## Mary Frances McMullin

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2764989/publications.pdf

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235 papers 11,751 citations

51 h-index 35168 102 g-index

237 all docs

237 docs citations

times ranked

237

10764 citing authors

#	Article	IF	CITATIONS
1	CALR type 1 mutations are associated with an increased incidence of myelofibrosis in young male patients. Irish Journal of Medical Science, 2023, 192, 591-593.	0.8	1
2	A randomized phase 3 trial of interferon- $\hat{l}_{\pm}$ vs hydroxyurea in polycythemia vera and essential thrombocythemia. Blood, 2022, 139, 2931-2941.	0.6	45
3	FEDORA: The Beginning of a Beautiful Friendship?. , 2022, 19, .		O
4	Spliceosome mutations are common in persons with myeloproliferative neoplasm-associated myelofibrosis with RBC-transfusion-dependence and correlate with response to pomalidomide. Leukemia, 2021, 35, 1197-1202.	3.3	9
5	Results of a national UK physician reported survey of COVID-19 infection in patients with a myeloproliferative neoplasm. Leukemia, 2021, 35, 2424-2430.	3.3	8
6	Long-term safety and efficacy of givinostat in polycythemia vera: 4-year mean follow up of three phase 1/2 studies and a compassionate use program. Blood Cancer Journal, 2021, 11, 53.	2.8	24
7	Defining the Optimal Total Number of Chemotherapy Courses in Younger Patients With Acute Myeloid Leukemia: A Comparison of Three Versus Four Courses. Journal of Clinical Oncology, 2021, 39, 890-901.	0.8	20
8	MOMENTUM: momelotinib vs danazol in patients with myelofibrosis previously treated with JAKi who are symptomatic and anemic. Future Oncology, 2021, 17, 1449-1458.	1.1	31
9	Protein tyrosine phosphatase receptor type C (PTPRC or CD45). Journal of Clinical Pathology, 2021, 74, 548-552.	1.0	90
10	Molecular pathogenesis of the myeloproliferative neoplasms. Journal of Hematology and Oncology, 2021, 14, 103.	6.9	49
11	Genetic Background of Congenital Erythrocytosis. Genes, 2021, 12, 1151.	1.0	8
12	Hereditary thrombocytosis: the genetic landscape. British Journal of Haematology, 2021, 194, 1098-1105.	1.2	6
13	Significance of NPM1 Gene Mutations in AML. International Journal of Molecular Sciences, 2021, 22, 10040.	1.8	18
14	Realâ€world tyrosine kinase inhibitor treatment pathways, monitoring patterns and responses in patients with chronic myeloid leukaemia in the United Kingdom: the UK TARGET CML study. British Journal of Haematology, 2021, 192, 62-74.	1.2	18
15	Correlation of Quality of Life between Treatment Outcomes in the Majic Study Which Compared Ruxolitinib to Best Available Therapy in Polycythemia Vera. Blood, 2021, 138, 3644-3644.	0.6	O
16	A Randomised Evaluation of Low-Dose Ara-C Plus BCT-100 Versus Low Dose Ara-C in Older Patients with Acute Myeloid Leukaemia: Results from the LI-1 Trial. Blood, 2021, 138, 2355-2355.	0.6	0
17	Exploring Genotype:Phenotype Correlations at Baseline and at One Year for ET and PV Patients in the Majic Study. Blood, 2021, 138, 1507-1507.	0.6	0
18	A molecular diagnostic algorithm for JAK2 V617F investigations in suspected myeloproliferative neoplasms. Irish Journal of Medical Science, 2020, 189, 621-626.	0.8	2

