Kanjaksha Ghosh

List of Publications by Year
in descending order

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544 papers	6,840 citations	101543 36 h-index	182427 51 g-index
557 all docs	557 docs citations	557 times ranked	7594 citing authors

#	Article	IF	Citations
1	Hereditary thrombophilia as a cause of Budd-Chiari syndrome: A study from Western India. Hepatology, 2001, 34, 666-670.	7.3	117
2	Acquired hemophilia A: Diagnosis, aetiology, clinical spectrum and treatment options. Autoimmunity Reviews, 2011, 10, 311-316.	5.8	103
3	Matrix metalloproteinase and its drug targets therapy in solid and hematological malignancies: An overview. Mutation Research - Reviews in Mutation Research, 2013, 753, 7-23.	5.5	88
4	Glanzmann's thrombasthenia: updated. Platelets, 2002, 13, 387-393.	2.3	84
5	Contribution of natural anticoagulant and fibrinolytic factors in modulating the clinical severity of haemophilia patients. British Journal of Haematology, 2007, 138, 541-544.	2.5	80
6	Asialoglycoprotein receptor targeted delivery of doxorubicin nanoparticles for hepatocellular carcinoma. Drug Delivery, 2017, 24, 20-29.	5.7	78
7	Reproducible methodology for the isolation of mesenchymal stem cells from human umbilical cord and its potential for cardiomyocyte generation. Journal of Tissue Engineering and Regenerative Medicine, 2008, 2, 394-399.	2.7	76
8	Fibrinolysis, inhibitors of blood coagulation, and monocyte derived coagulant activity in acute malaria. , 1997, 54, 23-29.		70
9	Osteoporosis in young haemophiliacs from western India. American Journal of Hematology, 2007, 82, 453-457.	4.1	70
10	Pathogenesis of anemia in malaria: a concise review. Parasitology Research, 2007, 101, 1463-1469.	1.6	70
11	Sickle cell disease in India. Current Opinion in Hematology, 2014, 21, 215-223.	2.5	68
12	Response to hydroxyurea in Î² thalassemia major and intermedia: Experience in western India. Clinica Chimica Acta, 2009, 407, 10-15.	1.1	67
13	JAK2Mutations Across a Spectrum of Venous Thrombosis Cases: Table 1. American Journal of Clinical Pathology, 2010, 134, 82-85.	0.7	66
14	Strongyloides stercoralis septicaemia following steroid therapy for eosinophilia: report of three cases. Transactions of the Royal Society of Tropical Medicine and Hygiene, 2007, 101, 1163-1165.	1.8	64
15	Venous Thromboembolism in Young Patients From Western India: A Study. Clinical and Applied Thrombosis/Hemostasis, 2001, 7, 158-165.	1.7	62
16	Changes in platelet glycoprotein receptors after smoking â€“ a flow cytometric study. Platelets, 2001, 12, 20-26.	2.3	60
17	Hydroxyurea in sickle cell diseaseâ€”A study of clinico-pharmacological efficacy in the Indian haplotype. Blood Cells, Molecules, and Diseases, 2009, 42, 25-31.	1.4	58
18	Regional heterogeneity of Î²-thalassemia mutations in the multi ethnic Indian population. Blood Cells, Molecules, and Diseases, 2009, 42, 241-246.	1.4	56

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19	Advances in autoimmune lymphoproliferative syndromes. European Journal of Haematology, 2011, 87, 1-9.	2.2	53
20	Pathophysiology of acquired von Willebrand disease: a concise review. European Journal of Haematology, 2011, 87, 99-106.	2.2	53
21	Epidemiology of Î²â€thalassaemia in Western India: mapping the frequencies and mutations in subâ€regions of Maharashtra and Gujarat. British Journal of Haematology, 2010, 149, 739-747.	2.5	52
22	Role of epsilon amino caproic acid in the management of haemophilic patients with inhibitors. Haemophilia, 2004, 10, 58-62.	2.1	50
23	Autoimmune lymphoproliferative syndrome caused by a homozygous null FAS ligand (FASLG) mutation. Journal of Allergy and Clinical Immunology, 2013, 131, 486-490.	2.9	50
24	Violence against doctors: A wake-up call. Indian Journal of Medical Research, 2018, 148, 130.	1.0	48
25	Immune Response to FVIII in Hemophilia A: An Overview of Risk Factors. Clinical Reviews in Allergy and Immunology, 2009, 37, 58-66.	6.5	46
