Can Boga

List of Publications by Year
in descending order

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Version: 2024-02-01

85	736	471061	610482
papers	citations	17 h-index	24 g-index

85	85	85	1037
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Pregnancy and sickle cell disease: A review of the current literature. Critical Reviews in Oncology/Hematology, 2016, 98, 364-374.	2.0	63
2	Prophylactic red blood cell exchange may be beneficial in the management of sickle cell disease in pregnancy. Transfusion, 2015, 55, 36-44.	0.8	42
3	Human bone marrow mesenchymal cells express NG2: possible increase in discriminative ability of flow cytometry during mesenchymal stromal cell identification. Cytotherapy, 2009, 11, 527-533.	0.3	41
4	Clinical significance of circulating blood and endothelial cell microparticles in sickle-cell disease. Journal of Thrombosis and Thrombolysis, 2014, 38, 167-175.	1.0	36
5	Flow cytometric evaluation of circulating endothelial cells: A new protocol for identifying endothelial cells at several stages of differentiation. American Journal of Hematology, 2007, 82, 706-711.	2.0	35
6	Pulmonary Function and Airway Hyperresponsiveness in Adults with Sickle Cell Disease. Lung, 2009, 187, 195-200.	1.4	34
7	Therapeutic Potential of Apigenin, a Plant Flavonoid, for Imatinib-Sensitive and Resistant Chronic Myeloid Leukemia Cells. Nutrition and Cancer, 2014, 66, 599-612.	0.9	28
8	Allogenic peripheral stem cell transplantation from HLA-matched related donors for adult sickle cell disease: remarkable outcomes from a single-center trial. Bone Marrow Transplantation, 2018, 53, 880-890.	1.3	28
9	Lemierre syndrome variant: Staphylococcus aureus associated with thrombosis of both the right internal jugular vein and the splenic vein after the exploration of a river cave. Journal of Thrombosis and Thrombolysis, 2007, 23, 151-154.	1.0	23
10	East Mediterranean region sickle cell disease mortality trial: retrospective multicenter cohort analysis of 735 patients. Annals of Hematology, 2016, 95, 993-1000.	0.8	23
11	Conventional and molecular cytogenetic findings of myelodysplastic syndrome patients. Clinical and Experimental Medicine, 2005, 5, 55-59.	1.9	21
12	Dental and periodontal health status of subjects with sickle cell disease. Journal of Dental Sciences, 2011, 6, 227-234.	1.2	21
13	Short-term central venous catheter complications in patients with sickle cell disease who undergo apheresis. Journal of Thrombosis and Thrombolysis, 2014, 37, 97-101.	1.0	20
14	Fatal cardiac tamponade in a patient with Kawasaki disease. Heart and Lung: Journal of Acute and Critical Care, 2005, 34, 257-259.	0.8	19
15	Plasma exchange in critically ill patients with sickle cell disease. Transfusion and Apheresis Science, 2007, 37, 17-22.	0.5	19
16	Automated red cell exchange procedures in patients with sickle cell disease. Transfusion and Apheresis Science, 2007, 36, 305-312.	0.5	19
17	Hematopoietic Stem Cell Transplantation in Adult Sickle Cell Disease: Problems and Solutions. Turkish Journal of Haematology, 2015, 32, 195-205.	0.2	17
18	Comparison of the clinical course of COVID-19 infection in sickle cell disease patients with healthcare professionals. Annals of Hematology, 2021, 100, 2195-2202.	0.8	15

