## Emanuele Angelucci

List of Publications by Year in descending order

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250 papers

13,346 citations

18465 62 h-index 27389 106 g-index

256 all docs

256 docs citations

256 times ranked

11925 citing authors

#	Article	IF	CITATIONS
1	Cardiovascular Events and Intensity of Treatment in Polycythemia Vera. New England Journal of Medicine, 2013, 368, 22-33.	13.9	664
2	Bone Marrow Transplantation in Patients with Thalassemia. New England Journal of Medicine, 1990, 322, 417-421.	13.9	580
3	Hepatic Iron Concentration and Total Body Iron Stores in Thalassemia Major. New England Journal of Medicine, 2000, 343, 327-331.	13.9	524
4	Hematopoietic stem cell transplantation in thalassemia major and sickle cell disease: indications and management recommendations from an international expert panel. Haematologica, 2014, 99, 811-820.	1.7	302
5	Effects of iron overload and hepatitis C virus positivity in determining progression of liver fibrosis in thalassemia following bone marrow transplantation. Blood, 2002, 100, 17-21.	0.6	278
6	Chemotherapy-Phased Imatinib Pulses Improve Long-Term Outcome of Adult Patients With Philadelphia Chromosome-Positive Acute Lymphoblastic Leukemia: Northern Italy Leukemia Group Protocol 09/00. Journal of Clinical Oncology, 2010, 28, 3644-3652.	0.8	250
7	Dexamethasone plus rituximab yields higher sustained response rates than dexamethasone monotherapy in adults with primary immune thrombocytopenia. Blood, 2010, 115, 2755-2762.	0.6	242
8	Marrow Transplantation in Patients with Thalassemia Responsive to Iron Chelation Therapy. New England Journal of Medicine, 1993, 329, 840-844.	13.9	211
9	Outcome of patients with hemoglobinopathies given either cord blood or bone marrow transplantation from an HLA-identical sibling. Blood, 2013, 122, 1072-1078.	0.6	210
10	Clinical Effects of Driver Somatic Mutations on the Outcomes of Patients With Myelodysplastic Syndromes Treated With Allogeneic Hematopoietic Stem-Cell Transplantation. Journal of Clinical Oncology, 2016, 34, 3627-3637.	0.8	204
11	Marrow transplantation for patients with thalassemia: results in class 3 patients. Blood, 1996, 87, 2082-2088.	0.6	202
12	Prophylaxis and treatment of hepatitis B in immunocompromised patients. Digestive and Liver Disease, 2007, 39, 397-408.	0.4	197
13	The MLL recombinome of acute leukemias. Leukemia, 2006, 20, 777-784.	3.3	196
14	New approach for bone marrow transplantation in patients with class 3 thalassemia aged younger than 17 years. Blood, 2004, 104, 1201-1203.	0.6	182
15	Italian Society of Hematology practice guidelines for the management of iron overload in thalassemia major and related disorders. Haematologica, 2008, 93, 741-752.	1.7	182
16	Long-term survival of ex-thalassemic patients with persistent mixed chimerism after bone marrow transplantation. Bone Marrow Transplantation, 2000, 25, 401-404.	1.3	177
17	Bone Marrow Transplantation in Adult Thalassemic Patients. Blood, 1999, 93, 1164-1167.	0.6	169
18	ABVD Versus Modified Stanford V Versus MOPPEBVCAD With Optional and Limited Radiotherapy in Intermediate- and Advanced-Stage Hodgkin's Lymphoma: Final Results of a Multicenter Randomized Trial by the Intergruppo Italiano Linfomi. Journal of Clinical Oncology, 2005, 23, 9198-9207.	0.8	167