26	Management of Haemophilia in Developing Countries: Challenges and Options. Indian Journal of Hematology and Blood Transfusion, 2016, 32, 347-355.	0.6	45
27	VKORC1 and CYP2C9 genotype distribution in Asian countries. Thrombosis Research, 2014, 134, 537-544.	1.7	44
28	Development of inhibitors in patients with haemophilia from India. Haemophilia, 2001, 7, 273-278.	2.1	43
29	Imaging the interaction between dengue 2 virus and human blood platelets using atomic force and electron microscopy. Journal of Electron Microscopy, 2008, 57, 113-118.	0.9	42
30	Iron deficiency as a risk factor for first febrile seizure. Indian Pediatrics, 2010, 47, 437-439.	0.4	42
31	Hereditary thrombophilia in cerebral venous thrombosis. Blood Coagulation and Fibrinolysis, 2013, 24, 540-543.	1.0	42
32	Down-regulation of miR-199b associated with imatinib drug resistance in 9q34.1 deleted BCR/ABL positive CML patients. Gene, 2014, 542, 109-112.	2.2	42
33	Tuberculosis and female reproductive health. Journal of Postgraduate Medicine, 2011, 57, 307-313.	0.4	40
34	Dengue 2 virus inhibitsin vitromegakaryocytic colony formation and induces apoptosis in thrombopoietin-inducible megakaryocytic differentiation from cord blood CD34+ cells. FEMS Immunology and Medical Microbiology, 2008, 53, 46-51.	2.7	38
35	Efficacy of fixed low dose hydroxyurea in Indian children with sickle cell anemia: A single centre experience. Indian Pediatrics, 2013, 50, 929-933.	0.4	38
36	Elevated Procoagulant Endothelial and Tissue Factor Expressing Microparticles in Women with Recurrent Pregnancy Loss. PLoS ONE, 2013, 8, e81407.	2.5	38

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37	Factor VIII and IX gene polymorphisms and carrier analysis in Indian population. , 1997, 54, 271-275.		37
38	Intracranial haemorrhage in severe haemophilia: prevalence and outcome in a developing country. Haemophilia, 2005, 11, 459-462.	2.1	37
39	Flow cytometric osmotic fragilityâ€"An effective screening approach for red cell membranopathies. Cytometry Part B - Clinical Cytometry, 2011, 80B, 186-190.	1.5	37
40	Guidelines for screening, diagnosis and management of hemoglobinopathies. Indian Journal of Human Genetics, 2014, 20, 101.	0.7	37
41	Sickle cell disease in tribal populations in India. Indian Journal of Medical Research, 2015, 141, 509-15.	1.0	37
42	Thrombophilic dimension of Budd chiari syndrome and portal venous thrombosis â€" A concise review. Thrombosis Research, 2011, 127, 505-512.	1.7	36
43	Erythropoiesis in Malaria Infections and Factors Modifying the Erythropoietic Response. Anemia, 2016, 2016, 1-8.	1.7	36
44	Blood coagulation in falciparum malariaâ€"a review. Parasitology Research, 2008, 102, 571-576.	1.6	35
45	First-time development of FVIII inhibitor in haemophilia patients during the postoperative period. Haemophilia, 2002, 8, 776-780.	2.1	34
46	Feasibility of a Newborn Screening and Follow-up Programme for Sickle Cell Disease among South Gujarat (India) Tribal Populations. Journal of Medical Screening, 2015, 22, 1-7.	2.3	34
47	Prevalence and Molecular Characterization of Î±-Thalassemia Syndromes among Indians. Genetic Testing and Molecular Biomarkers, 2008, 12, 177-180.	1.7	33
48	Newborn Screening Shows a High Incidence of Sickle Cell Anemia in Central India. Hemoglobin, 2012, 36, 316-322.	0.8	33
49	Effect of hydroxyurea on the transfusion requirements in patients with severe HbE-Â-thalassaemia: a genotypic and phenotypic study. Journal of Clinical Pathology, 2010, 63, 147-150.	2.0	32
50	Rare coagulation factor deficiencies: a countrywide screening data from India. Haemophilia, 2014, 20, 575-581.	2.1	32
51	Neonatal Screening and the Clinical Outcome in Children with Sickle Cell Disease in Central India. PLoS ONE, 2016, 11, e0147081.	2.5	32
52	Mycobacterium tuberculosis infection precipitates SLE in patients from endemic areas. Rheumatology International, 2009, 29, 1047-1050.	3.0	31
