Anil Pathare

List of Publications by Year
in descending order

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108	1,673	331259	360668
papers	citations	21 h-index	35 g-index

109	109	109	2047
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Cytokine profile of sickle cell disease in Oman. American Journal of Hematology, 2004, 77, 323-328.	2.0	123
2	Predictive Value of Neutrophil to Lymphocyte Ratio in Outcomes of Patients with Acute Coronary Syndrome. Archives of Medical Research, 2010, 41, 618-622.	1.5	81
3	Cytokines in Sickle Cell Disease. Hematology, 2003, 8, 329-337.	0.7	74
4	Combined therapy with desferrioxamine and deferiprone in beta thalassemia major patients with transfusional iron overload. Annals of Hematology, 2006, 85, 315-319.	0.8	70
5	Reduction in labile plasma iron during treatment with deferasirox, a onceâ€daily oral iron chelator, in heavily ironâ€overloaded patients with Î²â€thalassaemia. European Journal of Haematology, 2009, 82, 454-457.	1.1	70
6	Forecasting Hemoglobinopathy Burden Through Neonatal Screening in Omani Neonates. Hemoglobin, 2010, 34, 135-144.	0.4	64
7	Venous Thromboembolism in Young Patients From Western India: A Study. Clinical and Applied Thrombosis/Hemostasis, 2001, 7, 158-165.	0.7	62
8	Deferasirox (ExjadeÂ®) significantly improves cardiac T2* in heavily iron-overloaded patients with Î²-thalassemia major. Annals of Hematology, 2010, 89, 405-409.	0.8	53
9	Imatinib in pregnancy. European Journal of Haematology, 2005, 74, 535-537.	1.1	45
10	Orbital Infarction in Sickle Cell Disease. American Journal of Ophthalmology, 2008, 146, 595-601.e1.	1.7	43
11	Factor VIII and IX gene polymorphisms and carrier analysis in Indian population. , 1997, 54, 271-275.		37
12	Dose-ranging studies on liposomal amphotericin B (L-AMP-LRC-1) in the treatment of visceral leishmaniasis. Transactions of the Royal Society of Tropical Medicine and Hygiene, 1999, 93, 314-318.	0.7	35
13	Rifampicin-induced immune thrombocytopenia. Tubercle and Lung Disease, 1996, 77, 558-562.	2.1	33
14	Combined factor V and VIII deficiency in Indian population. Haemophilia, 2000, 6, 504-507.	1.0	30
15	The spectrum of bleeding disorders in women with menorrhagia: a report from Western India. Annals of Hematology, 2005, 84, 339-342.	0.8	28
16	Complications of PORT-A-CATHÂ® in patients with sickle cell disease. Journal of Infection and Public Health, 2012, 5, 57-62.	1.9	28
17	dRTA and hemolytic anemia: first detailed description of SLC4A1 A858D mutation in homozygous state. European Journal of Haematology, 2012, 88, 350-355.	1.1	26
18	Warfarin pharmacogenetics: development of a dosing algorithm for Omani patients. Journal of Human Genetics, 2012, 57, 665-669.	1.1	25

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19	Prevalence of methicillin resistant Staphylococcus aureus [MRSA] colonization or carriage among health-care workers. Journal of Infection and Public Health, 2016, 9, 571-576.	1.9	25
20	Complex t(8;13;21)(q22;q14;q22)â€“A Novel Variant of t(8;21) in a Patient with Acute Myeloid Leukemia (AMLâ€“M2). Archives of Medical Research, 2008, 39, 252-256.	1.5	23
21	Clinical and molecular findings of chronic granulomatous disease in Oman: family studies. Clinical Genetics, 2015, 87, 185-189.	1.0	23
22	Effect of <scp>CYP</scp>4F2,<scp>VKORC</scp>1, and <scp>CYP</scp>2C9 in Influencing Coumarin Dose: A Singleâ€Patient Data Metaâ€Analysis in More Than 15,000 Individuals. Clinical Pharmacology and Therapeutics, 2019, 105, 1477-1491.	2.3	23