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19	The cure of thalassemia by bone marrow transplantation. Blood Reviews, 2002, 16, 81-85.	2.8	166
20	Predictive factors for the outcome of allogeneic transplantation in patients with MDS stratified according to the revised IPSS-R. Blood, 2014, 123, 2333-2342.	0.6	162
21	Hemopoietic stem cell transplantation in thalassemia: a report from the European Society for Blood and Bone Marrow Transplantation Hemoglobinopathy Registry, 2000–2010. Bone Marrow Transplantation, 2016, 51, 536-541.	1.3	159
22	Analysis of Heritability and Shared Heritability Based on Genome-Wide Association Studies for Thirteen Cancer Types. Journal of the National Cancer Institute, 2015, 107, djv279.	3.0	152
23	Phlebotomy to Reduce Iron Overload in Patients Cured of Thalassemia by Bone Marrow Transplantation. Blood, 1997, 90, 994-998.	0.6	148
24	Genome-wide association study identifies multiple susceptibility loci for diffuse large B cell lymphoma. Nature Genetics, 2014, 46, 1233-1238.	9.4	147
25	Prognostic impact of pre-transplantation transfusion history and secondary iron overload in patients with myelodysplastic syndrome undergoing allogeneic stem cell transplantation: a GITMO study. Haematologica, 2010, 95, 476-484.	1.7	144
26	Accelerated programmed cell death (apoptosis) in erythroid precursors of patients with severe beta-thalassemia (Cooley's anemia) [see comments]. Blood, 1993, 82, 374-377.	0.6	129
27	Clearance of minimal residual disease after allogeneic stem cell transplantation and the prediction of the clinical outcome of adult patients with high-risk acute lymphoblastic leukemia. Haematologica, 2007, 92, 612-618.	1.7	128
28	Classification and Personalized Prognostic Assessment on the Basis of Clinical and Genomic Features in Myelodysplastic Syndromes. Journal of Clinical Oncology, 2021, 39, 1223-1233.	0.8	127
29	Needle liver biopsy in thalassaemia: analyses of diagnostic accuracy and safety in 1184 consecutive biopsies. British Journal of Haematology, 1995, 89, 757-761.	1.2	125
30	Marrow Transplantation in Patients with Advanced Thalassemia. New England Journal of Medicine, 1987, 316, 1050-1055.	13.9	123
31	Myocardial iron overload assessment by T2* magnetic resonance imaging in adult transfusion dependent patients with acquired anemias. Haematologica, 2008, 93, 1385-1388.	1.7	122
32	The importance of erythroid expansion in determining the extent of apoptosis in erythroid precursors in patients with $\hat{l}^2$ -thalassemia major. Blood, 2000, 96, 3624-3629.	0.6	121
33	MARROW TRANSPLANTATION FOR THALASSAEMIA FOLLOWING BUSULPHAN AND CYCLOPHOSPHAMIDE. Lancet, The, 1985, 325, 1355-1357.	6.3	119
34	Bone marrow transplantation from alternative donors for thalassemia: HLA-phenotypically identical relative and HLA-nonidentical sibling or parent transplants. Bone Marrow Transplantation, 2000, 25, 815-821.	1.3	115
35	Clinical management of myelodysplastic syndromes: update of SIE, SIES, GITMO practice guidelines. Leukemia Research, 2010, 34, 1576-1588.	0.4	112
36	Allogeneic bone marrow transplantation for chronic myelomonocytic leukemia in childhood: a report from the European Working Group on Myelodysplastic Syndrome in Childhood Journal of Clinical Oncology, 1997, 15, 566-573.	0.8	110