53	REVIEW ARTICLE: Antiâ€phospholipid Antibodies and Other Immunological Causes of Recurrent Foetal Loss â€" A Review of Literature of Various Therapeutic Protocols. American Journal of Reproductive Immunology, 2009, 62, 9-24.	1.2	31
54	An ELISA Assay for the Detection of Factor VIII Antibodies â€" Comparison with the Conventional Bethesda Assay in a Large Cohort of Haemophilia Samples. Acta Haematologica, 2003, 109, 18-22.	1.4	30

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55	Recent developments in drug resistance mechanism in chronic myeloid leukemia: a review. European Journal of Haematology, 2011, 87, 381-393.	2.2	30
56	Clinical and Autoimmune Profile of Scleroderma Patients from Western India. International Journal of Rheumatology, 2014, 2014, 1-6.	1.6	30
57	Sickle cell disease in India: A perspective. Indian Journal of Medical Research, 2016, 143, 21.	1.0	30
58	Impact of Î² globin gene mutations on the clinical phenotype of Î² thalassemia in India. Blood Cells, Molecules, and Diseases, 2004, 33, 153-157.	1.4	29
59	Clinical, hematologic and molecular variability of sickle cell-Î² thalassemia in western India. Indian Journal of Human Genetics, 2010, 16, 154.	0.7	29
60	Rapid Flow Cytometric Prenatal Diagnosis of Primary Immunodeficiency (PID) Disorders. Journal of Clinical Immunology, 2014, 34, 316-322.	3.8	29
61	Leukocyte Adhesion Deficiency-I with a Novel Intronic Mutation Presenting with Pyoderma Gangrenosum- Like Lesions. Journal of Clinical Immunology, 2015, 35, 431-434.	3.8	29
62	Association of clinical and serological parameters of systemic lupus erythematosus patients with Epsteinâ€Barr virus antibody profile. Journal of Medical Virology, 2018, 90, 559-563.	5.0	29
63	Non haematological effects of iron deficiency - A perspective. Indian Journal of Medical Sciences, 2006, 60, 30.	0.1	29
64	The spectrum of bleeding disorders in women with menorrhagia: a report from Western India. Annals of Hematology, 2005, 84, 339-342.	1.8	28
65	First-trimester prenatal diagnosis in haemophilia A and B familiesâ€”10 years experience from a centre in India. Prenatal Diagnosis, 2006, 26, 1015-1017.	2.3	28
66	Molecular and clinical heterogeneity in pyruvate kinase deficiency in India. Blood Cells, Molecules, and Diseases, 2013, 51, 133-137.	1.4	28
67	Evaluation of markers of endothelial damage in cases of young myocardial infarction. Atherosclerosis, 2005, 180, 375-380.	0.8	27
68	Ex vivo Expansion of Umbilical Cord Blood Stem Cells Using Different Combinations of Cytokines and Stromal Cells. Acta Haematologica, 2007, 118, 153-159.	1.4	27
69	Hemolytic anemia and distal renal tubular acidosis in two Indian patients homozygous for SLC4A1/AE1 mutation A858D. American Journal of Hematology, 2010, 85, 824-828.	4.1	27
70	Influence of CYP2C9 and VKORC1 gene polymorphisms on warfarin dosage, over anticoagulation and other adverse outcomes in Indian population. European Journal of Pharmacology, 2013, 710, 80-84.	3.5	27
71	Comparison of four commercially available activated partial thromboplastin time reagents using a semi-automated coagulometer. Blood Coagulation and Fibrinolysis, 2003, 14, 493-497.	1.0	26
72	Geography too determines the causes of inherited thrombophilia. Journal of Thrombosis and Haemostasis, 2004, 2, 363-364.	3.8	26

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73	Mutations in GPIIIa molecule as a cause for Glanzmann thrombasthenia in Indian patients. Journal of Thrombosis and Haemostasis, 2005, 3, 482-488.	3.8	26
74	Promising prognostic markers of Preeclampsia: New avenues in waiting. Thrombosis Research, 2015, 136, 189-195.	1.7	26
75	Genetic variations of hOCT1 gene and CYP3A4/A5 genes and their association with imatinib response in Chronic Myeloid Leukemia. European Journal of Pharmacology, 2015, 765, 124-130.	3.5	26
