# Alok Srivastava

List of Publications by Year
in descending order

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|---|---|---|---|
| 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 | Disease Status at Transplant has a Significant Impact on Outcomes of Autologous Transplantation (ASCT) in Patients with Hodgkin Lymphomaâ€"A Single Center Experience. Indian Journal of Hematology and Blood Transfusion, 2022, 38, 290-298. | 0.3 | 1 |
| 2 | 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 |
| 3 | Laboratory characterization of obligate carriers of type 3 von Willebrand disease with a potential role for Platelet Function Analyzer (PFAâ€200). International Journal of Laboratory Hematology, 2022, , . | 0.7 | 1 |
| 4 | Identification of novel HPFH-like mutations by CRISPR base editing that elevate the expression of fetal hemoglobin. ELife, 2022, 11, . | 2.8 | 29 |
| 5 | 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 |
| 6 | Haemophilia gene therapyâ€"Update on new country initiatives. Haemophilia, 2022, 28, 61-67. | 1.0 | 8 |
| 7 | Endothelial Activation and Stress Index-Measured Pretransplantation Predicts Transplantation-Related Mortality in Patients with Thalassemia Major Undergoing Transplantation with Thiotepa, Treosulfan, and Fludarabine Conditioning. Transplantation and Cellular Therapy, 2022, 28, 356.e1-356.e6. | 0.6 | 2 |
| 8 | Ageâ€stratified adenoâ€associated virus serotype 3 neutralizing and total antibody prevalence in hemophilia A patients from India. Journal of Medical Virology, 2022, 94, 4542-4547. | 2.5 | 2 |
| 9 | Determination of fibrin clot growth and spatial thrombin propagation in the presence of different types of phospholipid surfaces. Platelets, 2021, 32, 1031-1037. | 1.1 | 3 |
| 10 | 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 |
| 11 | Diagnosis of haemophilia and other inherited bleeding disorders â€•Is a new paradigm needed?. Haemophilia, 2021, 27, 14-20. | 1.0 | 10 |
| 12 | Safety of peripheral blood stem cell harvest in children under anaesthesia in the day care setting â€" A single centre experience. Transfusion and Apheresis Science, 2021, 60, 102962. | 0.5 | 1 |
| 13 | 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 |
| 14 | Management of COVIDâ€19â€associated coagulopathy in persons with haemophilia. Haemophilia, 2021, 27, 41-48. | 1.0 | 14 |
| 15 | Neighborhood poverty and pediatric allogeneic hematopoietic cell transplantation outcomes: a CIBMTR analysis. Blood, 2021, 137, 556-568. | 0.6 | 34 |
| 16 | Outcome of iron reduction therapy in ex-thalassemics. PLoS ONE, 2021, 16, e0238793. | 1.1 | 3 |
| 17 | Indian Society of Hematology and Blood Transfusion (ISHBT) Consensus Document on Hematological Practice During COVID-19 Pandemic. Indian Journal of Hematology and Blood Transfusion, 2021, 37, 1-9. | 0.3 | 1 |
| 18 | Ultrasound and magnetic resonance imaging for the detection of blood: An exâ€vivo study. Haemophilia, 2021, 27, 488-493. | 1.0 | 5 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | 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 |
| 20 | Worldwide Network for Blood and Marrow Transplantation (WBMT) Recommendations Regarding Essential Medications Required To Establish An Early Stage Hematopoietic Cell Transplantation Program. Transplantation and Cellular Therapy, 2021, 27, 267.e1-267.e5. | 0.6 | 6 |
| 21 | Direct Generation of Immortalized Erythroid Progenitor Cell Lines from Peripheral Blood Mononuclear Cells. Cells, 2021, 10, 523. | 1.8 | 10 |
| 22 | 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 |
| 23 | Mutation profile in BCR-ABL1-negative myeloproliferative neoplasms: A single-center experience from India. Hematology/ Oncology and Stem Cell Therapy, 2021, , . | 0.6 | 2 |
| 24 | 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 |
| 25 | The case for equitable haemophilia care. Lancet Haematology,the, 2021, 8, e626. | 2.2 | 2 |
| 26 | NUDT15 c.415C>T Polymorphism Predicts 6-MP Induced Early Myelotoxicity in Patients with Acute Lymphoblastic Leukemia Undergoing Maintenance Therapy. Pharmacogenomics and Personalized Medicine, 2021, Volume 14, 1303-1313. | 0.4 | 0 |
| 27 | 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 |
| 28 | 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 |
| 29 | 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 |
| 30 | 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 |
| 31 | Optimization of Robust Diagnostic Strategy for Patients with Fanconi Anaemia (FA)- an Indian Perspective. Blood, 2021, 138, 2187-2187. | 0.6 | 0 |
| 32 | 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 |
| 33 | Bone Marrow-Derived Mesenchymal Stem Cells Augment Regeneration of Intervertebral Disc in a Reproducible and Validated Mouse Intervertebral Disc Degeneration Model. Neurology India, 2021, 69, 1565. | 0.2 | 4 |
| 34 | Derivation of Clinical-Grade Induced Pluripotent Stem Cell Lines from Erythroid Progenitor Cells in Xenofree Conditions. Methods in Molecular Biology, 2021, , 1. | 0.4 | 1 |
| 35 | Endocrine Challenges and Metabolic Profile in Recipients of Allogeneic Haematopoietic Stem Cell Transplant: A Cross-Sectional Study from Southern India. Indian Journal of Hematology and Blood Transfusion, 2020, 36, 484-490. | 0.3 | 1 |
| 36 | Pharmacokinetics and Efficacy of Generic Melphalan Is Comparable to Innovator Formulation in Patients With Multiple Myeloma Undergoing Autologous Stem Cell Transplantation. Clinical Lymphoma, Myeloma and Leukemia, 2020, 20, 130-135.e1. | 0.2 | 2 |