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37	Bone Marrow Transplantation in Thalassemia: The Experience of Pesaro. Annals of the New York Academy of Sciences, 1998, 850, 270-275.	1.8	108
38	Long-Term Results of the FOLL05 Trial Comparing R-CVP Versus R-CHOP Versus R-FM for the Initial Treatment of Patients With Advanced-Stage Symptomatic Follicular Lymphoma. Journal of Clinical Oncology, 2018, 36, 689-696.	0.8	107
39	Valproic Acid at Therapeutic Plasma Levels May Increase 5-Azacytidine Efficacy in Higher Risk Myelodysplastic Syndromes. Clinical Cancer Research, 2009, 15, 5002-5007.	3.2	103
40	Incidence, Risk Factors and Outcome of Pre-engraftment Gram-Negative Bacteremia After Allogeneic and Autologous Hematopoietic Stem Cell Transplantation: An Italian Prospective Multicenter Survey. Clinical Infectious Diseases, 2017, 65, 1884-1896.	2.9	103
41	Consensus-based definition of unfitness to intensive and non-intensive chemotherapy in acute myeloid leukemia: a project of SIE, SIES and GITMO group on a new tool for therapy decision making. Leukemia, 2013, 27, 997-999.	<b>3.</b> 3	101
42	Rituximab-dose-dense chemotherapy with or without high-dose chemotherapy plus autologous stem-cell transplantation in high-risk diffuse large B-cell lymphoma (DLCL04): final results of a multicentre, open-label, randomised, controlled, phase 3 study. Lancet Oncology, The, 2017, 18, 1076-1088.	5.1	100
43	Haematopoietic stem cell transplantation trends in children over the last three decades: a survey by the paediatric diseases working party of the European Group for Blood and Marrow Transplantation. Bone Marrow Transplantation, 2007, 39, 89-99.	1.3	95
44	Deferasirox for transfusionâ€dependent patients with myelodysplastic syndromes: safety, efficacy, and beyond ( <scp>GIMEMA MDS</scp> 0306 <scp>T</scp> rial). European Journal of Haematology, 2014, 92, 527-536.	1.1	90
45	Fate of iron stores in thalassaemia after bone-marrow transplantation. Lancet, The, 1993, 342, 1388-1391.	6.3	89
46	Reversibility of Cirrhosis in Patients Cured of Thalassemia by Bone Marrow Transplantation. Annals of Internal Medicine, 2002, 136, 667.	2.0	89
47	Patient- versus physician-reporting of symptoms and health status in chronic myeloid leukemia. Haematologica, 2014, 99, 788-793.	1.7	85
48	Hepatitis B virus-related liver disease in isolated anti-hepatitis B-core positive lymphoma patients receiving chemo- or chemo-immune therapy. Haematologica, 2008, 93, 951-952.	1.7	82
49	Hematopoietic Stem Cell Transplantation in Thalassemia. Hematology American Society of Hematology Education Program, 2010, 2010, 456-462.	0.9	79
50	Management of chronic viral hepatitis in patients with thalassemia: recommendations from an international panel. Blood, 2010, 116, 2875-2883.	0.6	79
51	Iron Chelation in Transfusion-Dependent Patients With Low- to Intermediate-1–Risk Myelodysplastic Syndromes. Annals of Internal Medicine, 2020, 172, 513.	2.0	78
52	Achieving Molecular Remission before Allogeneic Stem Cell Transplantation in Adult Patients with Philadelphia Chromosome–Positive Acute Lymphoblastic Leukemia: Impact on Relapse and Long-Term Outcome. Biology of Blood and Marrow Transplantation, 2016, 22, 1983-1987.	2.0	77
53	Prognostic value of self-reported fatigue on overall survival in patients with myelodysplastic syndromes: a multicentre, prospective, observational, cohort study. Lancet Oncology, The, 2015, 16, 1506-1514.	5.1	76
54	Limitations of Magnetic Resonance Imaging in Measurement of Hepatic Iron. Blood, 1997, 90, 4736-4742.	0.6	73

