

Sukesh Chandran Nair

List of Publications by Year in descending order

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Version: 2024-02-01

48
papers

336
citations

1040056

9
h-index

940533

16
g-index

48
all docs

48
docs citations

48
times ranked

453
citing authors

#	ARTICLE	IF	CITATIONS
1	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.	2.5	43
2	Collection and Transport of Samples for Laboratory Testing in von Willebrand's Disease (VWD): Time for a Reappraisal?. Thrombosis and Haemostasis, 2001, 86, 1589-1590.	3.4	29
3	International Council for Standardization in Haematology (ICSH) recommendations for laboratory measurement of ADAMTS13. International Journal of Laboratory Hematology, 2020, 42, 685-696.	1.3	26
4	International Council for Standardization in Haematology (ICSH) recommendations for processing of blood samples for coagulation testing. International Journal of Laboratory Hematology, 2021, 43, 1272-1283.	1.3	26
5	Sensitivity and Robustness of Spatially Dependent Thrombin Generation and Fibrin Clot Propagation. Biophysical Journal, 2018, 115, 2461-2473.	0.5	23
6	International Council for Standardisation in Haematology (ICSH) recommendations for collection of blood samples for coagulation testing. International Journal of Laboratory Hematology, 2021, 43, 571-580.	1.3	17
7	Is pre-operative assessment of coagulation profile with Thrombelastography (TEG) useful in predicting venous thromboembolism (VTE) following orthopaedic surgery?. Journal of Clinical Orthopaedics and Trauma, 2016, 7, 225-229.	1.5	16
8	Low Volume Plasma Exchange and Low Dose Steroid Improve Survival in Patients With Alcohol-Related Acute on Chronic Liver Failure and Severe Alcoholic Hepatitis – Preliminary Experience. Journal of Clinical and Experimental Hepatology, 2022, 12, 372-378.	0.9	14
9	Diagnosis and Management of von Willebrand Disease: A Developing Country Perspective. Seminars in Thrombosis and Hemostasis, 2011, 37, 587-594.	2.7	13
10	Management of relapse in acute promyelocytic leukaemia treated with upfront arsenic trioxide-based regimens. British Journal of Haematology, 2021, 192, 292-299.	2.5	13
11	Patient blood management in India - Review of current practices and feasibility of applying appropriate standard of care guidelines. A position paper by an interdisciplinary expert group. Journal of Anaesthesiology Clinical Pharmacology, 2021, 37, 3.	0.7	10
12	Cytomorphometric Neutrophil and Monocyte Markers May Strengthen the Diagnosis of Sepsis. Journal of Intensive Care Medicine, 2018, 33, 656-662.	2.8	9
13	Prevalence of FVIII inhibitors in severe haemophilia A patients: Effect of treatment and genetic factors in an Indian population. Haemophilia, 2019, 25, 67-74.	2.1	9
14	International Society on Thrombosis and Haemostasis core curriculum project: Core competencies in laboratory thrombosis and hemostasis. Journal of Thrombosis and Haemostasis, 2019, 17, 1848-1859.	3.8	8
15	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.	1.4	8
16	Real world data with concurrent retinoic acid and arsenic trioxide for the treatment of acute promyelocytic leukemia. Blood Cancer Journal, 2022, 12, 22.	6.2	8
17	Coexistence of aberrant hematopoietic and stromal elements in myelodysplastic syndromes. Blood Cells, Molecules, and Diseases, 2017, 66, 37-46.	1.4	7
18	Plasma Von Willebrand Factor Levels Predict Survival in COVID-19 Patients Across the Entire Spectrum of Disease Severity. Indian Journal of Hematology and Blood Transfusion, 2022, 38, 333-340.	0.6	7