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19	Second cancers in MPN: Survival analysis from an international study. American Journal of Hematology, 2020, 95, 295-301.	2.0	34
20	Modification of the Histone Landscape with JAK Inhibition in Myeloproliferative Neoplasms. Cancers, 2020, 12, 2669.	1.7	6
21	Mutational profiling in suspected triple-negative essential thrombocythaemia using targeted next-generation sequencing in a real-world cohort. Journal of Clinical Pathology, 2020, 74, jclinpath-2020-206570.	1.0	10
22	Aetiology of Myeloproliferative Neoplasms. Cancers, 2020, 12, 1810.	1.7	16
23	Modifiable Lifestyle and Medical Risk Factors Associated With Myeloproliferative Neoplasms. HemaSphere, 2020, 4, e327.	1.2	18
24	Safety and efficacy of the maximum tolerated dose of givinostat in polycythemia vera: a two-part Phase lb/II study. Leukemia, 2020, 34, 2234-2237.	3.3	34
25	Arterial thrombosis in Philadelphia-negative myeloproliferative neoplasms predicts second cancer: a case-control study. Blood, 2020, 135, 381-386.	0.6	18
26	Ruxolitinib: gaining more than intended. Blood, 2020, 135, 983-984.	0.6	O
27	Effects of Tamoxifen on the Mutant Allele Burden and Disease Course in Patients with Myeloproliferative Neoplasms - Results of the Tamarin Study. Blood, 2020, 136, 33-35.	0.6	6
28	Presidential address to the Ulster Medical Society on 3rd October 2019. Ulster Medical Journal, 2020, 89, 72-76.	0.2	0
29	Somatic <i>SF3B1</i> mutations in myelodysplastic syndrome with ring sideroblasts and chronic lymphocytic leukaemia. Journal of Clinical Pathology, 2019, 72, 778-782.	1.0	17
30	Methylation age as a correlate for allele burden, disease status, and clinical response in myeloproliferative neoplasm patients treated with vorinostat. Experimental Hematology, 2019, 79, 26-34.	0.2	8
31	Second cancer in Philadelphia negative myeloproliferative neoplasms (MPN-K). A nested case-control study. Leukemia, 2019, 33, 1996-2005.	3.3	67
32	Facing erythrocytosis: Results of an international physician survey. American Journal of Hematology, 2019, 94, E225-E227.	2.0	10
33	The poor outcome in high molecular risk, hydroxycarbamide-resistant/intolerant ET is not ameliorated by ruxolitinib. Blood, 2019, 134, 2107-2111.	0.6	12
34	Reducing the burden of MPN. Blood, 2019, 134, 1483-1484.	0.6	0
35	Acute promyelocytic leukaemia ( <scp>APML</scp> ) with cryptic <i><scp>PML</scp>â€<scp>RARA</scp></i> fusion has a clinical course comparable to classical <scp>APML</scp> with t(15;17)(q24.1;q21.2) translocation. British Journal of Haematology, 2019, 186, 155-157.	1.2	3
36	A guideline for the management of specific situations in polycythaemia vera and secondary erythrocytosis. British Journal of Haematology, 2019, 184, 161-175.	1.2	76