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19	Long-term colchicine therapy in a patient with BehÃ§etâ€™s disease and acute promyelocytic leukemia. Rheumatology International, 2007, 27, 763-765.	1.5	14
20	Gonadal brucellar abscess: Imaging and clinical findings in 3 cases and review of the literature. Journal of Clinical Ultrasound, 2007, 35, 395-400.	0.4	13
21	A case of disseminated intravascular coagulation caused by Brucella melitensis. Journal of Thrombosis and Thrombolysis, 2008, 26, 71-73.	1.0	11
22	Corticosteroid-induced vaso-occlusive events may be prevented by lowering hemoglobin S levels in adults with sickle cell disease. Transfusion and Apheresis Science, 2017, 56, 717-718.	0.5	11
23	ANORECTAL COMPLICATIONS DURING NEUTROPENIC PERIOD IN PATIENTS WITH HEMATOLOGIC DISEASES. Mediterranean Journal of Hematology and Infectious Diseases, 2016, 8, 2016019.	0.5	10
24	Assessment of corrected QT interval in sickle cell disease patients who undergo erythroapheresis. Transfusion Medicine, 2007, 17, 466-472.	0.5	8
25	The association of pagophagia with Helicobacter pylori infection in patients with iron-deficiency anemia. International Journal of Hematology, 2009, 90, 28-32.	0.7	8
26	Alterations of circulating endothelial cells after apheresis in patients with sickle cell disease: A potential clue for restoration of pathophysiology. Transfusion and Apheresis Science, 2010, 43, 273-279.	0.5	8
27	Plasmaâ€exchange treatment for severe carbamazepine intoxication: A case study. Journal of Clinical Apheresis, 2014, 29, 178-180.	0.7	8
28	Tunnelled Central Venous Catheter-Related Problems in the Early Phase of Haematopoietic Stem Cell Transplantation and Effects on Transplant Outcome. Turkish Journal of Haematology, 2015, 32, 51-57.	0.2	8
29	Effectiveness of Visual Methods in Information Procedures for Stem Cell Recipients and Donors. Turkish Journal of Haematology, 2017, 34, 321-327.	0.2	8
30	A dramatic response to rituximab in a patient with resistant thrombotic thrombocytopenic purpura (TTP) who developed acute stroke. Journal of Thrombosis and Thrombolysis, 2007, 23, 147-150.	1.0	7
31	Safety, therapeutic effectiveness, and cost of parenteral iron therapy. International Journal of Hematology, 2009, 90, 24-27.	0.7	7
32	A Comparison of the BEAM and MITO/MEL Conditioning Regimens for Autologous Hematopoietic Stem Cell Transplantation in Hodgkin Lymphoma: An Analysis of Efficiency and Treatment-Related Toxicity. Clinical Lymphoma, Myeloma and Leukemia, 2020, 20, 652-660.	0.2	7
33	The apoptosis of blood polymorphonuclear leukocytes in sickle cell disease. Cytometry Part B - Clinical Cytometry, 2007, 72B, 276-280.	0.7	6
34	Cobalamin Deficiency Can Mask Depleted Body Iron Reserves. Indian Journal of Hematology and Blood Transfusion, 2015, 31, 255-258.	0.3	6
35	Organ damage mitigation with the Baskent Sickle Cell Medical Care Development Program (BASCARE). Medicine (United States), 2018, 97, e9844.	0.4	6
36	Recommendations for Risk Categorization and Prophylaxis of Invasive Fungal Diseases in Hematological Malignancies: A Critical Review of Evidence and Expert Opinion (TEO-4). Turkish Journal of Haematology, 2015, 32, 100-117.	0.2	6