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55	Differences among young adults, adults and elderly chronic myeloid leukemia patients. Annals of Oncology, 2015, 26, 185-192.	0.6	72
56	VEPEMB in elderly Hodgkin's lymphoma patients. Results from an Intergruppo Italiano Linfomi (IIL) study. Annals of Oncology, 2004, 15, 123-128.	0.6	71
57	Trichosporon beigelii: a life-threatening pathogen in immunocompromised hosts. Bone Marrow Transplantation, 2000, 25, 745-749.	1.3	70
58	GRAFT-VERSUS-HOST DISEASE AFTER BONE MARROW TRANSPLANTATION FOR THALASSEMIA. Transplantation, 1997, 63, 854-860.	0.5	70
59	Halting pro-survival autophagy by $TGF\hat{l}^2$ inhibition in bone marrow fibroblasts overcomes bortezomib resistance in multiple myeloma patients. Leukemia, 2016, 30, 640-648.	3.3	69
60	Myeloablative and Reduced-Intensity Conditioned Allogeneic Hematopoietic Stem Cell Transplantation in Myelofibrosis: A Retrospective Study by the Chronic Malignancies Working Party of the European Society for Blood and Marrow Transplantation. Biology of Blood and Marrow Transplantation, 2019, 25, 2167-2171.	2.0	69
61	Serial monitoring of isavuconazole blood levels during prolonged antifungal therapy. Journal of Antimicrobial Chemotherapy, 2019, 74, 2341-2346.	1.3	69
62	New Strategies in the Treatment of the Thalassemias. Annual Review of Medicine, 2005, 56, 157-171.	5.0	68
63	Post-transplant cyclophosphamide after matched sibling, unrelated and haploidentical donor transplants in patients with acute myeloid leukemia: a comparative study of the ALWP EBMT. Journal of Hematology and Oncology, 2020, 13, 46.	6.9	68
64	Current Results and Future Research Priorities in Late Effects after Hematopoietic Stem Cell Transplantation for Children with Sickle Cell Disease and Thalassemia: A Consensus Statement from the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects after Pediatric Hematopoietic Stem Cell Transplantation. Biology of Blood and Marrow	2.0	66
65	Transplantation, 2017, 23, 552-561.  Dose-dense and high-dose chemotherapy plus rituximab with autologous stem cell transplantation for primary treatment of diffuse large B-cell lymphoma with a poor prognosis: a phase II multicenter study. Haematologica, 2009, 94, 1250-1258.	1.7	65
66	Bone marrow transplantation in adult thalassemia. Blood, 1992, 80, 1603-1607.	0.6	64
67	Hematopoietic stem cell transplantation for paroxysmal nocturnal hemoglobinuria: long-term results of a retrospective study on behalf of the Gruppo Italiano Trapianto Midollo Osseo (GITMO). Haematologica, 2010, 95, 983-988.	1.7	64
68	The importance of erythroid expansion in determining the extent of apoptosis in erythroid precursors in patients with beta-thalassemia major. Blood, 2000, 96, 3624-9.	0.6	64
69	Optimal timing of allogeneic hematopoietic stem cell transplantation in patients with myelodysplastic syndrome. American Journal of Hematology, 2013, 88, 581-588.	2.0	61
70	Multiple Myeloma Treatment in Real-world Clinical Practice: Results of a Prospective, Multinational, Noninterventional Study. Clinical Lymphoma, Myeloma and Leukemia, 2018, 18, e401-e419.	0.2	61
71	Bone Marrow Transplantation in Thalassemia. Hematology/Oncology Clinics of North America, 1991, 5, 549-556.	0.9	59
72	Correction of anemia in a transfusion-dependent patient with primary myelofibrosis receiving iron chelation therapy with deferasirox (Exjade�, ICL670). European Journal of Haematology, 2007, 78, 540-542.	1.1	59

