choice of factor concentrates for haemophilia: a developing world perspective. Haemophilia, 2001, 7, 117-122.	1.0	22
85	Measuring activities and participation in persons with haemophilia: A systematic review of commonly used instruments. Haemophilia, 2018, 24, e33-e49.	1.0	21
86	Worldwide Network for Blood and Marrow Transplantation Recommendations for Establishing a Hematopoietic Cell Transplantation Program, Part I: Minimum Requirements and Beyond. Biology of Blood and Marrow Transplantation, 2019, 25, 2322-2329.	2.0	21
87	Arsenic Trioxide (As2O3) in the Treatment of Patients with Newly Diagnosed Acute Promyelocytic Leukemia (APML) - Toxicity and Outcome.. Blood, 2004, 104, 889-889.	0.6	21
88	Patient access to and ethical considerations of the application of the European Union hospital exemption rule for advanced therapy medicinal products. Cytotherapy, 2022, 24, 686-690.	0.3	21
89	Polymyositis - an unusual manifestation of chronic graft-versus-host disease. Rheumatology International, 2001, 20, 169-170.	1.5	20
90	Management of Hemophilia in Patients with Inhibitors: The Perspective from Developing Countries. Seminars in Thrombosis and Hemostasis, 2009, 35, 820-826.	1.5	20

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91	Population pharmacokinetics of Daunorubicin in adult patients with acute myeloid leukemia. Cancer Chemotherapy and Pharmacology, 2016, 78, 1051-1058.	1.1	20
92	Patientâ€relevant health outcomes for hemophilia care: Development of an international standard outcomes set. Research and Practice in Thrombosis and Haemostasis, 2021, 5, e12488.	1.0	20
93	Episodic replacement of clotting factor concentrates does not prevent bleeding or musculoskeletal damage â€“ the <scp>MUSFIH</scp> study. Haemophilia, 2017, 23, 538-546.	1.0	20
94	Dose and outcome of care in haemophilia - how do we define cost-effectiveness?. Haemophilia, 2004, 10, 216-220.	1.0	19
95	Cellular Immune Reconstitution and Its Impact on Clinical Outcome in Children with Î² Thalassemia Major Undergoing a Matched Related Myeloablative Allogeneic Bone Marrow Transplant. Biology of Blood and Marrow Transplantation, 2009, 15, 597-609.	2.0	19
96	Factor VIII/factor IX prophylaxis for severe hemophilia. Seminars in Hematology, 2016, 53, 3-9.	1.8	19
97	Randomized, Double-Blind, Pharmacokinetic Equivalence Trial Comparing DRL-Rituximab With MabThera in Patients With Diffuse Large B-Cell Lymphoma. Journal of Global Oncology, 2019, 5, 1-13.	0.5	19
98	Development of a Clinical Candidate AAV3 Vector for Gene Therapy of Hemophilia B. Human Gene Therapy, 2020, 31, 1114-1123.	1.4	19
99	Fracture neck of femur in haemophilia A ? experience from a cohort of 11 patients from a tertiary centre in India. Haemophilia, 2007, 13, 391-394.	1.0	18
100	Impact of pretransplant splenectomy on patients with Î²â€thalassemia major undergoing a matchedâ€related allogeneic stem cell transplantation. Pediatric Transplantation, 2009, 13, 171-176.	0.5	18
101	Differences between developed and developing countries in paediatric care in haemophilia. Haemophilia, 2012, 18, 94-100.	1.0	18
102	Haemophilia care â€“ beyond the treatment guidelines. Haemophilia, 2014, 20, 4-10.	1.0	18
103	Use of Non-Cryopreserved Peripheral Blood Stem Cells Is Associated with Adequate Engraftment in Patients with Multiple Myeloma Undergoing an Autologous Transplant. Biology of Blood and Marrow Transplantation, 2018, 24, e31-e35.	2.0	18
104	External fixators in haemophilia. Haemophilia, 2004, 10, 52-57.	1.0	17
105	External Quality Assessment Scheme for Hemostasis in India. Seminars in Thrombosis and Hemostasis, 2007, 33, 265-272.	1.5	17