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37	A Rare Complication Developing After Hematopoietic Stem Cell Transplantation: Wernickeâ€™s Encephalopathy. Turkish Journal of Haematology, 2016, 32, 367-370.	0.2	6
38	Effect of Hereditary Hemochromatosis Gene H63D and C282Y Mutations on Iron Overload in Sickle Cell Disease Patients. Turkish Journal of Haematology, 2016, 33, 320-325.	0.2	6
39	Cytogenetic findings and clinical outcomes of adult acute myeloid leukaemia patients. Clinical and Experimental Medicine, 2007, 7, 102-107.	1.9	5
40	Use of Mesenchymal Cells to Modulate Immune Suppression and Immune Reconstruction in a Patient with Aplastic Anemia Complicated by Invasive Sino-Orbital Aspergillosis. Turkish Journal of Haematology, 2014, 31, 180-183.	0.2	5
41	Quantum cell expansion system: Safe and rapid expansion. Cytotherapy, 2017, 19, 1246-1247.	0.3	4
42	Granulocyteâ€colony stimulating factor administration among hemoglobin S trait donors: A single center experience from the Eastern Mediterranean region. Journal of Clinical Apheresis, 2018, 33, 65-71.	0.7	4
43	Excellent outcomes of allogeneic transplantation from peripheral blood of HLA-matched related donors for adult sickle cell disease with ATLG and posttransplant cyclophosphamide-containing regimen: an update work. Bone Marrow Transplantation, 2020, 55, 1647-1651.	1.3	4
44	Tacrolimus-Induced Diabetic Ketoacidosis After Allogeneic Bone Marrow Transplant. Experimental and Clinical Transplantation, 2017, 15, 702-703.	0.2	4
45	Serum cancer antigen 15-3 concentrations in patients with sickle cell disease. British Journal of Haematology, 2006, 134, 546-547.	1.2	3
46	Platelet P-selectin Expression in Patients With Sickle Cell Disease Who Undergo Apheresis. Therapeutic Apheresis and Dialysis, 2007, 11, 255-261.	0.4	3
47	Nitric oxide in gingival crevicular fluid and nitric oxide synthase expression in the gingiva of patients with sickle cell disease. Turkish Journal of Haematology, 2011, 28, 115-124.	0.2	3
48	JACIE accreditation from the perspective of an accredited centre. Bone Marrow Transplantation, 2017, 52, 1352-1352.	1.3	3
49	QTc prolongation during peripheral stem cell apheresis in healthy volunteers. Journal of Clinical Apheresis, 2017, 32, 240-245.	0.7	3
50	An Unusual Giant Leg Ulcer as a Rare Presentation of Sweetâ€™s syndrome in a Patient with Hairy Cell Leukemia Succesfully Managed by Splenectomy. Turkish Journal of Haematology, 2017, 34, 270-271.	0.2	3
51	Use of Plerixafor to Mobilize a Healthy Donor Infected with Influenza A. Turkish Journal of Haematology, 2018, 35, 138-139.	0.2	3
52	A detachment technique based on the thermophysiologic responses of cultured mesenchymal cells exposed to cold. Cytotherapy, 2008, 10, 686-689.	0.3	2
53	Akut Myeloid LÃ¶semiye BaÄŸlÄ± Bilateral SerÃ¶z MakÃ¼la DekolmanÄ±. TÃ¼rk Oftalmoloji Dergisi, 2014, 44, 151-153.	0.4	2
54	Effectiveness of Fludarabine- and Busulfan-Based Conditioning Regimens in Patients With Acute Myeloblastic Leukemia: 8-Year Experience in a Single Center. Transplantation Proceedings, 2015, 47, 1217-1221.	0.3	2

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55	Development of Acute Promyelocytic Leukemia in a Patient With Gouty Arthritis on Long Term Colchicine. Indian Journal of Hematology and Blood Transfusion, 2016, 32, 80-81.	0.3	2
56	Effects of two doses of anti-T lymphocyte globulin-Fresenius given after full-match sibling stem cell transplantation in acute myeloblastic leukemia patients who underwent myeloablative fludarabine/busulfan conditioning. Hematology/ Oncology and Stem Cell Therapy, 2018, 11, 149-157.	0.6	2
57	Is Sickle Cell Trait Really Innocent?. Turkish Journal of Haematology, 2021, 38, 159-160.	0.2	2
58	Partial Splenic Embolization in Myelodysplastic Syndrome Associated with Immune Thrombocytopenia. Journal of Thrombosis and Thrombolysis, 2004, 18, 213-216.	1.0	1
59	Significance of electronic health records: A comparative study of vaccination rates in patients with sickle cell disease. Pakistan Journal of Medical Sciences, 2017, 33, 549-553.	0.3	1
60	Role of prophylactic and therapeutic red blood cell exchange in pregnancy with sickle cell disease: Maternal and perinatal outcomes. Journal of Clinical Apheresis, 2021, 36, 283-290.	0.7	1
61	Demodocidosis Accompanying Acute Cutaneous Graft-Versus-Host Disease After Allogeneic Stem Cell Transplantation. Turkish Journal of Haematology, 2018, 35, 313-314.	0.2	1
62	A rare hematological complication of visceral leishmaniasis: hemophagocytic syndrome. Ã‡ukurova Ãœniversitesi TÄ±p FakÃ¼ltesi Dergisi, 2016, 41, 161.	0.0	1
63	Implementation of ISBT 128 Compatible Medical Record System to Facilitate Traceability of Stem Cell Products. Turkish Journal of Haematology, 2017, 34, 280-281.	0.2	1
64	Frequency of Finding Family Donors: A Single Center Experience. Experimental and Clinical Transplantation, 2018, 16, 47-50.	0.2	1
65	Significance of Lymphocyte Count, Monocyte Count, and Lymphocyte-To-Monocyte Ratio in Predicting Molecular Response in Patients with Chronic Myeloid Leukemia: a Single-Centre Experience. Clinical Laboratory, 2020, 66, .	0.2	1
66	Problems With Unrelated Donors For Stem Cell Transplant and Proposed Solutions: A Single-Center Experience. Experimental and Clinical Transplantation, 2020, 18, 267-268.	0.2	1
67	Aplastic Anemia in a Professional Musician Exposed to Instrument Polish. International Journal of Hematology, 2005, 81, 304-306.	0.7	0
68	The First 2 Years of Clinical Experience With Peripheral Blood Stem Cell Transplantation for Various Hematological Malignancies: Results From a Single Baskent University Center. Transplantation Proceedings, 2007, 39, 1257-1260.	0.3	0
69	Autologous Serum For Expansion Of Human Bone Marrow Derivated Mesenchymal Stem Cell. Medical Journal of the Trakya University, 2009, , .	0.0	0
70	Weilâ€™s Disease: Four Cases from Ã‡ukurova, Turkey. Klimik Dergisi, 2011, 24, 52-56.	0.1	0
71	Percutaneous vertebroplasty for osteoporotic vertebral fracture in a patient with sickle cell disease. Turkish Journal of Haematology, 2012, 29, 193-194.	0.2	0
72	PP-067 EXPANSION OF BONE MARROW DERIVED MESENCHYMAL STEM CELLS USING A GMP COMPLIANT CLOSED SYSTEM BIOREACTOR AND QUALITY ANALYSIS OF THE EXPANDED CELLS. Leukemia Research, 2014, 38, S48.	0.4	0