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73	Desferrioxamine therapy accelerates clearance of iron deposits after bone marrow transplantation for thalassaemia. British Journal of Haematology, 1995, 89, 868-873.	1.2	59
74	Allogeneic stem cell transplantation for thalassemia major. Haematologica, 2008, 93, 1780-1784.	1.7	59
<b>7</b> 5	Prevalence, severity and correlates of fatigue in newly diagnosed patients with myelodysplastic syndromes. British Journal of Haematology, 2015, 168, 361-370.	1.2	59
76	A phase II, multicentre trial of decitabine in higher-risk chronic myelomonocytic leukemia. Leukemia, 2018, 32, 413-418.	3.3	58
77	Bendamustine in combination with Ofatumumab in relapsed or refractory chronic lymphocytic leukemia: a GIMEMA Multicenter Phase II Trial. Leukemia, 2014, 28, 642-648.	3.3	57
78	Evaluation of cardiac status in iron-loaded thalassaemia patients following bone marrow transplantation: improvement in cardiac function during reduction in body iron burden. British Journal of Haematology, 1998, 103, 916-921.	1.2	56
79	$HIF-1\hat{1}\pm$ of Bone Marrow Endothelial Cells Implies Relapse and Drug Resistance in Patients with Multiple Myeloma and May Act as a Therapeutic Target. Clinical Cancer Research, 2014, 20, 847-858.	3.2	54
80	Genetically predicted longer telomere length is associated with increased risk of B-cell lymphoma subtypes. Human Molecular Genetics, 2016, 25, 1663-1676.	1.4	52
81	A storm in the niche: Iron, oxidative stress and haemopoiesis. Blood Reviews, 2018, 32, 29-35.	2.8	52
82	Decision analysis of allogeneic hematopoietic stem cell transplantation for patients with myelodysplastic syndrome stratified according to the revised International Prognostic Scoring System. Leukemia, 2017, 31, 2449-2457.	3.3	51
83	Outcome of haploidentical versus matched sibling donors in hematopoietic stem cell transplantation for adult patients with acute lymphoblastic leukemia: a study from the Acute Leukemia Working Party of the European Society for Blood and Marrow Transplantation. Journal of Hematology and Oncology. 2021, 14, 53.	6.9	51
84	Relationship of plasma pharmacokinetics of high-dose oral busulfan to the outcome of allogeneic bone marrow transplantation in children with thalassemia. Bone Marrow Transplantation, 1997, 20, 915-920.	1.3	50
85	A Modified Post-Transplant Cyclophosphamide Regimen, for Unmanipulated Haploidentical Marrow Transplantation, in Acute Myeloid Leukemia: A Multicenter Study. Biology of Blood and Marrow Transplantation, 2018, 24, 1243-1249.	2.0	49
86	Marrow transplantation for patients with thalassemia: results in class 3 patients. Blood, 1996, 87, 2082-8.	0.6	46
87	The role of oxidant injury in the pathophysiology of human thalassemias. Redox Report, 2003, 8, 241-245.	1.4	45
88	Rituximab plus bendamustine as front-line treatment in frail elderly (>70 years) patients with diffuse large B-cell non-Hodgkin lymphoma: a phase II multicenter study of the <i>Fondazione Italiana Linfomi</i>	1.7	45
89	Interleukin-1B (IL1B) and interleukin-6 (IL6) gene polymorphisms are associated with risk of chronic lymphocytic leukaemia. Hematological Oncology, 2008, 26, 98-103.	0.8	44
90	Bone marrow transplantation in adult thalassemic patients. Blood, 1999, 93, 1164-7.	0.6	44

