Sukesh Chandran Nair

List of Publications by Year
in descending order

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48	336	1040056	940533
papers	citations	9	16
		h-index	g-index

48	48	48	453
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	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
2	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
3	Low Volume Plasma Exchange and Low Dose Steroid Improve Survival in Patients With Alcohol-Related Acute on Chronic Liver Failure and Severe Alcoholic Hepatitis â€“ Preliminary Experience. Journal of Clinical and Experimental Hepatology, 2022, 12, 372-378.	0.9	14
4	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, , .	1.3	1
5	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
6	Determination of fibrin clot growth and spatial thrombin propagation in the presence of different types of phospholipid surfaces. Platelets, 2021, 32, 1031-1037.	2.3	3
7	Management of relapse in acute promyelocytic leukaemia treated with upâ€front arsenic trioxideâ€based regimens. British Journal of Haematology, 2021, 192, 292-299.	2.5	13
8	Sitosterolemia: Four Cases of an Uncommon Cause of Hemolytic Anemia (Mediterranean) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 467 Td (2021, 37, 157-161.	0.6	2
9	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. International Journal of Laboratory Hematology, 2021, 43, 76-83.	1.3	5
10	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
11	Does hemodialysis need to be initiated to improve platelet function in CKD G5 patients? A pilot prospective, observational cohort study. Indian Journal of Nephrology, 2021, 31, 43.	0.5	1
12	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. International Journal of Laboratory Hematology, 2021, 43, O156-O160.	1.3	1
13	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
14	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
15	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
16	Real World Data of Concurrent Arsenic Trioxide and All-Trans Retinoic Acid with Minimal Use of Anthracycline in the Treatment of Acute Promyelocytic Leukemia. Blood, 2021, 138, 2338-2338.	1.4	1
17	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
18	Resource utilization and cost effectiveness of treating acute promyelocytic leukaemia using generic arsenic trioxide. British Journal of Haematology, 2020, 189, 269-278.	2.5	4

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19	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
20	Ehl Factors at Lower Than Standard Dose Achieve Satisfactory Surgical Haemostatsis in Haemophilia. Blood, 2020, 136, 25-26.	1.4	0
21	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
22	Plasma Coagulation Tests for Detection of Antiphospholipid Antibodies: Whatâ€™s Good, and What Might Be Improved?. Indian Journal of Hematology and Blood Transfusion, 2019, 35, 407-408.	0.6	0
23	Evaluation of nonneutralizing antibodies against factor VIII in severe haemophilia A patients from India. Blood Coagulation and Fibrinolysis, 2019, 30, 337-340.	1.0	3
24	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
25	Heterogeneity of Mesenchymal Stromal Cells in Myelodysplastic Syndrome-with Multilineage Dysplasia (MDS-MLD). Indian Journal of Hematology and Blood Transfusion, 2019, 35, 223-232.	0.6	5
26	D-dimer levels in patients with thromboangiitis obliterans. The National Medical Journal of India, 2019, 32, 134.	0.3	2
27	Lack of grading agreement among international hemostasis external quality assessment programs. Blood Coagulation and Fibrinolysis, 2018, 29, 111-119.	1.0	6
28	Cytomorphometric Neutrophil and Monocyte Markers May Strengthen the Diagnosis of Sepsis. Journal of Intensive Care Medicine, 2018, 33, 656-662.	2.8	9
29	Sensitivity and Robustness of Spatially Dependent Thrombin Generation and Fibrin Clot Propagation. Biophysical Journal, 2018, 115, 2461-2473.	0.5	23
30	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.	1.4	2
31	Coexistence of aberrant hematopoietic and stromal elements in myelodysplastic syndromes. Blood Cells, Molecules, and Diseases, 2017, 66, 37-46.	1.4	7
32	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
33	Analytical performance of a point-of-care device in monitoring patients on oral anticoagulation with vitamin K antagonists. Phlebology, 2016, 31, 660-667.	1.2	1
34	Evaluation of Factor VIII as a Risk Factor in Indian Patients with DVT. Surgery Research and Practice, 2015, 2015, 1-4.	0.5	2
35	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
36	Coagulopathy in Acute Promyelocytic Leukemia: Strategies to Improve Assessment of Hemostatic Risk. Blood, 2015, 126, 3758-3758.	1.4	0

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37	Molecular Basis of Von Willebrand Disease in Patients from India. Blood, 2015, 126, 1102-1102.	1.4	0
38	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
39	Genetic Diagnosis of Inherited Bleeding Disorders in 1250 Probands From India: A Single Centre Experience. Blood, 2012, 120, 1127-1127.	1.4	0
40	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
41	Diagnosis and Management of<scp>v</scp>on Willebrand Disease: A Developing Country Perspective. Seminars in Thrombosis and Hemostasis, 2011, 37, 587-594.	2.7	13
42	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
43	A Polymorphism In Interferon Gamma Gene Impacts the Extent of Joint Damage In Patients with Severe Hemophilia. Blood, 2010, 116, 546-546.	1.4	1
44	Two Mutations (1717 Tã€‰C, GP IX;124del145, GPIb beta) Occur Frequently among Patients with Bernard Soulier Syndrome in India.. Blood, 2006, 108, 1096-1096.	1.4	0
45	Polymorphisms in Coagulant and Inflammatory Genes Modify the Phenotype of Severe Hemophilia A and B.. Blood, 2006, 108, 1009-1009.	1.4	3
46	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.. Blood, 2004, 104, 1042-1042.	1.4	6
47	Molecular Genetics of Hereditary Prothrombin Deficiency in Indian Patients: Identification of a Novel Ala362â†'Thr (Prothrombin Vellore 1) Mutation by Conformation Sensitive Gel Electrophoresis.. Blood, 2004, 104, 1037-1037.	1.4	0
48	Collection and Transport of Samples for Laboratory Testing in von Willebrandâ€™s Disease (VWD): Time for a Reappraisal?. Thrombosis and Haemostasis, 2001, 86, 1589-1590.	3.4	29