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37	A guideline for the diagnosis and management of polycythaemia vera. A British Society for Haematology Guideline. British Journal of Haematology, 2019, 184, 176-191.	1.2	102
38	Final Results of Prospective Treatment with Pegylated Interferon Alfa-2a for Patients with Polycythemia Vera and Essential Thrombocythemia in First and Second-Line Settings. Blood, 2019, 134, 2943-2943.	0.6	4
39	Frequency of Thrombosis Is Higher in MPN Patients Who Develop Second Cancer Than in Controls. Blood, 2019, 134, 4170-4170.	0.6	2
40	Diagnostic workflow for hereditary erythrocytosis and thrombocytosis. Hematology American Society of Hematology Education Program, 2019, 2019, 391-396.	0.9	12
41	<i><scp>HFE</scp></i> mutations in idiopathic erythrocytosis. British Journal of Haematology, 2018, 181, 270-272.	1.2	13
42	The MOSAICC study: Assessing feasibility for biological sample collection in epidemiology studies and comparison of DNA yields from saliva and whole blood samples. Annals of Human Genetics, 2018, 82, 114-118.	0.3	6
43	A spotlight on the management of complications associated with myeloproliferative neoplasms: a clinician $\hat{a} \in \mathbb{R}^{M}$ s perspective. Expert Review of Hematology, 2018, 11, 25-35.	1.0	0
44	Hydroxycarbamide Plus Aspirin Versus Aspirin Alone in Patients With Essential Thrombocythemia Age 40 to 59 Years Without High-Risk Features. Journal of Clinical Oncology, 2018, 36, 3361-3369.	0.8	54
45	The ruxolitinib effect: understanding how molecular pathogenesis and epigenetic dysregulation impact therapeutic efficacy in myeloproliferative neoplasms. Journal of Translational Medicine, 2018, 16, 360.	1.8	50
46	Splanchnic venous thrombosis in JAK2 V617F mutation positive myeloproliferative neoplasms – long term follow-up of a regional case series. Thrombosis Journal, 2018, 16, 33.	0.9	11
47	Classification and Personalized Prognosis in Myeloproliferative Neoplasms. New England Journal of Medicine, 2018, 379, 1416-1430.	13.9	442
48	Absence of <b><i>CALR</i></b> Mutations in Idiopathic Erythrocytosis Patients with Low Serum Erythropoietin Levels. Acta Haematologica, 2018, 139, 217-219.	0.7	3
49	Impact on MPN Symptoms and Quality of Life of Front Line Pegylated Interferon Alpha-2a Vs. Hydroxyurea in High Risk Polycythemia Vera and Essential Thrombocythemia: Results of Myeloproliferative Disorders Research Consortium (MPD-RC) 112 Global Phase III Trial. Blood, 2018, 132, 3032-3032.	0.6	6
50	Results of the Myeloproliferative Neoplasms - Research Consortium (MPN-RC) 112 Randomized Trial of Pegylated Interferon Alfa-2a (PEG) Versus Hydroxyurea (HU) Therapy for the Treatment of High Risk Polycythemia Vera (PV) and High Risk Essential Thrombocythemia (ET). Blood, 2018, 132, 577-577.	0.6	39
51	Risk Factors for Secondary Cancer in a Case-Control Study on 1,259 Patients with Myeloproliferative Neoplasms. Blood, 2018, 132, 4279-4279.	0.6	1
52	Spliceosome Mutations Are Common in MPN-Associated Myelofibrosis with RBC-Transfusion-Dependence and Correlate with Response to Pomalidomide. Blood, 2018, 132, 3037-3037.	0.6	0
53	Longitudinal Mutational Analysis in Hydroxycarbamide-Resistant/Intolerant Essential Thrombocythemia Treated on the Majic-ET Study. Blood, 2018, 132, 1784-1784.	0.6	O
54	Myeloproliferative Neoplasm Quality of Life (MPN-QOL) Study Group: MPN Experimental Assessment of Symptoms By Utilizing Repetitive Evaluation (MEASURE) Trial. Blood, 2018, 132, 1762-1762.	0.6	1

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55	Myeloproliferative Neoplasms in Patients below 25 Years Old at Diagnosis: A Retrospective International Cooperative Work. Blood, 2018, 132, 1759-1759.	0.6	О
56	Equivalence of BCSH and WHO diagnostic criteria for ET. Leukemia, 2017, 31, 527-528.	3.3	6
57	70 years of the <i>JCP</i> -highly cited papers: Proposals for the classification of chronic (mature) B and T lymphoid leukaemias. Journal of Clinical Pathology, 2017, 70, 909-910.	1.0	1
58	Epigenetics in Myeloproliferative Neoplasms. Journal of Cellular and Molecular Medicine, 2017, 21, 1660-1667.	1.6	29
59	Outcome of Azacitidine Therapy in Acute Myeloid Leukemia Is not Improved by Concurrent Vorinostat Therapy but Is Predicted by a Diagnostic Molecular Signature. Clinical Cancer Research, 2017, 23, 6430-6440.	3.2	74
60	Ruxolitinib vs best available therapy for ET intolerant or resistant to hydroxycarbamide. Blood, 2017, 130, 1889-1897.	0.6	130
61	Which patients with myelofibrosis should receive ruxolitinib therapy? ELN-SIE evidence-based recommendations. Leukemia, 2017, 31, 882-888.	3.3	40
62	Management of polycythaemia vera: a critical review of current data. British Journal of Haematology, 2016, 172, 337-349.	1.2	28
63	LNK mutations and myeloproliferative disorders. American Journal of Hematology, 2016, 91, 248-251.	2.0	31
64	TheCalreticulingene and myeloproliferative neoplasms. Journal of Clinical Pathology, 2016, 69, 841-845.	1.0	12
65	Coexistence of inversion 16 in chronic myeloid leukaemia in blast crisis. Journal of Hematopathology, 2016, 9, 155-160.	0.2	О
66	Congenital erythrocytosis. International Journal of Laboratory Hematology, 2016, 38, 59-65.	0.7	13
67	Antiplatelet therapy versus observation in low-risk essential thrombocythemia with a CALR mutation. Haematologica, 2016, 101, 926-931.	1.7	118
68	Investigation and Management of Erythrocytosis. Current Hematologic Malignancy Reports, 2016, 11, 342-347.	1.2	36
69	Gene panel sequencing improves the diagnostic work-up of patients with idiopathic erythrocytosis and identifies new mutations. Haematologica, 2016, 101, 1306-1318.	1.7	66
70	Experience of Myeloproliferative Neoplasms Guidelines in the United Kingdom: Perspective and International Context. Journal of the National Comprehensive Cancer Network: JNCCN, 2016, 14, e1626-e1631.	2.3	0
71	Minor allele frequency of myeloproliferative neoplasm mutations in the Irish blood donor population. Hematological Oncology, 2016, 34, 161-164.	0.8	О
72	The prevalence of CALR mutations in a cohort of patients with myeloproliferative neoplasms. International Journal of Laboratory Hematology, 2016, 38, 102-106.	0.7	3