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91	The importance of erythroid expansion in determining the extent of apoptosis in erythroid precursors in patients with $\hat{l}^2$ -thalassemia major. Blood, 2000, 96, 3624-3629.	0.6	43
92	Bone marrow transplantation in thalassemia: modifications of hepatic iron overload and associated lesions after long-term engrafting. Liver, 2008, 14, 14-24.	0.1	42
93	Pre-Engraftment Bloodstream Infections after Allogeneic Hematopoietic Cell Transplantation: Impact of T Cell-Replete Transplantation from a Haploidentical Donor. Biology of Blood and Marrow Transplantation, 2018, 24, 109-118.	2.0	41
94	Haploidentical transplantation is associated with better overall survival when compared to single cord blood transplantation: an EBMT-Eurocord study of acute leukemia patients conditioned with thiotepa, busulfan, and fludarabine. Journal of Hematology and Oncology, 2018, 11, 110.	6.9	41
95	Sudden cardiac tamponade after chemotherapy for marrow transplantation in thalassaemia. Lancet, The, 1992, 339, 287-289.	6.3	40
96	Role of BCL2L10 methylation and TET2 mutations in higher risk myelodysplastic syndromes treated with 5-Azacytidine. Leukemia, 2011, 25, 1910-1913.	3.3	40
97	High cure rates in Burkitt lymphoma and leukemia: a Northern Italy Leukemia Group study of the German short intensive rituximab-chemotherapy program. Haematologica, 2013, 98, 1718-1725.	1.7	40
98	Late Effects Screening Guidelines after Hematopoietic Cell Transplantation (HCT) for Hemoglobinopathy: Consensus Statement from the Second Pediatric Blood and Marrow Transplant Consortium International Conference on Late Effects after Pediatric HCT. Biology of Blood and Marrow Transplantation, 2018, 24, 1313-1321.	2.0	40
99	Impact of spleen size and splenectomy on outcomes of allogeneic hematopoietic cell transplantation for myelofibrosis: A retrospective analysis by the chronic malignancies working party on behalf of European society for blood and marrow transplantation (EBMT). American Journal of Hematology, 2021. 96. 69-79.	2.0	40
100	Management of iron overload before, during, and after hematopoietic stem cell transplantation for thalassemia major. Annals of the New York Academy of Sciences, 2016, 1368, 115-121.	1.8	39
101	Comparing transplant outcomes in ALL patients after haploidentical with PTCy or matched unrelated donor transplantation. Blood Advances, 2020, 4, 2073-2083.	2.5	39
102	Preference for involvement in treatment decisions and request for prognostic information in newly diagnosed patients with higher-risk myelodysplastic syndromes. Annals of Oncology, 2014, 25, 447-454.	0.6	38
103	Impact of HLA Disparity in Haploidentical Bone Marrow Transplantation Followed by High-Dose Cyclophosphamide. Biology of Blood and Marrow Transplantation, 2018, 24, 119-126.	2.0	37
104	$\hat{l}_{\pm}$ -Interferon treatment of chronic hepatitis C after bone marrow transplantation for homozygous $\hat{l}^2$ -thalassemia. Bone Marrow Transplantation, 1997, 20, 767-772.	1.3	36
105	Treatment of Iron Overload in the "Ex-Thalassemic": Report from the Phlebotomy Programa. Annals of the New York Academy of Sciences, 1998, 850, 288-293.	1.8	36
106	De-Escalation and Discontinuation of Empirical Antibiotic Treatment in a Cohort of Allogeneic Hematopoietic Stem Cell Transplantation Recipients during the Pre-Engraftment Period. Biology of Blood and Marrow Transplantation, 2018, 24, 1721-1726.	2.0	36
107	Pentraxin 3 (PTX3) inhibits plasma cell/stromal cell crossâ€ŧalk in the bone marrow of multiple myeloma patients. Journal of Pathology, 2013, 229, 87-98.	2.1	34
108	Determinants of survival in myelofibrosis patients undergoing allogeneic hematopoietic cell transplantation. Leukemia, 2021, 35, 215-224.	3.3	34