76	Diagnostic approach to primary immunodeficiency disorders. Indian Pediatrics, 2013, 50, 579-586.	0.4	25
77	Combined protein- and nucleic acid-level effects of rs1143679 (R77H), a lupus-predisposing variant within ITGAM. Human Molecular Genetics, 2014, 23, 4161-4176.	2.9	25
78	Combination of copeptin, placental growth factor and total annexin V microparticles for prediction of preeclampsia at 10â€“14 weeks of gestation. Placenta, 2017, 58, 67-73.	1.5	25
79	PTPN22 gene polymorphisms in autoimmune diseases with special reference to systemic lupus erythematosus disease susceptibility. Journal of Postgraduate Medicine, 2010, 56, 239-242.	0.4	25
80	Nanoimaging in cardiovascular diseases: Current state of the art. Indian Journal of Medical Research, 2015, 141, 285.	1.0	25
81	Mutations in theMCFD2 gene and a novel mutation in theLMAN1 gene in Indian families with combined deficiency of factor V and VIII. American Journal of Hematology, 2005, 79, 262-266.	4.1	24
82	Robustness of factor assays following cordocentesis in the prenatal diagnosis of haemophilia and other bleeding disorders. Haemophilia, 2007, 13, 172-177.	2.1	24
83	Evolution and selection of human leukocyte antigen alleles by Plasmodium falciparum infection. Human Immunology, 2008, 69, 856-860.	2.4	24
84	Menorrhagia and reproductive health in rare bleeding disorders: a study from the Indian subcontinent. Haemophilia, 2009, 15, 199-202.	2.1	24
85	Combined effects of the UGT1A1 and OATP2 gene polymorphisms as major risk factor for unconjugated hyperbilirubinemia in Indian neonates. Gene, 2014, 547, 18-22.	2.2	24
86	Genetic Variations in Bilirubin Metabolism Genes and Their Association with Unconjugated Hyperbilirubinemia in Adults. Annals of Human Genetics, 2017, 81, 11-19.	0.8	24
87	Molecular basis of weak D expression in the Indian population and report of a novel, predominant variant RHD allele. Transfusion, 2018, 58, 1540-1549.	1.6	24
88	Red cell distribution width and its association with mortality in neonatal sepsis. Journal of Maternal-Fetal and Neonatal Medicine, 2019, 32, 1925-1930.	1.5	24
89	Evolution of BCR/ABL Gene Mutation in CML Is Time Dependent and Dependent on the Pressure Exerted by Tyrosine Kinase Inhibitor. PLoS ONE, 2015, 10, e0114828.	2.5	24
90	Do high sensitivity C-reactive protein and serum interleukin-6 levels correlate with disease activity in systemic lupus erythematosuspatients?. Journal of Postgraduate Medicine, 2017, 63, 92-95.	0.4	24

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91	Immune regulatory gene polymorphisms as predisposing risk factors for the development of factor VIII inhibitors in Indian severe haemophilia A patients. Haemophilia, 2012, 18, 794-797.	2.1	23
92	Bone health in persons with haemophilia: a review. European Journal of Haematology, 2012, 89, 95-102.	2.2	23
93	Experimental animal model to study iron overload and iron chelation and review of other such models. Blood Cells, Molecules, and Diseases, 2015, 55, 194-199.	1.4	23
94	Chronic synovitis and HLA B27 in patients with severe haemophilia. Lancet, The, 2003, 361, 933-934.	13.7	22
95	Î²â€Globin Gene Cluster Haplotypes Linked to the Î²SGene in Western India. Hemoglobin, 2004, 28, 157-161.	0.8	22
96	Phenotypic and genotypic characterization of Factor VII deficiency patients from Western India. Clinica Chimica Acta, 2009, 409, 106-111.	1.1	22
97	Anti-nucleosome antibodies as a disease marker in systemic lupus erythematosus and its correlation with disease activity and other autoantibodies. Indian Journal of Dermatology, Venereology and Leprology, 2010, 76, 145.	0.6	22
98	X-linked hyper IgM syndrome: Clinical, immunological and molecular features in patients from India. Blood Cells, Molecules, and Diseases, 2014, 53, 99-104.	1.4	22
99	Neuropsychiatric manifestations and associated autoantibodies in systemic lupus erythematosus patients from Western India. Rheumatology International, 2015, 35, 541-545.	3.0	22