23	T2* cardiovascular magnetic resonance in the management of thalassemia patients in Oman. Haematologica, 2009, 94, 140-141.	1.7	22
24	Intron 22 Inversions in Factor VIII Gene in Indian Hemophiliacs. Thrombosis and Haemostasis, 1998, 79, 881-881.	1.8	19
25	A stepwise Î±-thalassemia screening strategy in high-prevalence areas. European Journal of Haematology, 2013, 91, 164-169.	1.1	19
26	Bleeding score in Type 1 von Willebrand disease patients using the <scp>ISTH</scp>â€<scp>BAT</scp> questionnaire. International Journal of Laboratory Hematology, 2018, 40, 175-180.	0.7	19
27	Hereditary thrombophilia in ethnic omani patients. American Journal of Hematology, 2006, 81, 101-106.	2.0	18
28	Pattern and outcome of vascular involvement of Omani patients with Behcetâ€™s disease. Rheumatology International, 2011, 31, 731-735.	1.5	18
29	Sensorineural hearing loss in sickle cell diseaseâ€”A prospective study from Oman. Laryngoscope, 2011, 121, 392-396.	1.1	18
30	Systemic Capillary Leak Syndrome Preceding Plasma Cell Leukaemia. Acta Haematologica, 2001, 106, 118-121.	0.7	17
31	Arterial and Venous Thrombotic Complications with Thalidomide in Multiple Myeloma. Archives of Medical Research, 2008, 39, 257-258.	1.5	17
32	Necrotizing Enterocolitis in a Term Neonate Following Intravenous Immunoglobulin Therapy. Indian Journal of Pediatrics, 2011, 78, 743-744.	0.3	17
33	Use of the dual force system to correct chronic knee deformities due to severe haemophilia. Haemophilia, 2000, 6, 177-180.	1.0	16
34	Cytogenetic, morphological, and immunophenotypic patterns in Omani patients with de novo acute myeloid leukemia. Cancer Genetics and Cytogenetics, 2007, 177, 89-94.	1.0	16
35	Warfarin Pharmacogenetics: Polymorphisms of theCYP2C9, CYP4F2, and VKORC1Loci in a Genetically Admixed Omani Population. Human Biology, 2012, 84, 67-77.	0.4	16
36	ALLOIMMUNIZATION IN PATIENTS WITH SICKLE CELL DISEASE AND THALASSAEMIA: EXPERIENCE OF SINGLE CENTRE FROM OMAN. Mediterranean Journal of Hematology and Infectious Diseases, 2016, 9, e2017013.	0.5	16

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37	A Patient with Congenital Dyserythropoietic Anaemia Type III Presenting with Stillbirths. Acta Haematologica, 1998, 99, 31-33.	0.7	15
38	Novel spectrum of perforin gene mutations in familial hemophagocytic lymphohistiocytosis in ethnic omani patients. American Journal of Hematology, 2007, 82, 1099-1102.	2.0	15
39	Predictors of impending acute chest syndrome in patients with sickle cell anaemia. Scientific Reports, 2020, 10, 2470.	1.6	15
40	Perforin A91V polymorphism and putative susceptibility to hematological malignancies. Leukemia, 2006, 20, 2178-2178.	3.3	14
41	Hematopoietic stem cell transplantation in Oman. Bone Marrow Transplantation, 2008, 42, S109-S113.	1.3	14
42	Carrier detection in haemophilia A families: comparison of conventional coagulation parameters with DNA polymorphism analysis - first report from India. Haemophilia, 1999, 5, 243-246.	1.0	13
43	Clinical and Genetic Studies of Familial Hemophagocytic Lymphohistiocytosis in Oman: Need for Early Treatment. Pediatric Hematology and Oncology, 2003, 20, 603-609.	0.3	13
44	Role of antiidiotypic antibodies on the clinical course of idiopathic thrombocytopenic purpura. Translational Research, 2003, 142, 113-120.	2.4	12
45	Postvaricella Thrombosisâ€"Report of Two Cases and Literature Review. Pediatric Infectious Disease Journal, 2012, 31, 985-987.	1.1	12