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|---|---|---|---|
| 37 | 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 |
| 38 | 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 |
| 39 | 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 |
| 40 | Do Bone Density, Bone Microarchitecture, and Body Composition Differ in Recipients of Allogeneic Hematopoietic Stem Cell Transplant? A Cross-Sectional Study from Southern India. Biology of Blood and Marrow Transplantation, 2020, 26, 540-545. | 2.0 | 2 |
| 41 | Impact of imaging modality on clinical outcome in Hodgkin lymphoma in a resource constraint setting. British Journal of Haematology, 2020, 188, 930-934. | 1.2 | 2 |
| 42 | Development of a Clinical Candidate AAV3 Vector for Gene Therapy of Hemophilia B. Human Gene Therapy, 2020, 31, 1114-1123. | 1.4 | 19 |
| 43 | WFH Guidelines for the Management of Hemophilia, 3rd edition. Haemophilia, 2020, 26, 1-158. | 1.0 | 915 |
| 44 | 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 |
| 45 | 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 |
| 46 | 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 |
| 47 | 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 |
| 48 | 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 |
| 49 | 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 |
| 50 | 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 |
| 51 | 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 |
| 52 | Ehl Factors at Lower Than Standard Dose Achieve Satisfactory Surgical Haemostatsis in Haemophilia. Blood, 2020, 136, 25-26. | 0.6 | 0 |
| 53 | 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 |
| 54 | 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 |

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|---|---|---|---|
| 55 | Outcomes Following Allogeneic Stem Cell Transplantation Using Non-sibling Family Donors. Indian Journal of Hematology and Blood Transfusion, 2019, 35, 43-49. | 0.3 | 4 |
| 56 | 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 |
| 57 | 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 |
| 58 | 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 |
| 59 | Evaluation of nonneutralizing antibodies against factor VIII in severe haemophilia A patients from India. Blood Coagulation and Fibrinolysis, 2019, 30, 337-340. | 0.5 | 3 |
| 60 | Fundamentals for a Systematic Approach to Mild and Moderate Inherited Bleeding Disorders: An EHA Consensus Report. HemaSphere, 2019, 3, e286. | 1.2 | 43 |
| 61 | 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 |
| 62 | 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 |
| 63 | 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 |
| 64 | Heterogeneity of Mesenchymal Stromal Cells in Myelodysplastic Syndrome-with Multilineage Dysplasia (MDS-MLD). Indian Journal of Hematology and Blood Transfusion, 2019, 35, 223-232. | 0.3 | 5 |
| 65 | Challenges in managing graft-versus-host disease in developing countries: a perspective. Bone Marrow Transplantation, 2019, 54, 641-647. | 1.3 | 6 |
| 66 | Fludarabine and Cyclophosphamide Based Conditioning Is Associated with Good Outcomes in Patients Undergoing Matched Sibling Donor Transplants for Aplastic Anaemia. Blood, 2019, 134, 3272-3272. | 0.6 | 1 |
| 67 | Impact of Graft Versus Host Disease on Outcome of Allogeneic Haematopoietic Stem Cell Transplantation for Thalassemia Major - Comparison of Bone Marrow Vs Peripheral Blood Stem Cell Grafts. Blood, 2019, 134, 4537-4537. | 0.6 | 0 |
| 68 | Treosulfan Metabolite (S, S-EBDM) Pharmacokinetics Influences Regimen Related Toxicity in Patients with Beta Thalassaemia Major Undergoing HSCT. Blood, 2019, 134, 1977-1977. | 0.6 | 0 |
| 69 | 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 |
| 70 | Principles of haemophilia care: The Asiaâ€Pacific perspective. Haemophilia, 2018, 24, 366-375. | 1.0 | 15 |
| 71 | 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 |
| 72 | Oxidation of factor VIII increases its immunogenicity in mice with severe hemophilia A. Cellular Immunology, 2018, 325, 64-68. | 1.4 | 4 |