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73	Vorinostat Does Not Improve Outcome in Patients with Acute Myeloid Leukemia and High Risk Myelodysplasia Treated with Azacitidine: Results of the UK Trials Acceleration Programme Ravva Trial. Blood, 2016, 128, 1065-1065.	0.6	3
74	A Two-Part Study of Givinostat in Patients with Polycythemia Vera: Maximum Tolerated Dose Definition and Preliminary Efficacy Results. Blood, 2016, 128, 4261-4261.	0.6	6
75	Impact on MPN Symptoms and Quality of Life of Front Line Pegylated Interferon Alpha-2a Vs. Hydroxyurea in High Risk Polycythemia Vera and Essential Thrombocythemia: Interim Analysis Results of Myeloproliferative Disorders Research Consortium (MPD-RC) 112 Global Phase III Trial. Blood, 2016, 128. 4271-4271.	0.6	5
76	Interim Analysis of the Myeloproliferative Disorders Research Consortium (MPD-RC) 112 Global Phase III Trial of Front Line Pegylated Interferon Alpha-2a Vs. Hydroxyurea in High Risk Polycythemia Vera and Essential Thrombocythemia. Blood, 2016, 128, 479-479.	0.6	32
77	Correlation Between Treatment Outcomes, Baseline Characteristics and Molecular Responses in the Majic Study Which Compared Ruxolitinib to Best Available Therapy in Essential Thrombocythemia. Blood, 2016, 128, 1929-1929.	0.6	O
78	Myeloproliferative Neoplasm Quality of Life (MPN-QOL) Study Group: Interim Results from the MPN Experimental Assessment of Symptoms By Utilizing Repetitive Evaluation (MEASURE) Trial. Blood, 2016, 128, 5479-5479.	0.6	0
79	(32)P in the treatment of myeloproliferative disorders. Ulster Medical Journal, 2016, 85, 83-5.	0.2	2
80	Outcomes of pregnancy in patients with congenital erythrocytosis. British Journal of Haematology, 2015, 170, 586-588.	1.2	3
81	Molecular diagnostics of myeloproliferative neoplasms. European Journal of Haematology, 2015, 95, 270-279.	1.1	67
82	Patient perspectives of a diagnosis of myeloproliferative neoplasm in a case control study. Experimental Hematology and Oncology, 2015, 5, 14.	2.0	3
83	A Japanese Family with Congenital Erythrocytosis Caused by Haemoglobin Bethesda. Internal Medicine, 2015, 54, 2389-2393.	0.3	O
84	The use of erythropoiesis-stimulating agents with ruxolitinib in patients with myelofibrosis in COMFORT-II: an open-label, phase 3 study assessing efficacy and safety of ruxolitinib versus best available therapy in the treatment of myelofibrosis. Experimental Hematology and Oncology, 2015, 4, 26.	2.0	36
85	Myeloproliferative neoplasm patient symptom burden and quality of life: Evidence of significant impairment compared to controls. American Journal of Hematology, 2015, 90, 864-870.	2.0	33
86	Management of newly diagnosed chronic myeloid leukaemia during a twin pregnancy using leucapheresis: Case report and review of the literature. Transfusion and Apheresis Science, 2015, 52, 199-203.	0.5	5
87	Protein deregulation associated with breast cancer metastasis. Cytokine and Growth Factor Reviews, 2015, 26, 415-423.	3.2	7
88	Factors influencing success of clinical genome sequencing across a broad spectrum of disorders. Nature Genetics, 2015, 47, 717-726.	9.4	310
89	Arsenic trioxide and all-trans retinoic acid treatment for acute promyelocytic leukaemia in all risk groups (AML17): results of a randomised, controlled, phase 3 trial. Lancet Oncology, The, 2015, 16, 1295-1305.	5.1	433
90	Assessment and Validation of the EQ-5D Among a Population of Myeloproliferative Neoplasm Patients. Blood, 2015, 126, 5179-5179.	0.6	2