46	Red Cell Alloimmunization to Rhesus Antigen Among Pregnant Women Attending a Tertiary Care Hospital in Oman. Oman Medical Journal, 2016, 31, 73-76.	0.3	12
47	Spectrum of inherited bleeding disorders from Western India. Haematologia, 2002, 32, 39-47.	0.2	11
48	COMPARISON OF METHICILLIN RESISTANT STAPHYLOCOCCUS AUREUS IN HEALTHY COMMUNITY HOSPITAL VISITORS[CA-MRSA] AND HOSPITAL STAFF [HA-MRSA]. Mediterranean Journal of Hematology and Infectious Diseases, 2015, 7, e2015053.	0.5	10
49	Molecular characterization of haemophilia A & B in Indians. Haemophilia, 1998, 4, 802-805.	1.0	9
50	Cytogenetic Profile of Childhood Acute Lymphoblastic Leukemia in Oman. Archives of Medical Research, 2007, 38, 305-312.	1.5	9
51	Comparative Assessment of Deferiprone and Deferasirox in Thalassemia Major Patients in the First Two Decades-Single Centre Experience. Pediatric Hematology and Oncology, 2013, 30, 104-112.	0.3	9
52	SERUM TOTAL BILIRUBIN, NOT CHOLELITHIASIS, IS INFLUENCED BY UGT1A1 POLYMORPHISM, ALPHA THALASSEMIA AND ï¢S GENOTYPE: FIRST REPORT ON COMPARISON BETWEEN ARAB-INDIAN AND AFRICAN ï¢S GENES. Mediterranean Journal of Hematology and Infectious Diseases, 2015, 7, e2015060.	0.5	9
53	Aetiological profile of women presenting with premature ovarian failure to a single tertiary care center in Oman. Post Reproductive Health, 2015, 21, 63-68.	0.3	9
54	Epsilon-Aminocaproic Acid Inhibits the Activity of Factor VIII Inhibitors in Patients with Severe Haemophilia A in vivo and in vitro. Acta Haematologica, 2000, 103, 67-72.	0.7	8

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55	Clinical and laboratory characteristics of patients with sickle-cell and autoimmune/connective tissue diseases. Rheumatology International, 2012, 32, 373-378.	1.5	8
56	Predictive Value of Red Cell Indices and High Performance Liquid Chromatography in the Diagnosis of Haemoglobinopathies: Interim Results of the National Neonatal Cord Blood Screening Programme.. Blood, 2006, 108, 3352-3352.	0.6	8
57	Heterozygous methylene tetrahydrofolate reductase mutation with mild hyperhomocysteinemia associated with deep vein thrombosis. International Journal of Laboratory Hematology, 2004, 26, 143-146.	0.2	7
58	Successful outcome with anagrelide in pregnancy. Annals of Hematology, 2005, 84, 758-759.	0.8	7
59	Complete recovery following sudden sensorineural hearing loss in a patient with sickle cell disease. Hematology/ Oncology and Stem Cell Therapy, 2011, 4, 97-99.	0.6	7
60	Transcranial Doppler Ultrasonography in Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2013, 35, 18-23.	0.3	7
61	First report of the spectrum of Î´-globin gene mutations in Omani subjects - identification of novel mutations. International Journal of Laboratory Hematology, 2015, 37, 238-243.	0.7	7
62	Respiratory Viral Infections in Sickle Cell Anemia: Special Emphasis on H1N1 Co-infection. Oman Medical Journal, 2020, 35, e197-e197.	0.3	7
63	Neonatal Screening: Mean haemoglobin and red cell indices in cord blood from Omani neonates. Sultan Qaboos University Medical Journal, 2011, 11, 462-9.	0.3	7
64	Recurrent priapism in sickle cell trait with protein S deficiency. JPMA the Journal of the Pakistan Medical Association, 2008, 58, 701-2.	0.1	7
65	Clinically Significant Inhibitors in Hemophilia A Patients from India Tend to Persist. Acta Haematologica, 2000, 103, 175-176.	0.7	6
66	Von Willebrand Factor 1 and Factor 2 Alleles (Intron 40) Are Suitable Markers for Carrier Detection in von Willebrand Disease Families in the Indian Population. Acta Haematologica, 2006, 115, 64-67.	0.7	6
