

William Vainchenker

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

154
papers

15,307
citations

65
h-index

123
g-index

164
ext. papers

17,194
ext. citations

5.9
avg, IF

6.04
L-index

#	Paper	IF	Citations
154	Macrophage migration inhibitory factor is overproduced through EGR1 in TET2 resting monocytes.. <i>Communications Biology</i> , 2022 , 5, 110	6.7	0
153	JAK2V617F myeloproliferative neoplasm eradication by a novel interferon/arsenic therapy involves PML. <i>Journal of Experimental Medicine</i> , 2021 , 218,	16.6	7
152	Impact of NFE2 mutations on AML transformation and overall survival in patients with myeloproliferative neoplasms. <i>Blood</i> , 2021 , 138, 2142-2148	2.2	3
151	CALR mutant protein rescues the response of MPL p.R464G variant associated with CAMT to eltrombopag. <i>Blood</i> , 2021 , 138, 480-485	2.2	2
150	Dual role of EZH2 in megakaryocyte differentiation. <i>Blood</i> , 2021 , 138, 1603-1614	2.2	0
149	Germline ATG2B/GSKIP-containing 14q32 duplication predisposes to early clonal hematopoiesis leading to myeloid neoplasms. <i>Leukemia</i> , 2021 ,	10.7	1
148	Induced Pluripotent Stem Cells Enable Disease Modeling and Drug Screening in Calreticulin del52 and ins5 Myeloproliferative Neoplasms. <i>HemaSphere</i> , 2021 , 5, e593	0.3	2
147	Functional Consequences of Mutations in Myeloproliferative Neoplasms. <i>HemaSphere</i> , 2021 , 5, e578	0.3	4
146	CCND2 mutations are infrequent events in BCR-ABL1 negative myeloproliferative neoplasm patients. <i>Haematologica</i> , 2021 , 106, 863-864	6.6	2
145	IFN: Jekyll and Hyde. <i>Blood</i> , 2021 , 137, 291-293	2.2	
144	in acute myeloid leukemia (AML): high prevalence of germline predisposition in French West Indies. <i>Leukemia and Lymphoma</i> , 2021 , 62, 1770-1773	1.9	2
143	Role of Rho-GTPases in megakaryopoiesis. <i>Small GTPases</i> , 2021 , 12, 399-415	2.7	1
142	Inferring the dynamics of mutated hematopoietic stem and progenitor cells induced by IFN in myeloproliferative neoplasms. <i>Blood</i> , 2021 , 138, 2231-2243	2.2	8
141	Lyl-1 regulates primitive macrophages and microglia development. <i>Communications Biology</i> , 2021 , 4, 1382	6.7	1
140	Regulation of Platelet Production and Life Span: Role of Bcl-xL and Potential Implications for Human Platelet Diseases. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	7
139	A p53-JAK-STAT connection involved in myeloproliferative neoplasm pathogenesis and progression to secondary acute myeloid leukemia. <i>Blood Reviews</i> , 2020 , 42, 100712	11.1	7
138	Germline genetic factors in the pathogenesis of myeloproliferative neoplasms. <i>Blood Reviews</i> , 2020 , 42, 100710	11.1	6