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91	Myeloproliferative Neoplasm Patient Symptom Burden and Quality of Life: Evidence of Significant Impairment Compared to Controls Using Multivariate Analysis. Blood, 2015, 126, 1620-1620.	0.6	O
92	Myeloproliferative Neoplasms: An in-Depth Case-Control (MOSAICC) Study. Blood, 2015, 126, 1621-1621.	0.6	12
93	Conditional Deletion of the Hoxa Cluster in MLL-AF9 Is Incompatible with Leukemia Maintenance. Blood, 2015, 126, 3630-3630.	0.6	O
94	A Comparison of $1$ or $2$ Courses of High Dose Cytarabine As Consolidation in Younger Patients with AML: First Results of the UK NCRI AML17 Trial. Blood, 2015, 126, 221-221.	0.6	1
95	The role of PHD2 mutations in the pathogenesis of erythrocytosis. Hypoxia (Auckland, N Z ), 2014, 2, 71.	1.9	39
96	Clinical utility gene card for: Hereditary thrombocythemia. European Journal of Human Genetics, 2014, 22, 293-293.	1.4	8
97	Epidemiology of MPN: What Do We Know?. Current Hematologic Malignancy Reports, 2014, 9, 340-349.	1.2	27
98	Genetic Basis of Congenital Erythrocytosis: Mutation Update and Online Databases. Human Mutation, 2014, 35, 15-26.	1.1	101
99	Assessment of <i>CALR </i> mutations in myelofibrosis patients, post-allogeneic stem cell transplantation. British Journal of Haematology, 2014, 166, 800-802.	1.2	30
100	Use of <scp>JAK</scp> inhibitors in the management of myelofibrosis: a revision of the <scp>B</scp> ritish <scp>C</scp> ommittee for <scp>S</scp> tandards in <scp>H</scp> aematology <scp>G</scp> uidelines for <scp>I</scp> nvestigation and <scp>M</scp> anagement of <scp>M</scp> yelofibrosis 2012. British Journal of Haematology, 2014, 167, 418-420.	1.2	37
101	Secondary erythrocytosis. Hematology, 2014, 19, 183-184.	0.7	10
102	Update in the myeloproliferative neoplasms. Clinical Medicine, 2014, 14, s66-s70.	0.8	0
103	Retrospective study of alemtuzumab vs ATG-based conditioning without irradiation for unrelated and matched sibling donor transplants in acquired severe aplastic anemia: a study from the British Society for Blood and Marrow Transplantation. Bone Marrow Transplantation, 2014, 49, 42-48.	1.3	65
104	How common are myeloproliferative neoplasms? A systematic review and metaâ€analysis. American Journal of Hematology, 2014, 89, 581-587.	2.0	141
105	Polycythaemia-inducing mutations in the erythropoietin receptor (EPOR): mechanism and function as elucidated by epidermal growth factor receptor-EPOR chimeras. British Journal of Haematology, 2014, 165, 519-528.	1.2	13
106	Nilotinib 300 mg BID as frontline treatment of CML: Prospective analysis of the Xpert BCR-ABL Monitor system and significance of 3-month molecular response. Leukemia Research, 2014, 38, 310-315.	0.4	12
107	Modification of British Committee for Standards in Haematology diagnostic criteria for essential thrombocythaemia. British Journal of Haematology, 2014, 167, 421-423.	1.2	40
108	Circulating YKL-40 in patients with essential thrombocythemia and polycythemia vera treated with the novel histone deacetylase inhibitor vorinostat. Leukemia Research, 2014, 38, 816-821.	0.4	12