106	Worldwide Network for Blood and Marrow Transplantation (WBMT) recommendations for establishing a hematopoietic stem cell transplantation program in countries with limited resources (Part II): Clinical, technical and socio-economic considerations. Hematology/ Oncology and Stem Cell Therapy, 2020, 13, 7-16.	0.6	17
107	von Willebrand disease in the developing world. Seminars in Hematology, 2005, 42, 36-41.	1.8	16
108	Achieving and maintaining quality in the laboratory. Haemophilia, 2006, 12, 61-67.	1.0	16

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109	Population pharmacokinetics of cyclophosphamide in patients with thalassemia major undergoing HSCT. Bone Marrow Transplantation, 2012, 47, 1178-1185.	1.3	15
110	Advancing Stem Cell Biology toward Stem Cell Therapeutics. Cell Stem Cell, 2012, 10, 149-150.	5.2	15
111	Principles of haemophilia care: The Asiaâ€Pacific perspective. Haemophilia, 2018, 24, 366-375.	1.0	15
112	Towards a global multidisciplinary consensus framework on haemophilia gene therapy: Report of the 2nd World Federation of Haemophilia Gene Therapy Round Table. Haemophilia, 2020, 26, 443-449.	1.0	15
113	Association betweenCYP1A2gene single nucleotide polymorphisms and clinical responses to clozapine in patients with treatment-resistant schizophrenia. Acta Neuropsychiatrica, 2013, 25, 2-11.	1.0	14
114	Arsenic Trioxide Enhances the NK Cell Cytotoxicity Against Acute Promyelocytic Leukemia While Simultaneously Inhibiting Its Bio-Genesis. Frontiers in Immunology, 2018, 9, 1357.	2.2	14
115	Utility of tissue elafin as an immunohistochemical marker for diagnosis of acute skin graft-versus-host disease: a pilot study. Clinical and Experimental Dermatology, 2019, 44, 161-168.	0.6	14
116	Worldwide Network for Blood and Marrow Transplantation (WBMT) recommendations for establishing a hematopoietic cell transplantation program (Part I): Minimum requirements and beyond. Hematology/ Oncology and Stem Cell Therapy, 2020, 13, 131-142.	0.6	14
117	Comparison of Human Platelet Lysate versus Fetal Bovine Serum for Expansion of Human Articular Cartilageâ€"Derived Chondroprogenitors. Cartilage, 2021, 13, 107S-116S.	1.4	14
118	Hemophilia gene therapy knowledge and perceptions: Results of an international survey. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 644-651.	1.0	14
119	Management of COVIDâ€19â€associated coagulopathy in persons with haemophilia. Haemophilia, 2021, 27, 41-48.	1.0	14
120	Endothelial cells do not express GSTA1: potential relevance to busulfanâ€mediated endothelial damage during haematopoietic stem cell transplantation. European Journal of Haematology, 2008, 80, 299-302.	1.1	13
121	Diagnosis and Management of<scp>v</scp>on Willebrand Disease: A Developing Country Perspective. Seminars in Thrombosis and Hemostasis, 2011, 37, 587-594.	1.5	13
122	World bleeding disorders registry: The pilot study. Haemophilia, 2018, 24, e113-e116.	1.0	13
123	Prevalence of Adeno-Associated Virus 3 Capsid Binding and Neutralizing Antibodies in Healthy and Hemophilia B Individuals from India. Human Gene Therapy, 2021, 32, 451-457.	1.4	12
124	NK Cell Mediated Cytotoxicity Against Malignant Promyelocytes Enhanced By Arsenic Trioxide: Potential Clinical Relevance. Blood, 2013, 122, 1455-1455.	0.6	12
125	Efficacy and Safety of Fitusiran Prophylaxis, an siRNA Therapeutic, in a Multicenter Phase 3 Study (ATLAS-INH) in People with Hemophilia A or B, with Inhibitors (PwHI). Blood, 2021, 138, 4-4.	0.6	12
126	Fitusiran, an Investigational siRNA Therapeutic Targeting Antithrombin for the Treatment of Hemophilia: First Results from a Phase 3 Study to Evaluate Efficacy and Safety in People with Hemophilia a or B without Inhibitors (ATLAS-A/B). Blood, 2021, 138, LBA-3-LBA-3.	0.6	12