137	Different impact of calreticulin mutations on human hematopoiesis in myeloproliferative neoplasms. <i>Oncogene</i> , 2020 , 39, 5323-5337	9.2	7
136	TET2 haploinsufficiency alters reprogramming into induced pluripotent stem cells. <i>Stem Cell Research</i> , 2020 , 44, 101755	1.6	3
135	The Pediatric Acute Leukemia Fusion Oncogene ETO2-GLIS2 Increases Self-Renewal and Alters Differentiation in a Human Induced Pluripotent Stem Cells-Derived Model. <i>HemaSphere</i> , 2020 , 4, e319	0.3	4
134	A new efficient tool for non-invasive diagnosis of fetomaternal platelet antigen incompatibility. <i>British Journal of Haematology</i> , 2020 , 190, 787-798	4.5	2
133	Immunosuppression by Mutated Calreticulin Released from Malignant Cells. <i>Molecular Cell</i> , 2020 , 77, 748-760.e9	17.6	45
132	Calreticulin del52 and ins5 knock-in mice recapitulate different myeloproliferative phenotypes observed in patients with MPN. <i>Nature Communications</i> , 2020 , 11, 4886	17.4	12
131	Megakaryocytes tame erythropoiesis with TGF β . <i>Blood</i> , 2020 , 136, 1016-1017	2.2	1
130	Multilayer intraclonal heterogeneity in chronic myelomonocytic leukemia. <i>Haematologica</i> , 2020 , 105, 112-123	6.6	8
129	Megakaryocyte polyploidization: role in platelet production. <i>Platelets</i> , 2020 , 31, 707-716	3.6	10
128	Knock-in of murine Calr del52 induces essential thrombocythemia with slow-rising dominance in mice and reveals key role of Calr exon 9 in cardiac development. <i>Leukemia</i> , 2020 , 34, 510-521	10.7	23
127	The role of the thrombopoietin receptor MPL in myeloproliferative neoplasms: recent findings and potential therapeutic applications. <i>Expert Review of Hematology</i> , 2019 , 12, 437-448	2.8	14
126	Calreticulin mutants as oncogenic rogue chaperones for TpoR and traffic-defective pathogenic TpoR mutants. <i>Blood</i> , 2019 , 133, 2669-2681	2.2	45
125	Remodeling of Bone Marrow Hematopoietic Stem Cell Niches Promotes Myeloid Cell Expansion during Premature or Physiological Aging. <i>Cell Stem Cell</i> , 2019 , 25, 407-418.e6	18	114
124	Description of a knock-in mouse model of JAK2V617F MPN emerging from a minority of mutated hematopoietic stem cells. <i>Blood</i> , 2019 , 134, 2383-2387	2.2	13
123	Disrupted filamin A/Interaction induces macrothrombocytopenia by increasing RhoA activity. <i>Blood</i> , 2019 , 133, 1778-1788	2.2	19
122	Rare type 1-like and type 2-like calreticulin mutants induce similar myeloproliferative neoplasms as prevalent type 1 and 2 mutants in mice. <i>Oncogene</i> , 2019 , 38, 1651-1660	9.2	5
121	New pathogenic mechanisms induced by germline erythropoietin receptor mutations in primary erythrocytosis. <i>Haematologica</i> , 2018 , 103, 575-586	6.6	9
120	Myelodysplastic Syndromes: Mechanisms, Diagnosis, and Treatment 2018 , 563-563		

119	Secreted Mutant Calreticulins As Rogue Cytokines Trigger Thrombopoietin Receptor Activation Specifically in CALR Mutated Cells: Perspectives for MPN Therapy. <i>Blood</i> , 2018 , 132, 4-4	2.2	24
118	Megakaryocyte and polyploidization. <i>Experimental Hematology</i> , 2018 , 57, 1-13	3.1	42
117	JAK inhibitors for the treatment of myeloproliferative neoplasms and other disorders. <i>F1000Research</i> , 2018 , 7, 82	3.6	83
116	P53 deletion and NrasG12D cooperate for AML. <i>Blood</i> , 2017 , 129, 271-273	2.2	
115	Acquired TET2 mutation in one patient with familial platelet disorder with predisposition to AML led to the development of pre-leukaemic clone resulting in T2-ALL and AML-M0. <i>Journal of Cellular and Molecular Medicine</i> , 2017 , 21, 1237-1242	5.6	8
114	Genetic basis and molecular pathophysiology of classical myeloproliferative neoplasms. <i>Blood</i> , 2017 , 129, 667-679	2.2	275
113	Critical role of the HDAC6-cortactin axis in human megakaryocyte maturation leading to a proplatelet-formation defect. <i>Nature Communications</i> , 2017 , 8, 1786	17.4	28
112	CXCL12/CXCR4 pathway is activated by oncogenic JAK2 in a PI3K-dependent manner. <i>Oncotarget</i> , 2017 , 8, 54082-54095	3.3	22
111	Genetic Alterations of the Thrombopoietin/MPL/JAK2 Axis Impacting Megakaryopoiesis. <i>Frontiers in Endocrinology</i> , 2017 , 8, 234	5.7	23
110	Identification of R102P Mutation in Hereditary Thrombocytosis. <i>Frontiers in Endocrinology</i> , 2017 , 8, 235	5.7	15
109	Downregulation of GATA1 drives impaired hematopoiesis in primary myelofibrosis. <i>Journal of Clinical Investigation</i> , 2017 , 127, 1316-1320	15.9	47
108	Calreticulin mutants in mice induce an MPL-dependent thrombocytosis with frequent progression to myelofibrosis. <i>Blood</i> , 2016 , 127, 1317-24	2.2	178
107	Thrombopoietin receptor activation by myeloproliferative neoplasm associated calreticulin mutants. <i>Blood</i> , 2016 , 127, 1325-35	2.2	204
106	Mutation allele burden remains unchanged in chronic myelomonocytic leukaemia responding to hypomethylating agents. <i>Nature Communications</i> , 2016 , 7, 10767	17.4	140
105	TET2-mediated 5-hydroxymethylcytosine induces genetic instability and mutagenesis. <i>DNA Repair</i> , 2016 , 43, 78-88	4.3	16
104	ATG2B and GSKIP: 2 new genes predisposing to myeloid malignancies. <i>Molecular and Cellular Oncology</i> , 2016 , 3, e1094564	1.2	7
103	P53 activation inhibits all types of hematopoietic progenitors and all stages of megakaryopoiesis. <i>Oncotarget</i> , 2016 , 7, 31980-92	3.3	23
102	Recent advances in understanding myelofibrosis and essential thrombocythemia. <i>F1000Research</i> , 2016 , 5,	3.6	29