100	Glucose-6-phosphate dehydrogenase (G6PD) deficiency among tribal populations of India - Country scenario. Indian Journal of Medical Research, 2015, 141, 516-20.	1.0	22
101	Background noise of infection for using ANCA as a diagnostic tool for vasculitis in tropical and developing countries. Parasitology Research, 2008, 102, 1093-1095.	1.6	21
102	Comprehensive Report of Primary Immunodeficiency Disorders from a Tertiary Care Center in India. Journal of Clinical Immunology, 2013, 33, 507-512.	3.8	21
103	Nucleic acid amplification testing in Indian blood banks: A review with perspectives. Indian Journal of Pathology and Microbiology, 2017, 60, 313.	0.2	21
104	Fractures of long bones in severe haemophilia. Haemophilia, 2007, 13, 337-339.	2.1	20
105	Molecular characterization of leukocyte adhesion deficiency-I in Indian patients: Identification of 9 novel mutations. Blood Cells, Molecules, and Diseases, 2015, 54, 217-223.	1.4	20
106	Epidemiology of hepatocellular carcinoma (HCC) in hemophilia. Critical Reviews in Oncology/Hematology, 2016, 99, 129-133.	4.4	20
107	Haemoglobinopathies in tribal populations of India. Indian Journal of Medical Research, 2015, 141, 505-8.	1.0	20
108	Intron 22 Inversions in Factor VIII Gene in Indian Hemophiliacs. Thrombosis and Haemostasis, 1998, 79, 881-881.	3.4	19

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109	Correlation of thromboelastographic patterns with clinical presentation and rationale for use of antifibrinolytics in severe haemophilia patients. Haemophilia, 2007, 13, 734-739.	2.1	18
110	Thrombophilic dimension of recurrent fetal loss in Indian patients. Blood Coagulation and Fibrinolysis, 2008, 19, 581-584.	1.0	18
111	Association of (GT)n Repeats Promoter Polymorphism of Heme Oxygenase-1 Gene with Serum Bilirubin Levels in Healthy Indian Adults. Genetic Testing and Molecular Biomarkers, 2011, 15, 215-218.	0.7	18
112	First case of Hb Fontainebleau with sickle haemoglobin and other non-deletional Î± gene variants identified in neonates during newborn screening for sickle cell disorders. Journal of Clinical Pathology, 2012, 65, 654-659.	2.0	18
113	Spectrum of perforin gene mutations in familial hemophagocytic lymphohistiocytosis (FHL) patients in India. Blood Cells, Molecules, and Diseases, 2015, 54, 250-257.	1.4	18
114	Clinical spectrum and molecular basis of recessive congenital methemoglobinemia in India. Clinical Genetics, 2015, 87, 62-67.	2.0	18
115	Warfarin Dose Model for the Prediction of Stable Maintenance Dose in Indian Patients. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 353-359.	1.7	18
116	The phenotypic and molecular diversity of hemoglobinopathies in India: A review of 15Âyears at a referral center. International Journal of Laboratory Hematology, 2019, 41, 218-226.	1.3	18
117	Systemic Capillary Leak Syndrome Preceding Plasma Cell Leukaemia. Acta Haematologica, 2001, 106, 118-121.	1.4	17
118	Human platelet alloantigen polymorphism in Glanzmann's thrombasthenia and its impact on the severity of the disease. British Journal of Haematology, 2002, 119, 348-353.	2.5	17
119	Molecular Diversity of Hemoglobin H Disease in India. American Journal of Clinical Pathology, 2010, 133, 491-494.	0.7	17
120	HLA involvement in nevirapine-induced dermatological reaction in antiretroviral-treated HIV-1 patients. Journal of Pharmacology and Pharmacotherapeutics, 2011, 2, 114-115.	0.4	17
121	Homeopathic medicines substantially reduce the need for clotting factor concentrates in haemophilia patients: results of a blinded placebo controlled cross over trial. Homeopathy, 2012, 101, 38-43.	1.0	17
122	<scp>DNA</scp> interstrand crossâ€link repair: understanding role of <scp>F</scp>anconi anemia pathway and therapeutic implications. European Journal of Haematology, 2013, 91, 381-393.	2.2	17