67	Follicular dendritic cell hyperplasia in plasma cell variant of Castleman's disease with interfollicular Hodgkin's disease. Pathology Research and Practice, 2007, 203, 479-484.	1.0	6
68	Haematological and clinical features of Î²â€‹thalassaemia associated with Hb Dhofar. European Journal of Haematology, 2008, 80, 67-70.	1.1	6
69	Hydroxyurea or Chronic Exchange Transfusions in Patients With Sickle Cell Disease. Journal of Pediatric Hematology/Oncology, 2009, 31, 42-44.	0.3	6
70	Predicting risk factors for thromboembolic complications in patients with sickle cell anaemia â€“ lessons learned for prophylaxis. Journal of International Medical Research, 2021, 49, 030006052110553.	0.4	6
71	Low cost autologous peripheral blood stem cell transplantation performed in a municipal hospital for a patient with plasma cell leukaemia. International Journal of Laboratory Hematology, 2002, 24, 187-190.	0.2	5
72	CD45 Gene C77G Mutation in Haemophagocytic Lymphohistiocytosis. Acta Haematologica, 2007, 118, 160-161.	0.7	5

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73	Trisomy 21 as a Sole Acquired Abnormality in an Adult Omani Patient with CD7- and CD9-Positive Acute Myeloid Leukemia. Archives of Medical Research, 2007, 38, 797-802.	1.5	5
74	Coexistence of immune thrombocytopenic purpura and idiopathic membranous glomerulonephritis successfully treated with rituximab. Platelets, 2010, 21, 575-577.	1.1	5
75	PREVALENCE OF HEPATITIS B, HEPATITIS C AND HIV IN MULTIPLY TRANSFUSED SICKLE CELL DISEASE PATIENTS FROM OMAN. Mediterranean Journal of Hematology and Infectious Diseases, 2019, 11, e2019058.	0.5	5
76	Fresh frozen plasma (Octaplas) and topical heparin in the management of ligneous conjunctivitis. Journal of AAPOS, 2019, 23, 42-45.e1.	0.2	5
77	Disseminated strongyloidiasis and cytomegalovirus infection in a patient with anaplastic large cell lymphoma. Annals of Hematology, 2007, 86, 925-926.	0.8	4
78	Protein Z levels in pregnant Omani women: correlation with pregnancy outcome. Journal of Thrombosis and Thrombolysis, 2011, 32, 453-458.	1.0	4
79	Rituximab Leads to Long Remissions in Patients with Chronic Immune Thrombocytopenia. Oman Medical Journal, 2015, 30, 111-114.	0.3	4
80	INCREASED VASOOCCLUSIVE CRISIS IN â€œOâ€•BLOOD GROUP SICKLE CELL DISEASE PATIENTS: ASSOCIATION WITH UNDERLYING THROMBOSPONDIN LEVELS.. Mediterranean Journal of Hematology and Infectious Diseases, 2016, 9, e2017028.	0.5	4
81	Bleeding score in type 1 von Willebrand disease patients using the condensed MCMDMâ€¡ vWD validated questionnaire. International Journal of Laboratory Hematology, 2018, 40, 515-520.	0.7	4
82	Clinical and laboratory parameters, risk factors predisposing to the development of priapism in sickle cell patients. Experimental Biology and Medicine, 2020, 245, 79-83.	1.1	4
83	A Novel Deletional Î²-Thalassemic Variant in an Ethnic Qatari Patient. Hemoglobin, 2009, 33, 214-219.	0.4	3
84	IMPACT OF MANNOSE-BINDING PROTEIN GENE POLYMORPHISMS IN OMANI SICKLE CELL DISEASE PATIENTS. Mediterranean Journal of Hematology and Infectious Diseases, 2016, 8, 2016013.	0.5	3
85	THE USE OF HPLC AS A TOOL FOR NEONATAL CORD BLOOD SCREENING OF HAEMOGLOBINOPATHY - A VALIDATION STUDY. Mediterranean Journal of Hematology and Infectious Diseases, 2019, 11, e2019005.	0.5	3
86	Permanent Alopecia in Children Following Busulfan Based Conditioning Is Associated with Glutathione M1 Null Genotype.. Blood, 2005, 106, 2740-2740.	0.6	3