#	Article	IF	Citations
109	Anemia in -thalassemia patients targets hepatic hepcidin transcript levels independently of iron metabolism genes controlling hepcidin expression. Haematologica, 2008, 93, 111-115.	1.7	33
110	Azacitidine improves the T-cell repertoire in patients with myelodysplastic syndromes and acute myeloid leukemia with multilineage dysplasia. Leukemia Research, 2015, 39, 957-963.	0.4	32
111	High dose sequential chemotherapy with autologous transplantation versus dose-dense chemotherapy MegaCEOP as first line treatment in poor-prognosis diffuse large cell lymphoma: an "Intergruppo Italiano Linfomi" randomized trial. Haematologica, 2005, 90, 793-801.	1.7	31
112	Impact of donor age and kinship on clinical outcomes after T-cell–replete haploidentical transplantation with PT-Cy. Blood Advances, 2020, 4, 3900-3912.	2.5	30
113	Bone marrow versus mobilized peripheral blood stem cell graft in T-cell-replete haploidentical transplantation in acute lymphoblastic leukemia. Leukemia, 2020, 34, 2766-2775.	3.3	30
114	Second marrow transplants for graft failure in patients with thalassemia. Bone Marrow Transplantation, 1999, 24, 1299-1306.	1.3	29
115	Mutations of TP53 gene in adult acute lymphoblastic leukemia at diagnosis do not affect the achievement of hematologic response but correlate with early relapse and very poor survival. Haematologica, 2016, 101, e245-e248.	1.7	29
116	Reduced intensity <scp>VEPEMB</scp> regimen compared with standard <scp>ABVD</scp> in elderly Hodgkin lymphoma patients: results from a randomized trial on behalf of the Fondazione Italiana Linfomi ( <scp>FIL</scp> ). British Journal of Haematology, 2016, 172, 879-888.	1.2	29
117	Risk of malignant lymphoma following viral hepatitis infection. International Journal of Hematology, 2008, 87, 474-483.	0.7	28
118	Reduced Left Ventricular Contractile Reserve Identified by Low Dose Dobutamine Echocardiography as an Early Marker of Cardiac Involvement in Asymptomatic Patients with Thalassemia Major. Echocardiography, 1996, 13, 463-472.	0.3	27
119	Transplantation in thalassemia: Revisiting the Pesaro risk factors 25 years later. American Journal of Hematology, 2017, 92, 411-413.	2.0	27
120	HLA-Mismatched Donors in Patients with Myelodysplastic Syndrome: An EBMT Registry Analysis. Biology of Blood and Marrow Transplantation, 2019, 25, 114-120.	2.0	27
121	Post-transplant cyclophosphamide containing regimens after matched sibling, matched unrelated and haploidentical donor transplants in patients with acute lymphoblastic leukemia in first complete remission, a comparative study of the ALWP of the EBMT. Journal of Hematology and Oncology, 2021, 14.84.	6.9	27
122	Cytarabine and clofarabine after highâ€dose cytarabine in relapsed or refractory AML patients. American Journal of Hematology, 2012, 87, 1047-1051.	2.0	26
123	Intravenous chelation therapy during transplantation for thalassemia. Haematologica, 1995, 80, 300-4.	1.7	26
124	Kaposi's sarcoma after allogeneic bone marrow transplantation. Bone Marrow Transplantation, 1997, 19, 629-631.	1.3	25
125	CD20 expression has no prognostic role in Philadelphia-negative B-precursor acute lymphoblastic leukemia: new insights from the molecular study of minimal residual disease. Haematologica, 2012, 97, 568-571.	1.7	25
126	Unraveling the mechanisms behind iron overload and ineffective hematopoiesis in myelodysplastic syndromes. Leukemia Research, 2017, 62, 108-115.	0.4	25