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127	Low-dose activated factor IX complex concentrates (FEIBAR) for post-operative haemostasis in a patient with high responding factor VIII inhibitors. Haemophilia, 1995, 1, 274-276.	1.0	11
128	Drugâ€resistant organisms are common in fecal surveillance cultures, predict bacteremia and correlate with poorer outcomes in patients undergoing allogeneic stem cell transplants. Transplant Infectious Disease, 2020, 22, e13273.	0.7	11
129	Risk Stratification without a Liver Biopsy of Patients with Î² Thalassemia Major Undergoing a Matched Related Allogeneic Bone Marrow Transplant.. Blood, 2009, 114, 659-659.	0.6	11
130	A patient-prioritized ability assessment in haemophilia: the Canadian Occupational Performance Measure. Haemophilia, 2011, 17, 605-611.	1.0	10
131	Clinical Outcomes in Multiple Myeloma Post-Autologous Transplantationâ€"A Single Centre Experience. Indian Journal of Hematology and Blood Transfusion, 2019, 35, 215-222.	0.3	10
132	A new and simplified comprehensive ultrasound protocol of haemophilic joints: the Universal Simplified Ultrasound (US-US) protocol. Clinical Radiology, 2019, 74, 897.e9-897.e16.	0.5	10
133	Prognostic plasma biomarkers of early complications and graftâ€versusâ€host disease in patients undergoing allogeneic hematopoietic stem cell transplantation. EJHaem, 2020, 1, 219-229.	0.4	10
134	Clinical outcomes in hemophilia: Towards development of a core set of standardized outcome measures for research. Research and Practice in Thrombosis and Haemostasis, 2020, 4, 652-658.	1.0	10
135	Diagnosis of haemophilia and other inherited bleeding disorders â€Is a new paradigm needed?. Haemophilia, 2021, 27, 14-20.	1.0	10
136	Direct Generation of Immortalized Erythroid Progenitor Cell Lines from Peripheral Blood Mononuclear Cells. Cells, 2021, 10, 523.	1.8	10
137	Surgery in patients with congenital coagulation disorders. The National Medical Journal of India, 1994, 7, 8-12.	0.1	10
138	Generation of an induced pluripotent stem cell line that mimics the disease phenotypes from a patient with Fanconi anemia by conditional complementation. Stem Cell Research, 2017, 20, 54-57.	0.3	9
139	Second Hematopoietic Stem Cell Transplant for Thalassemia Major: Improved Clinical Outcomes with a Treosulfan-Based Conditioning Regimen. Biology of Blood and Marrow Transplantation, 2018, 24, 103-108.	2.0	9
140	Prevalence of FVIII inhibitors in severe haemophilia A patients: Effect of treatment and genetic factors in an Indian population. Haemophilia, 2019, 25, 67-74.	1.0	9
141	Plasma imatinib levels and ABCB1 polymorphism influences early molecular response and failure-free survival in newly diagnosed chronic phase CML patients. Scientific Reports, 2020, 10, 20640.	1.6	9
142	Vaccination against COVIDâ€19: Rationale, modalities and precautions for patients with haemophilia and other inherited bleeding disorders. Haemophilia, 2021, 27, 515-518.	1.0	9
143	The t(8;14)(q24.1;q32) and its variant translocations: A study of 34 cases. Hematology/ Oncology and Stem Cell Therapy, 2017, 10, 126-134.	0.6	8
144	Typeâ€3 von Willebrand disease in Indiaâ€"Clinical spectrum and molecular profile. Haemophilia, 2018, 24, 930-940.	1.0	8

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145	Dendritic Cell Type 2 Counts on Day 28 in HLA-Matched Related Allogeneic PBSCT Predicts the Incidence of Acute and Chronic GVHD.. Blood, 2006, 108, 2893-2893.	0.6	8