87	Haemostatic Parameters in Patients with BehÃ§et's Disease. Sultan Qaboos University Medical Journal, 2014, 14, e190-6.	0.3	3
88	Iron Overload in Patients With Heavily Transfused Sickle Cell Diseaseâ€"Correlation of Serum Ferritin With Cardiac T2* MRI (CMRTools), Liver T2* MRI, and R2-MRI (FerriscanÂ®). Frontiers in Medicine, 2021, 8, 731102.	1.2	2
89	Isochromosome 9q as a sole anomaly in an Omani boy with acute lymphoblastic leukaemia. BMJ Case Reports, 2009, 2009, bcr0920080890-bcr0920080890.	0.2	2
90	Heterozygous methylene tetrahydrofolate reductase mutation with mild hyperhomocysteinemia associated with deep vein thrombosis. Haematologia, 2002, 32, 551-6.	0.2	2

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91	Gas gangrene in a patient with severe haemophilia A. Haemophilia, 1999, 5, 450-452.	1.0	1
92	Biphenotypic leukemia with interstitial del(9)(q22q32) as a sole abnormality. Cancer Genetics and Cytogenetics, 2007, 178, 170-172.	1.0	1
93	Acute massive splenic infarction with complete liquefaction of the spleen in sickle cell disease. Hematology/ Oncology and Stem Cell Therapy, 2011, 4, 142-144.	0.6	1
94	A profile of rare bloods in Oman. Asian Journal of Transfusion Science, 2013, 7, 162.	0.1	1
95	Association between cardiac T2* magnetic resonance imaging values and endocrine function tests in patients with Î²-thalassemia major. Blood Cells, Molecules, and Diseases, 2014, 52, 50-51.	0.6	1
96	Suboptimal Busulphan Exposure Is Associated with Mixed Hematopoietic Chimerism, a Risk Factor for Rejection in Bone Marrow Transplantation for Homozygous beta Thalassemia.. Blood, 2004, 104, 1149-1149.	0.6	1
97	Long Term Follow up of Patients with Immune Thrombocytopenia Receiving Rituximab. Blood, 2011, 118, 4675-4675.	0.6	1
98	Combined effect of <i>CYP2C9</i> and <i>VKORC1</i> polymorphisms on warfarin maintenance dose in Omani patients. Open Journal of Genetics, 2012, 02, 184-189.	0.1	1
99	Novel PKLR missense mutation (A300P) causing pyruvate kinase deficiency in an Omani Kindredâ€"PK deficiency masquerading as congenital dyserythropoietic anemia. Clinical Case Reports (discontinued), 2022, 10, e05315.	0.2	1
100	Prenatal diagnosis of haemophilia: a preliminary report. The National Medical Journal of India, 1998, 11, 218-9.	0.1	1
101	Novel human pathological mutations. Gene symbol: PROC. Disease: Protein C deficiency. Human Genetics, 2009, 126, 336-7.	1.8	1
102	Clinical and Genetic Studies of Familial Hemophagocytic Lymphohistiocytosis in Oman: Need for Early Treatment. Pediatric Hematology and Oncology, 2003, 20, 603-609.	0.3	1
103	Orbital infarction in sickle cell disease: A clinicogenetic study from Oman. Journal of AAPOS, 2007, 11, 88-89.	0.2	0
104	PO12-TU-02 Transcranial Doppler ultrasound in sickle cell disease: an experience from Oman. Journal of the Neurological Sciences, 2009, 285, S222-S223.	0.3	0
105	A novel p.Pro353His SERPINC1 mutation in the thrombinâ€binding region affecting stability of Antithrombin molecule in an extended Omani family. International Journal of Laboratory Hematology, 2018, 40, e49-e51.	0.7	0
106	Newborn Haemoglobinopathy Carrier Screening: Towards Comprehensive and Improved Patient Care of Sickle Cell Disease in Oman, a Cost Effective Analysis.. Blood, 2007, 110, 5155-5155.	0.6	0
107	Inherited Thrombophilia In Omani Women with Recurrent Pregnancy Loss. Blood, 2010, 116, 5124-5124.	0.6	0
108	Pulmonary Hypertension in Sickle Cell Disease Patients: Correlation of TRV Jet with Serum NT-Pro BNP Concentration. National Journal of Health Sciences, 2016, 1, 5-10.	0.1	0