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19	Lack of grading agreement among international hemostasis external quality assessment programs. <i>Blood Coagulation and Fibrinolysis</i> , 2018, 29, 111-119.	1.0	6
20	Six Novel Mutations Including Triple Heterozygosity for Phe31Ser, 514delT and 516Tâ†’G Mutations in Factor X Gene Is Responsible for Congenital Factor X Deficiency in Patients of Indian and Nepali Origin.. <i>Blood</i> , 2004, 104, 1042-1042.	1.4	6
21	Heterogeneity of Mesenchymal Stromal Cells in Myelodysplastic Syndrome-with Multilineage Dysplasia (MDS-MLD). <i>Indian Journal of Hematology and Blood Transfusion</i> , 2019, 35, 223-232.	0.6	5
22	A study to compare Hematopoietic Progenitor Cell count determined on a nextâ€’generation automated cell counter with flow cytometric CD34 count in peripheral blood and the harvested peripheral blood stem cell graft from autologous and allogenic donors. <i>International Journal of Laboratory Hematology</i> , 2021, 43, 76-83.	1.3	5
23	Resource utilization and cost effectiveness of treating acute promyelocytic leukaemia using generic arsenic trioxide. <i>British Journal of Haematology</i> , 2020, 189, 269-278.	2.5	4
24	Evaluation of nonneutralizing antibodies against factor VIII in severe haemophilia A patients from India. <i>Blood Coagulation and Fibrinolysis</i> , 2019, 30, 337-340.	1.0	3
25	Determination of fibrin clot growth and spatial thrombin propagation in the presence of different types of phospholipid surfaces. <i>Platelets</i> , 2021, 32, 1031-1037.	2.3	3
26	Polymorphisms in Coagulant and Inflammatory Genes Modify the Phenotype of Severe Hemophilia A and B.. <i>Blood</i> , 2006, 108, 1009-1009.	1.4	3
27	Evaluation of Factor VIII as a Risk Factor in Indian Patients with DVT. <i>Surgery Research and Practice</i> , 2015, 2015, 1-4.	0.5	2
28	Sitosterolemia: Four Cases of an Uncommon Cause of Hemolytic Anemia (Mediterranean) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 387 Td 2021, 37, 157-161.	0.6	2
29	Outcome of Immune Tolerance Induction Using an Extended Half-Life Clotting Factor Concentrate â€” Recombinant Factor VIII Fc (Eloctateâ„¢) â€” a Report from India. <i>Blood</i> , 2018, 132, 2494-2494.	1.4	2
30	D-dimer levels in patients with thromboangiitis obliterans. <i>The National Medical Journal of India</i> , 2019, 32, 134.	0.3	2
31	Analytical performance of a point-of-care device in monitoring patients on oral anticoagulation with vitamin K antagonists. <i>Phlebology</i> , 2016, 31, 660-667.	1.2	1
32	Does hemodialysis need to be initiated to improve platelet function in CKD G5 patients? A pilot prospective, observational cohort study. <i>Indian Journal of Nephrology</i> , 2021, 31, 43.	0.5	1
33	High fluorescent lymphocyte cell count and scattergram patterns on the Sysmex XN series cell countersâ€”Novel parameters in early and reliable diagnosis of dengue on a background of acute febrile illness. <i>International Journal of Laboratory Hematology</i> , 2021, 43, O156-O160.	1.3	1
34	A Polymorphism In Interferon Gamma Gene Impacts the Extent of Joint Damage In Patients with Severe Hemophilia. <i>Blood</i> , 2010, 116, 546-546.	1.4	1
35	Real World Data of Concurrent Arsenic Trioxide and All-Trans Retinoic Acid with Minimal Use of Anthracycline in the Treatment of Acute Promyelocytic Leukemia. <i>Blood</i> , 2021, 138, 2338-2338.	1.4	1
36	Laboratory characterization of obligate carriers of type 3 von Willebrand disease with a potential role for Platelet Function Analyzer (PFAâ€”200). <i>International Journal of Laboratory Hematology</i> , 2022, , .	1.3	1

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37	Clinical utility of activated partial thromboplastin time clot waveform analysis and thrombin generation test in the evaluation of bleeding phenotype in Hemophilia A. Indian Journal of Pathology and Microbiology, 2021, 64, 117-122.	0.2	1
38	Plasma Coagulation Tests for Detection of Antiphospholipid Antibodies: What's Good, and What Might Be Improved?. Indian Journal of Hematology and Blood Transfusion, 2019, 35, 407-408.	0.6	0
39	Unusual Morphological and Automated Hematology Analyzer Features in 3 Cases of B-cell Malignancy-associated Type I Cryoglobulinemic Vasculitis. Indian Journal of Hematology and Blood Transfusion, 2021, 37, 658-663.	0.6	0
40	Utility of International Society on Thrombosis and Hemostasis Bleeding Assessment Tool (ISTH-BAT) in Patients with Inherited Bleeding Disorders: A Cross-Sectional Study from Southern India. Indian Journal of Hematology and Blood Transfusion, 2022, 38, 122-131.	0.6	0
41	Molecular Genetics of Hereditary Prothrombin Deficiency in Indian Patients: Identification of a Novel Ala362→Thr (Prothrombin Vellore 1) Mutation by Conformation Sensitive Gel Electrophoresis.. Blood, 2004, 104, 1037-1037.	1.4	0
42	Two Mutations (1717 T→C, GP IX;124del145, GPIb beta) Occur Frequently among Patients with Bernard Soulier Syndrome in India.. Blood, 2006, 108, 1096-1096.	1.4	0
43	Identifying Myelodysplastic Syndrome Among Patients with Anemia: Neutrophil Characteristics Computed by a Cell Counter (Beckman Coulter LH750), An Effective Screening Tool. Blood, 2011, 118, 5027-5027.	1.4	0
44	Community Based Evaluation of Prevalence of Inhibitors in Patients with Severe Hemophilia A in India and Their Correlation with Environmental and Genetic Factors. Blood, 2012, 120, 3380-3380.	1.4	0
45	Genetic Diagnosis of Inherited Bleeding Disorders in 1250 Proband From India: A Single Centre Experience. Blood, 2012, 120, 1127-1127.	1.4	0
46	Coagulopathy in Acute Promyelocytic Leukemia: Strategies to Improve Assessment of Hemostatic Risk. Blood, 2015, 126, 3758-3758.	1.4	0
47	Molecular Basis of Von Willebrand Disease in Patients from India. Blood, 2015, 126, 1102-1102.	1.4	0
48	Ehl Factors at Lower Than Standard Dose Achieve Satisfactory Surgical Haemostasis in Haemophilia. Blood, 2020, 136, 25-26.	1.4	0