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109	Community-acquired infections and their association with myeloid malignancies. Cancer Epidemiology, 2014, 38, 56-61.	0.8	36
110	Mutational spectrum defines primary and secondary myelofibrosis. Haematologica, 2014, 99, 2-3.	1.7	10
111	Erythrocytosis associated with a novel missense mutation in the BPGM gene. Haematologica, 2014, 99, e201-e204.	1.7	35
112	A Comparison of Single Dose Gemtuzumab Ozogamicin 3mg/m2 and 6mg/m2 Combined with Induction Chemotherapy in Younger Patients with AML: Data from the UK NCRI AML17 Trial. Blood, 2014, 124, 2308-2308.	0.6	2
113	A phase II study of vorinostat ( <scp>MK</scp> â€0683) in patients with polycythaemia vera and essential thrombocythaemia. British Journal of Haematology, 2013, 162, 498-508.	1.2	65
114	Establishing optimal quantitative-polymerase chain reaction assays for routine diagnosis and tracking of minimal residual disease in JAK2-V617F-associated myeloproliferative neoplasms: a joint European LeukemiaNet/MPN&MPNr-EuroNet (COST action BM0902) study. Leukemia, 2013, 27, 2032-2039.	3.3	96
115	Molecular diagnosis of the myeloproliferative neoplasms: <scp>UK</scp> guidelines for the detection of <i><scp>JAK</scp>2 </i> <scp>V</scp> 617 <scp>F</scp> and other relevant mutations. British Journal of Haematology, 2013, 160, 25-34.	1.2	87
116	Prognostic and therapeutic relevance of câ€ <scp>FLIP</scp> in acute myeloid leukaemia. British Journal of Haematology, 2013, 160, 188-198.	1.2	39
117	Inclusion of chemotherapy in addition to anthracycline in the treatment of acute promyelocytic leukaemia does not improve outcomes: results of the MRC AML15 trial. Leukemia, 2013, 27, 843-851.	3.3	74
118	The diagnosis and management of erythrocytosis. BMJ, The, 2013, 347, f6667-f6667.	3.0	67
119	Clofarabine doubles the response rate in older patients with acute myeloid leukemia but does not improve survival. Blood, 2013, 122, 1384-1394.	0.6	123
120	Diagnostic pathway for the investigation of thrombocytosis. British Journal of Haematology, 2013, 161, 604-606.	1.2	6
121	Characteristics and outcomes of haematology patients admitted to the intensive care unit. Nursing in Critical Care, 2013, 18, 193-199.	1.1	6
122	The addition of gemtuzumab ozogamicin to low-dose Ara-C improves remission rate but does not significantly prolong survival in older patients with acute myeloid leukaemia: results from the LRF AML14 and NCRI AML16 pick-a-winner comparison. Leukemia, 2013, 27, 75-81.	3.3	146
123	Revised response criteria for polycythemia vera and essential thrombocythemia: an ELN and IWG-MRT consensus project. Blood, 2013, 121, 4778-4781.	0.6	219
124	Erythrocytosis in children and adolescents-classification, characterization, and consensus recommendations for the diagnostic approach. Pediatric Blood and Cancer, 2013, 60, 1734-1738.	0.8	26
125	HOXA/PBX3 knockdown impairs growth and sensitizes cytogenetically normal acute myeloid leukemia cells to chemotherapy. Haematologica, 2013, 98, 1216-1225.	1.7	39
126	Cooperativity of imprinted genes inactivated by acquired chromosome 20q deletions. Journal of Clinical Investigation, 2013, 123, 2169-2182.	3.9	36