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73	Comparative efficacy in red blood cell exchange transfusions with different apheresis machines in patients with sickle cell disease. Indian Journal of Hematology and Blood Transfusion, 2018, 34, 495-500.	0.3	0
74	A Rare and Successfully Managed Complication of Stem Cell Transplantation in an Adult Patient With Sickle Cell Disease: Bone Marrow Necrosis. Experimental and Clinical Transplantation, 2021, , .	0.2	0
75	Is It Possible to be a Stem Cell Donor for the Second Time: A Single-Center Report of 12 Consecutive Procedures. Experimental and Clinical Transplantation, 2021, , .	0.2	0
76	Detection of vitamin B12 levels with the aid of some hematological and biochemical parameters that are more sensitive. Marmara Pharmaceutical Journal, 2010, 3, 125-129.	0.5	0
77	Prevention of recurrence of stroke in a patient with sickle cell disease who has Moyamoya Syndrome. Gaziantep Medical Journal, 2013, 19, 205.	0.2	0
78	Clinical Relevance of Apheretic Graft Composition in Patients With Acute Myeloblastic Leukemia Who Received a Busulfan-Fludarabine-Antithymocyte Globulin Conditioning Regimen for Allogeneic Transplant. Experimental and Clinical Transplantation, 2015, 13, 453-60.	0.2	0
79	TÃ¼rkiyeâ€™ de orak hÃ¼cre hastalÄ±ÄŸÄ±na sahip hastalarda eritrosit alloimmÃ¼nizasyonu: tek merkez geriye dÃ¶nÃ¼k kohort Ã§alÄ±ÅŸmasÄ±. Ã‡ukurova Ãœniversitesi TÄ±p FakÃ¼ltesi Dergisi, 2016, 41, 622-627.	0.0	0
80	Is the Game Over or Starting Again? The Role of the Transplant Team in Genetic Counseling for Adult Sickle Cell Disease Recipients. Turkish Journal of Haematology, 2017, 34, 196-197.	0.2	0
81	Survival of AML Patients Relapsing after Hematopoetic Stem Cell Transplantation: A Single Center Experience. Blood, 2018, 132, 5581-5581.	0.6	0
82	Survival Outcomes of Young Multiple Myeloma Patients: A Single Center Experience. Blood, 2018, 132, 5766-5766.	0.6	0
83	The Impact of the Ferric Carboxymaltose on Hemoglobin and Ferritin Levels. Clinical Laboratory, 2020, 66, .	0.2	0
84	The Clinicopathologic Features and the Factors Associated with the Survival in Light -Chain Amyloidosis Patients: A Single Center Descriptive Study. Medical Journal of Bakirkoy, 2020, , .	0.0	0
85	Rational use of chronic graft-versus-host treatment alternatives: A systematic review. Transfusion and Apheresis Science, 2022, , 103371.	0.5	0