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|---|---|---|---|
| 73 | World bleeding disorders registry: The pilot study. Haemophilia, 2018, 24, e113-e116. | 1.0 | 13 |
| 74 | 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 |
| 75 | Management of Hemophilic Cysts and Pseudotumors ofÂthe Hand in Bleeding Disorders: A Case Series. Journal of Hand Surgery, 2018, 43, 486.e1-486.e9. | 0.7 | 5 |
| 76 | Lack of grading agreement among international hemostasis external quality assessment programs. Blood Coagulation and Fibrinolysis, 2018, 29, 111-119. | 0.5 | 6 |
| 77 | 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 |
| 78 | 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 |
| 79 | 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 |
| 80 | Measuring activities and participation in persons with haemophilia: A systematic review of commonly used instruments. Haemophilia, 2018, 24, e33-e49. | 1.0 | 21 |
| 81 | 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 |
| 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 | Performance of an In-House Real-Time PCR Assay for Detecting Cytomegalovirus Infection among Transplant Patients from a Tertiary Care Centre. Indian Journal of Medical Microbiology, 2018, 36, 241-246. | 0.3 | 4 |
| 84 | Sensitivity and Robustness of Spatially Dependent Thrombin Generation and Fibrin Clot Propagation. Biophysical Journal, 2018, 115, 2461-2473. | 0.2 | 23 |
| 85 | Quality control guidelines for clinical-grade human induced pluripotent stem cell lines. Regenerative Medicine, 2018, 13, 859-866. | 0.8 | 147 |
| 86 | 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 |
| 87 | 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 |
| 88 | 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 |
| 89 | Typeâ€3 von Willebrand disease in Indiaâ€"Clinical spectrum and molecular profile. Haemophilia, 2018, 24, 930-940. | 1.0 | 8 |
| 90 | A Low Incidence of Cytomegalo Virus Infection Following Allogeneic Hematopoietic Stem Cell Transplantation Despite a High Seroprevalence. Indian Journal of Hematology and Blood Transfusion, 2018, 34, 636-642. | 0.3 | 6 |