123	Masking of a Î²-Thalassemia Determinant by a Novel Î´-Globin Gene Defect [Hb A₂-Saurashtra or Î´100(G2)Proâ†'Ser;HBD: c.301C>T] inCis. Hemoglobin, 2014, 38, 24-27.	0.8	17
124	Novel therapeutic approaches for haemophilia. Haemophilia, 2015, 21, 152-161.	2.1	17
125	Tyrosine kinase inhibitors: New class of antimalarials on the horizon?. Blood Cells, Molecules, and Diseases, 2015, 55, 119-126.	1.4	17
126	Investigation of Plasminogen Activator Inhibitorâ€1 (<scp>PAI</scp>â€1) 4G/5G promoter polymorphism in Indian venous thrombosis patients: A caseâ€control study. European Journal of Haematology, 2017, 99, 249-254.	2.2	17

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127	Differential role of Kruppel like factor 1 (KLF1) gene in red blood cell disorders. Genomics, 2019, 111, 1771-1776.	2.9	17
128	Use of the dual force system to correct chronic knee deformities due to severe haemophilia. Haemophilia, 2000, 6, 177-180.	2.1	16
129	High Frequency of Factor V Leiden Mutation in Parsis â€“ a Highly Endogamous Population in India. Thrombosis and Haemostasis, 2000, 83, 965-965.	3.4	16
130	Carrier detection and prenatal diagnosis in haemophilia in India: realities and challenges. Haemophilia, 2002, 8, 51-55.	2.1	16
131	HLA A*02 allele frequencies and B haplotype associations in Western Indians. Human Immunology, 2003, 64, 562-566.	2.4	16
132	Frequency of partial D in Western India. Transfusion Medicine, 2008, 18, 91-96.	1.1	16
133	Paroxysmal nocturnal haemoglobinuria: diagnostic tests, advantages, & limitations. European Journal of Haematology, 2009, 83, 503-511.	2.2	16
134	Diverse phenotypes and transfusion requirements due to interaction of Î²-thalassemias with triplicated Î±-globin genes. Annals of Hematology, 2015, 94, 1953-1958.	1.8	16
135	A functional SNP MCP-1 (â^'2518A/G) predispose to renal disorder in Indian Systemic Lupus Erythematosus patients. Cytokine, 2017, 96, 189-194.	3.2	16
136	Hydroxyurea Could Be a Good Clinically Relevant Iron Chelator. PLoS ONE, 2013, 8, e82928.	2.5	16
137	A Patient with Congenital Dyserythropoietic Anaemia Type III Presenting with Stillbirths. Acta Haematologica, 1998, 99, 31-33.	1.4	15
138	Feasibility of Antenatal Screening of Î²-Thalassemia in Mumbai, India. Acta Haematologica, 2001, 105, 252-252.	1.4	15
139	A novel R198H mutation in the glucose-6-phosphate dehydrogenase gene in the tribal groups of the Nilgiris in Southern India. Journal of Human Genetics, 2008, 53, 181-184.	2.3	15
140	A comprehensive screening analysis of antiphospholipid antibodies in Indian women with fetal loss. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2008, 137, 136-140.	1.1	15
141	Spectrum of novel mutations in the human PKLR gene in pyruvate kinaseâ€deficient Indian patients with heterogeneous clinical phenotypes. Clinical Genetics, 2009, 75, 157-162.	2.0	15
142	Severe mental retardation and recessive congenital methemoglobinemia in three Indian patients: Compound heterozygous for NADHâ€cytochrome b5 reductase gene mutations. American Journal of Hematology, 2011, 86, 327-329.	4.1	15
143	Epidemiology, Diagnosis, and Management of<scp>v</scp>on Willebrand Disease in India. Seminars in Thrombosis and Hemostasis, 2011, 37, 595-601.	2.7	15
144	Prenatal Diagnosis of LAD-I on Cord Blood by Flowcytometry. Indian Journal of Pediatrics, 2012, 79, 1605-1609.	0.8	15

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145	The Epidemiology of FVIII Inhibitors in Indian Haemophilia A Patients. Indian Journal of Hematology and Blood Transfusion, 2014, 30, 356-363.	0.6	15
146	Clinical and hematological presentation among Indian patients with common hemoglobin variants. Clinica Chimica Acta, 2014, 431, 46-51.	1.1	15
147	Variable phenotypes of sickle cell disease in <scp>I</scp>ndia with the <scp>A</scp>rabâ€«<scp>I</scp>ndian haplotype. British Journal of Haematology, 2015, 168, 156-159.	2.5	15
148	Does HbF induction by hydroxycarbamide work through <scp>MIR</scp>210 in sickle cell anaemia patients?. British Journal of Haematology, 2016, 173, 801-803.	2.5	15