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127	Reasons For Survival Improvement In Core Binding Factor AML: A 25 Year Analysis Of The UK MRC/NCRI AML Trials. Blood, 2013, 122, 358-358.	0.6	9
128	Phase 3 Study Of Pomalidomide In Myeloproliferative Neoplasm (MPN)-Associated Myelofibrosis With RBC-Transfusion-Dependence. Blood, 2013, 122, 394-394.	0.6	29
129	Polycythemia-Inducing Mutations In The Erythropoietin Receptor (EPOR): Mechanism and Function Elucidated By EGFR– EPOR Chimeras. Blood, 2013, 122, 2174-2174.	0.6	O
130	Diagnosis and management of congenital and idiopathic erythrocytosis. Therapeutic Advances in Hematology, 2012, 3, 391-398.	1.1	24
131	Clinical utility gene card for: familial erythrocytosis. European Journal of Human Genetics, 2012, 20, 4-4.	1.4	10
132	Commentary. Clinical Chemistry, 2012, 58, 335-335.	1.5	1
133	Management of myelofibrosis: a survey of current practice in the United Kingdom. Journal of Clinical Pathology, 2012, 65, 1124-1127.	1.0	2
134	JAK2V617F homozygosity arises commonly and recurrently in PV and ET, but PV is characterized by expansion of a dominant homozygous subclone. Blood, 2012, 120, 2704-2707.	0.6	94
135	<scp>J</scp> anus kinase Inhibition and its effect upon the therapeutic landscape for myelofibrosis: from palliation to cure?. British Journal of Haematology, 2012, 157, 426-437.	1.2	19
136	Guideline for the diagnosis and management of myelofibrosis. British Journal of Haematology, 2012, 158, 453-471.	1.2	89
137	Environmental, lifestyle, and familial/ethnic factors associated with myeloproliferative neoplasms. American Journal of Hematology, 2012, 87, 175-182.	2.0	35
138	Two new mutations in the <i>HIF2A</i> gene associated with erythrocytosis. American Journal of Hematology, 2012, 87, 439-442.	2.0	37
139	Symptom Burden Among PV and ET Patients Receiving A Novel Histone Deacetylase Inhibitor: Findings From a Open-Label Phase II Study. Blood, 2012, 120, 1736-1736.	0.6	1
140	Expand: a Phase 1b, Open-Label, Dose-Finding Study of Ruxolitinib in Patients with Myelofibrosis and Baseline Platelet Counts Between 50 × 109/L and 99 × 109/L. Blood, 2012, 120, 177-177.	0.6	7
141	The Use of Erythropoietic-Stimulating Agents (ESAs) with Ruxolitinib in Patients with Primary Myelofibrosis (PMF), Post-Polycythemia Vera Myelofibrosis (PPV-MF), and Post-Essential Thrombocythemia Myelofibrosis (PET-MF) Blood, 2012, 120, 2838-2838.	0.6	7
142	A Phase II Study of Vorinostat (MK-0683) in Patients with Polycythemia Vera and Essential Thrombocythemia. Blood, 2012, 120, 803-803.	0.6	4
143	A Randomised Comparison of Clofarabine Versus Low Dose Ara-C As First Line Treatment for Older Patients with AML. Blood, 2012, 120, 889-889.	0.6	4
144	Critical Issues About the Diagnosis of Myeloproliferative Neoplasms: World Health Organization Classification., 2012,, 37-45.		0

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145	Molecular and clinical features of the myeloproliferative neoplasm associated with JAK2 exon 12 mutations. Blood, 2011, 117, 2813-2816.	0.6	190
146	Philadelphia-Negative Classical Myeloproliferative Neoplasms: Critical Concepts and Management Recommendations From European LeukemiaNet. Journal of Clinical Oncology, 2011, 29, 761-770.	0.8	724
147	A nonsynonymous <i>LNK</i> polymorphism associated with idiopathic erythrocytosis. American Journal of Hematology, 2011, 86, 962-964.	2.0	30
148	Cardiopulmonary function in two human disorders of the hypoxiaâ€inducible factor (HIF) pathway: von Hippelâ€Lindau disease and HIFâ€2α gainâ€ofâ€function mutation. FASEB Journal, 2011, 25, 2001-2011.	0.2	86
149	A Novel Base Change Leading to Hb Vanderbilt [β89(F5)Serâ†'Arg, AG <i>T</i> >AG <i>A</i> ]. Hemoglobin, 2011, 35, 428-429.	0.4	1
150	Increased MSI2 expression Is Associated with Aggressive CML and AML. Blood, 2011, 118, 2516-2516.	0.6	0
151	First Achievements of MPN& MPNr-EuroNet (COST Action BM0902), a New European Network Dedicated to the Diagnosis of Myeloproliferative Neoplasms and Hereditary Erythrocytosis and Thrombocytosis. Blood, 2011, 118, 2809-2809.	0.6	6
152	Two routes to leukemic transformation after a JAK2 mutation–positive myeloproliferative neoplasm. Blood, 2010, 115, 2891-2900.	0.6	269
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