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|---|---|---|---|
| 91 | 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 |
| 92 | Principles of haemophilia care: The asia pacific perspective. Response. Haemophilia, 2018, 24, e243-e244. | 1.0 | 0 |
| 93 | Emerging therapies for haemophilia â€•Global perspective. Haemophilia, 2018, 24, 15-21. | 1.0 | 24 |
| 94 | Very Long Term Follow-up Data of Pediatric Acute Promyelocytic Leukemia Treated with Upfront Arsenic-Trioxide Based Regimens. Blood, 2018, 132, 1400-1400. | 0.6 | 3 |
| 95 | Higher Incidence of Graft Rejection in Non-Sibling Fully Matched Related Donor Stem Cell Transplants for Thalassemia Major: A Cautionary Note. Blood, 2018, 132, 2178-2178. | 0.6 | 4 |
| 96 | Haplo-Identical Transplants Using Post Transplant Cyclophosphamide (PTCy) Are Associated with Good Outcomes If Transplanted with Early Disease - a Single Centre Analysis from India. Blood, 2018, 132, 4652-4652. | 0.6 | 2 |
| 97 | Outcome of Immune Tolerance Induction Using an Extended Half-Life Clotting Factor Concentrate â€" Recombinant Factor VIII Fc (Eloctateâ„¢) â€" a Report from India. Blood, 2018, 132, 2494-2494. | 0.6 | 2 |
| 98 | Peripheral T cell lymphoma: Clinico-pathological characteristics & outcome from a tertiary care centre in south India. Indian Journal of Medical Research, 2018, 147, 464. | 0.4 | 5 |
| 99 | Iron Reduction Therapy in Ex-Thalassemics - Long Term Outcome. Blood, 2018, 132, 4591-4591. | 0.6 | 0 |
| 100 | SMALL Molecules Mediated Hematopoietic STEM and Progenitor CELLS Expansion for GENE Editing Application. Blood, 2018, 132, 5803-5803. | 0.6 | 0 |
| 101 | Targeted IV Vs Oral Busulfan in Very Young Children with Thalassemia Major Undergoing Matched Allogeneic Haematopoietic Stem Cell Transplantation. Blood, 2018, 132, 5707-5707. | 0.6 | 0 |
| 102 | Management of Relapse in Acute Promyelocytic Leukemia Treated with Upfront Arsenic Trioxide Based Regimens. Blood, 2018, 132, 666-666. | 0.6 | 3 |
| 103 | Systematic evaluation of markers used for the identification of human induced pluripotent stem cells. Biology Open, 2017, 6, 100-108. | 0.6 | 22 |
| 104 | ATP-binding casette transporter expression in acute myeloid leukemia: association within vitrocytotoxicity and prognostic markers. Pharmacogenomics, 2017, 18, 235-244. | 0.6 | 24 |
| 105 | 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 |
| 106 | Population pharmacokinetics of fludarabine in patients with aplastic anemia and Fanconi anemia undergoing allogeneic hematopoietic stem cell transplantation. Bone Marrow Transplantation, 2017, 52, 977-983. | 1.3 | 5 |
| 107 | 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 |
| 108 | Efficacy and safety of a <scp>VWF</scp>/<scp>FVIII</scp> concentrate (wilate<sup>Â®</sup>) in inherited von Willebrand disease patients undergoing surgical procedures. Haemophilia, 2017, 23, 264-272. | 1.0 | 29 |

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|---|---|---|---|
| 109 | 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 |
| 110 | Cure for thalassemia major â€“ from allogeneic hematopoietic stem cell transplantation to gene therapy. Haematologica, 2017, 102, 214-223. | 1.7 | 57 |
| 111 | Invasive fungal infection following chemotherapy for acute myeloid leukaemiaâ€"Experience from a developing country. Mycoses, 2017, 60, 686-691. | 1.8 | 26 |
| 112 | Coexistence of aberrant hematopoietic and stromal elements in myelodysplastic syndromes. Blood Cells, Molecules, and Diseases, 2017, 66, 37-46. | 0.6 | 7 |
| 113 | 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 |
| 114 | Choosing outcome assessment tools in haemophilia care and research: a multidisciplinary perspective. Haemophilia, 2017, 23, 11-24. | 1.0 | 63 |
| 115 | Allogeneic stem cell transplantation for thalassemia major in India. Pediatric Hematology Oncology Journal, 2017, 2, 114-120. | 0.1 | 7 |
| 116 | Where thereâ€™s a will, thereâ€™s a way: establishing hematopoietic stem cell transplantation in Myanmar. Blood Advances, 2017, 1, 65-69. | 2.5 | 3 |
| 117 | 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 |
| 118 | Comparative double-blind randomized trial of 2 rituximab products in patients with CD20+ diffuse large B-cell lymphoma (DLBCL).. Journal of Clinical Oncology, 2017, 35, 7550-7550. | 0.8 | 2 |
| 119 | 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 |
| 120 | Reply to the letter of O'Mahoney et al.. Haemophilia, 2016, 22, e209-11. | 1.0 | 0 |
| 121 | Part 2: Making the â€œunprovenâ€•â€œprovenâ€• Cytotherapy, 2016, 18, 120-123. | 0.3 | 6 |
| 122 | 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 |
| 123 | Treatment rates of paediatric acute myeloid leukaemia: a view from three tertiary centres in India â€“ response to Gupta etÂal. British Journal of Haematology, 2016, 175, 347-349. | 1.2 | 3 |
| 124 | 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 |
| 125 | Correlating clinical and radiological assessment of joints in haemophilia: results of a cross sectional study. Haemophilia, 2016, 22, 925-933. | 1.0 | 44 |
| 126 | Role of endovascular embolization in treatment of acute bleeding complications in haemophilia patients. British Journal of Radiology, 2016, 89, 20151064. | 1.0 | 5 |