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127	Early mortality in myeloma patients treated with first-generation novel agents thalidomide, lenalidomide, bortezomib at diagnosis: A pooled analysis. Critical Reviews in Oncology/Hematology, 2018, 130, 27-35.	2.0	25
128	Comparison of Haploidentical Bone Marrow versus Matched Unrelated Donor Peripheral Blood Stem Cell Transplantation with Posttransplant Cyclophosphamide in Patients with Acute Leukemia. Clinical Cancer Research, 2021, 27, 843-851.	3.2	25
129	Secondary acute myeloid leukaemia: results of conventional treatments. Experience of GIMEMA trials. Annals of Oncology, 2005, 16, 228-233.	0.6	24
130	Ironâ€chelating therapy with deferasirox in transfusionâ€dependent, higher risk myelodysplastic syndromes: a retrospective, multicentre study. British Journal of Haematology, 2017, 177, 741-750.	1.2	23
131	Bone marrow transplantation in thalassemia. Cancer Treatment and Research, 1997, 77, 305-315.	0.2	23
132	Treosulfan/fludarabine as an allogeneic hematopoietic stem cell transplant conditioning regimen for highâ€risk patients. American Journal of Hematology, 2008, 83, 717-720.	2.0	22
133	Healthâ€related quality of life in patients with chronic myeloid leukemia receiving firstâ€line therapy with nilotinib. Cancer, 2018, 124, 2228-2237.	2.0	22
134	Graft versus host disease in unmanipulated haploidentical marrow transplantation with a modified post-transplant cyclophosphamide (PT-CY) regimen: an update on 425 patients. Bone Marrow Transplantation, 2019, 54, 708-712.	1.3	22
135	Acute Pancreatitis Following Brentuximab Vedotin Therapy for Refractory Hodgkin Lymphoma: A Case Report. Drugs in R and D, 2014, 14, 9-11.	1.1	21
136	Treatment of hepatitis C in patients with thalassemia. Haematologica, 2008, 93, 1121-1123.	1.7	20
137	Haploidentical bone marrow transplantation from mother to child with advanced leukemia. Bone Marrow Transplantation, 1995, 16, 529-35.	1.3	20
138	Bone marrow transplantation for transfused patients with severe aplastic anemia using cyclophosphamide and total lymphoid irradiation as conditioning therapy: long-term follow-up from a single center. Bone Marrow Transplantation, 1999, 24, 253-257.	1.3	19
139	Updated recommendations on the management of gastrointestinal disturbances during iron chelation therapy with Deferasirox in transfusion dependent patients with myelodysplastic syndrome – Emphasis on optimized dosing schedules and new formulations. Leukemia Research, 2015, 39, 1028-1033.	0.4	19
140	Allogeneic Hemopoietic Stem Cell Transplants in Patients with Acute Myeloid Leukemia (AML) Prepared with Busulfan and Fludarabine (BUFLU) or Thiotepa, Busulfan, and Fludarabine (TBF): A Retrospective Study. Biology of Blood and Marrow Transplantation, 2020, 26, 698-703.	2.0	19
141	The impact of comorbidity on health-related quality of life in elderly patients with chronic myeloid leukemia. Annals of Hematology, 2016, 95, 211-219.	0.8	18
142	Validation of the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire Core 30 Summary Score in Patients With Hematologic Malignancies. Value in Health, 2019, 22, 1303-1310.	0.1	18
143	Use of Aspergillus fumigatus real-time PCR in bronchoalveolar lavage samples (BAL) for diagnosis of invasive aspergillosis, including azole-resistant cases, in high risk haematology patients: the need for a combined use with galactomannan. Medical Mycology, 2019, 57, 987-996.	0.3	18
144	Pubertal development in thalassaemic patients after allogenic bone marrow transplantation. European Journal of Pediatrics, 1993, 152, 993-997.	1.3	17

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145	Accuracy of physician assessment of treatment preferences and health status in elderly patients with higher-risk myelodysplastic syndromes. Leukemia Research, 2015, 39, 859-865.	0.4	17
146	Haploidentical Transplants with Post-Transplant Cyclophosphamide for Relapsed or Refractory Hodgkin Lymphoma: The Role of Comorbidity Index and Pretransplant Positron Emission Tomography. Biology of Blood and Marrow Transplantation, 2018, 24, 2501-2508.	2.0	17
147	Hepatitis C virus infection in thalassemia patients undergoing allogeneic bone marrow transplantation. Bone Marrow Transplantation, 1994, 14, 369-72.	1.3	17
148	Fate of chronic myeloid leukemia patients treated with allogeneic bone marrow transplantation or chemotherapy and/or interferon at a single center: long-term results. Bone Marrow Transplantation, 2002, 29, 1-8.	1.3	16
149	Dramatic erythroid response to lowâ€dose thalidomide in two patients with transfusion independent thalassemia and severe postâ€transfusional alloimmune hemolysis. American Journal of Hematology, 2015, 90, E141.	2.0	16
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