146	Clinical, Cellular and Molecular Differences Between Newly Diagnosed and Relapsed Patients with Acute Promyelocytic Leukemia: Insights Into Mechanisms of Resistance. Blood, 2012, 120, 1390-1390.	0.6	8
147	Haemophilia gene therapyâ€"Update on new country initiatives. Haemophilia, 2022, 28, 61-67.	1.0	8
148	A new multiplex PCR and conformation-sensitive gel electrophoresis strategy for mutation detection in the platelet glycoprotein Î±IIb and Î²3 genes. Journal of Thrombosis and Haemostasis, 2007, 5, 206-209.	1.9	7
149	Standardizing patient outcomes measurement to improve haemophilia care. Haemophilia, 2016, 22, 651-653.	1.0	7
150	Coexistence of aberrant hematopoietic and stromal elements in myelodysplastic syndromes. Blood Cells, Molecules, and Diseases, 2017, 66, 37-46.	0.6	7
151	Allogeneic stem cell transplantation for thalassemia major in India. Pediatric Hematology Oncology Journal, 2017, 2, 114-120.	0.1	7
152	Antibodies to human platelet antigens form a significant proportion of platelet antibodies detected in Indian patients with refractoriness to platelet transfusions. Transfusion Medicine, 2018, 28, 392-397.	0.5	7
153	Comparison of the Efficacy of Innovator Rituximab and its Biosimilars in Diffuse Large B Cell Lymphoma Patients: A Retrospective Analysis. Indian Journal of Hematology and Blood Transfusion, 2020, 36, 71-77.	0.3	7
154	Limited utility of plasma elafin as a biomarker for skin graftâ€versusâ€host disease following allogeneic stem cell transplantation. Clinical and Experimental Dermatology, 2021, 46, 1482-1487.	0.6	7
155	Coagulation factor IX gene transfer to non-human primates using engineered AAV3 capsid and hepatic optimized expression cassette. Molecular Therapy - Methods and Clinical Development, 2021, 23, 98-107.	1.8	7
156	Acute Myeloid Leukemia: Challenges and Real World Data from India. Blood, 2014, 124, 3685-3685.	0.6	7
157	The safety of activated eptacog beta in the management of bleeding episodes and perioperative haemostasis in adult and paediatric haemophilia patients with inhibitors. Haemophilia, 2021, 27, 921-931.	1.0	7
158	Comparison of Clinical Outcomes of Relapsed APL Patients Induced with ATO and Consolidated with Either an Autologous SCT or ATO Based Treatment Regimen.. Blood, 2007, 110, 2880-2880.	0.6	7
159	Part 2: Making the â€œunprovenâ€•â€œprovenâ€• Cytotherapy, 2016, 18, 120-123.	0.3	6
160	Efficacy of narrow band UVB in the treatment of cutaneous GvHD: an Indian experience. Bone Marrow Transplantation, 2016, 51, 988-990.	1.3	6
161	Economic Challenges in Hematopoietic Cell Transplantation: How Will New and Established Programs Face the Growing Costs?. Biology of Blood and Marrow Transplantation, 2017, 23, 1815-1816.	2.0	6
162	Generation of an integration-free iPSC line (CSCRi005-A) from erythroid progenitor cells of a healthy Indian male individual. Stem Cell Research, 2018, 29, 148-151.	0.3	6

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163	Lack of grading agreement among international hemostasis external quality assessment programs. Blood Coagulation and Fibrinolysis, 2018, 29, 111-119.	0.5	6
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320	Endothelial Activation and Stress Index (EASIX) Measured Pre-Transplant Identifies a Subgroup with High Transplant Related Mortality in Patients with Thalassemia Undergoing Stem Cell Transplantation Using Thiotepa-Treosulfan-Fludarabine Conditioning. Blood, 2021, 138, 1781-1781.	0.6	0
321	Optimization of Robust Diagnostic Strategy for Patients with Fanconi Anaemia (FA)- an Indian Perspective. Blood, 2021, 138, 2187-2187.	0.6	0
322	Ehl Factors at Lower Than Standard Dose Achieve Satisfactory Surgical Haemostatsis in Haemophilia. Blood, 2020, 136, 25-26.	0.6	0