| # | Article | IF | Citations |
|---|---|---|---|
| 127 | Population pharmacokinetics of Daunorubicin in adult patients with acute myeloid leukemia. Cancer Chemotherapy and Pharmacology, 2016, 78, 1051-1058. | 1.1 | 20 |
| 128 | Standardizing patient outcomes measurement to improve haemophilia care. Haemophilia, 2016, 22, 651-653. | 1.0 | 7 |
| 129 | Clinical, Hematological and Molecular Analysis of Homozygous Hb E (HBB: c.79Gâ€‰>â€‰A) in the Indian Population. Hemoglobin, 2016, 40, 16-19. | 0.4 | 4 |
| 130 | Factor VIII/factor IX prophylaxis for severe hemophilia. Seminars in Hematology, 2016, 53, 3-9. | 1.8 | 19 |
| 131 | Efficacy of narrow band UVB in the treatment of cutaneous GvHD: an Indian experience. Bone Marrow Transplantation, 2016, 51, 988-990. | 1.3 | 6 |
| 132 | AtypicalBCR-ABL1fusion transcripts in adult B-acute lymphoblastic leukemia, including a novel fusion transcript-e8a1. Leukemia and Lymphoma, 2016, 57, 2481-2484. | 0.6 | 5 |
| 133 | Part 1: Defining unproven cellular therapies. Cytotherapy, 2016, 18, 117-119. | 0.3 | 33 |
| 134 | Management of Relapsed Acute Promyelocytic Leukemia Post ATO Upfront Therapy: Open-Labeled Phase II Study Evaluating Role of Proteasome Inhibition. Blood, 2016, 128, 446-446. | 0.6 | 1 |
| 135 | A 5'UTR Polymorphism in NT5E Gene Influences Outcome in Patients with Acute Myeloid Leukemia Undergoing Hematopoietic Stem Cell Transplantation with Fludarabine Based Conditioning Regimen. Blood, 2016, 128, 984-984. | 0.6 | 2 |
| 136 | Comparison of Pharmacokinetics and Pharmacodynamics of Two Anti-CD20 Monoclonal Antibodies | | |

| # | Article | IF | Citations |
|---|---|---|---|
| 145 | Assessments of outcome in haemophilia â€“ what is the added value of <scp>QoL</scp> tools?. Haemophilia, 2015, 21, 430-435. | 1.0 | 25 |
| 146 | Blood-Induced Arthropathy in Hemophilia: Mechanisms and Heterogeneity. Seminars in Thrombosis and Hemostasis, 2015, 41, 832-837. | 1.5 | 31 |
| 147 | Genetic modifiers of secondary iron overload in beta thalassemia major. Blood Cells, Molecules, and Diseases, 2015, 54, 242-243. | 0.6 | 1 |
| 148 | Diagnostic Accuracy of Ultrasound for Assessment of Hemophilic Arthropathy: MRI Correlation. American Journal of Roentgenology, 2015, 204, W336-W347. | 1.0 | 98 |
| 149 | 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 |
| 150 | Cryotherapy for acute haemarthrosis in haemophilia â€“ attempts to understand the â€˜ice ageâ€™ practice. Haemophilia, 2015, 21, e103-5. | 1.0 | 5 |
| 151 | Positioning a Scientific Community on Unproven Cellular Therapies: The 2015 International Society for Cellular Therapy Perspective. Cytotherapy, 2015, 17, 1663-1666. | 0.3 | 44 |
| 152 | Understanding the Molecular Basis of Heterogeneity in Induced Pluripotent Stem Cells. Cellular Reprogramming, 2015, 17, 427-440. | 0.5 | 2 |
| 153 | Prognostic Significance of Immunophenotypic Composition of B Cell Acute Lymphoblastic Leukemia at Diagnosis: A Novel Immunophenotype Based Risk Score. Blood, 2015, 126, 2620-2620. | 0.6 | 1 |
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