149	Differential antigen expression and aberrant signaling via PI3/AKT, MAP/ERK, JAK/STAT, and Wnt/Î² catenin pathways in Linâ^'/CD38â^'/CD34+ cells in acute myeloid leukemia. European Journal of Haematology, 2016, 96, 309-317.	2.2	15
150	Microparticles as prognostic biomarkers in dengue virus infection. Acta Tropica, 2018, 181, 21-24.	2.0	15
151	FANCA Gene Mutations with 8 Novel Molecular Changes in Indian Fanconi Anemia Patients. PLoS ONE, 2016, 11, e0147016.	2.5	15
152	Common human leucocyte antigen haplotypes in Indians ? its implications in finding unrelated compatible bone marrow donors. Transfusion Medicine, 2002, 12, 43-48.	1.1	14
153	Haematuria and urolithiasis in patients with haemophilia. European Journal of Haematology, 2003, 70, 410-412.	2.2	14
154	Congenital methemoglobinemia caused by Hb-MRatnagiri (Î²-63CATâ†'TAT, Hisâ†'Tyr) in an Indian family. American Journal of Hematology, 2005, 79, 168-170.	4.1	14
155	Second Trimester Antenatal Diagnosis in Rare Coagulation Factor Deficiencies. Journal of Pediatric Hematology/Oncology, 2007, 29, 137-139.	0.6	14
156	G6PD Namoru (208 T? C) is the major polymorphic variant in the tribal populations in southern India. British Journal of Haematology, 2007, 136, 512-513.	2.5	14
157	Successful Pregnancy Outcome in Women With Bad Obstetric History and Recurrent Fetal Loss Due to Thrombophilia: Effect of Unfractionated Heparin and Lowâ€"Molecular Weight Heparin. Clinical and Applied Thrombosis/Hemostasis, 2008, 14, 174-179.	1.7	14
158	Clinical, genetic and cytogenetic study of Fanconi anemia in an Indian population. Hematology, 2010, 15, 58-62.	1.5	14
159	Five Î± globin chain variants identified during screening for haemoglobinopathies. European Journal of Clinical Investigation, 2010, 40, 226-232.	3.4	14
160	Immunological disturbances associated with malarial infection. Journal of Parasitic Diseases, 2013, 37, 11-15.	1.0	14
161	Genetic Heterogeneity in a Large Cohort of Indian Type 3 von Willebrand Disease Patients. PLoS ONE, 2014, 9, e92575.	2.5	14
162	Molecular Pathology of Rare Bleeding Disorders (RBDs) in India: A Systematic Review. PLoS ONE, 2014, 9, e108683.	2.5	14

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163	Impact of TNF-Î± and LTÎ± gene polymorphisms on genetic susceptibility in Indian SLE patients. Human Immunology, 2017, 78, 201-208.	2.4	14
164	Predisposition of IL-1Î² (-511 C/T) polymorphism to renal and hematologic disorders in Indian SLE patients. Gene, 2018, 641, 41-45.	2.2	14
165	Investigating Cell Surface Markers on Normal Hematopoietic Stem Cells in Three Different Niche Conditions. International Journal of Stem Cells, 2013, 6, 129-133.	1.8	14
166	Fasting plasma homocysteine levels are increased in young patients with acute myocardial infarction from Western India. Indian Heart Journal, 2007, 59, 242-5.	0.5	14
167	Carrier detection in haemophilia A families: comparison of conventional coagulation parameters with DNA polymorphism analysis - first report from India. Haemophilia, 1999, 5, 243-246.	2.1	13
168	Clinical Diversity of Sickle Cell Disease in Western India â€“ Influence of Genetic Factors. Acta Haematologica, 2000, 103, 122-123.	1.4	13
169	Inversion of intron is a rare cause of severe hemophilia A in Indian population. Journal of Thrombosis and Haemostasis, 2004, 2, 1481-1482.	3.8	13
170	Frequency distribution of human platelet antigens in the Indian population. Transfusion Medicine, 2005, 15, 119-124.	1.1	13
171	Evaluation of the Use of Monoclonal Antibodies and Nested PCR for Noninvasive Prenatal Diagnosis of Hemoglobinopathies in India. American Journal of Clinical Pathology, 2008, 130, 202-209.	0.7	13
172	Novel mutations in GP IIb gene in Glanzmann's thrombasthenia from India. Platelets, 2009, 20, 35-40.	2.3	13
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