101	Uncoupling of the Hippo and Rho pathways allows megakaryocytes to escape the tetraploid checkpoint. <i>Haematologica</i> , 2016 , 101, 1469-1478	6.6	12
100	An incomplete trafficking defect to the cell-surface leads to paradoxical thrombocytosis for human and murine MPL P106L. <i>Blood</i> , 2016 , 128, 3146-3158	2.2	13
99	Activity of nonmuscle myosin II isoforms determines localization at the cleavage furrow of megakaryocytes. <i>Blood</i> , 2016 , 128, 3137-3145	2.2	14
98	Presence of atypical thrombopoietin receptor (MPL) mutations in triple-negative essential thrombocythemia patients. <i>Blood</i> , 2016 , 127, 333-42	2.2	113
97	EZH2: a molecular switch of the MPN phenotype. <i>Blood</i> , 2016 , 127, 3297-8	2.2	2
96	Germline duplication of ATG2B and GSKIP predisposes to familial myeloid malignancies. <i>Nature Genetics</i> , 2015 , 47, 1131-40	36.3	83
95	Concise Review: Induced Pluripotent Stem Cells as New Model Systems in Oncology. <i>Stem Cells</i> , 2015 , 33, 2887-92	5.8	8
94	JAK/STAT Signalling and Haematological Malignancies 2015 , 1-20		
93	Level of RUNX1 activity is critical for leukemic predisposition but not for thrombocytopenia. <i>Blood</i> , 2015 , 125, 930-40	2.2	66
92	TET2 loss, a rescue of JAK2V617F HSCs. <i>Blood</i> , 2015 , 125, 212-3	2.2	1
91	A CALR mutation preceding BCR-ABL1 in an atypical myeloproliferative neoplasm. <i>New England Journal of Medicine</i> , 2015 , 372, 688-90	59.2	35
90	Germ-line JAK2 mutations in the kinase domain are responsible for hereditary thrombocytosis and are resistant to JAK2 and HSP90 inhibitors. <i>Blood</i> , 2014 , 123, 1372-83	2.2	59
89	Acquired initiating mutations in early hematopoietic cells of CLL patients. <i>Cancer Discovery</i> , 2014 , 4, 1088-101	2.2	172
88	TET2 deficiency inhibits mesoderm and hematopoietic differentiation in human embryonic stem cells. <i>Stem Cells</i> , 2014 , 32, 2084-97	5.8	32
87	The formin DIAPH1 (mDia1) regulates megakaryocyte proplatelet formation by remodeling the actin and microtubule cytoskeletons. <i>Blood</i> , 2014 , 124, 3967-77	2.2	49
86	Defective endomitosis during megakaryopoiesis leads to thrombocytopenia in Fanca ^{-/-} mice. <i>Blood</i> , 2014 , 124, 3613-23	2.2	17
85	A new form of macrothrombocytopenia induced by a germ-line mutation in the PRKACG gene. <i>Blood</i> , 2014 , 124, 2554-63	2.2	59
84	JAK2 and MPL protein levels determine TPO-induced megakaryocyte proliferation vs differentiation. <i>Blood</i> , 2014 , 124, 2104-15	2.2	34

83	Emergence of a BCR-ABL translocation in a patient with the JAK2V617F mutation: evidence for secondary acquisition of BCR-ABL in the JAK2V617F clone. <i>Journal of Clinical Oncology</i> , 2014 , 32, e76-9	2.2	16
82	p19 INK4d controls hematopoietic stem cells in a cell-autonomous manner during genotoxic stress and through the microenvironment during aging. <i>Stem Cell Reports</i> , 2014 , 3, 1085-102	8	21
81	Genetic basis of congenital erythrocytosis: mutation update and online databases. <i>Human Mutation</i> , 2014 , 35, 15-26	4.7	82
80	Myeloproliferative neoplasms: JAK2 signaling pathway as a central target for therapy. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2014 , 14 Suppl, S23-35	2	21
79	Thrombocytopenia-associated mutations in the ANKRD26 regulatory region induce MAPK hyperactivation. <i>Journal of Clinical Investigation</i> , 2014 , 124, 580-91	15.9	119
78	Calr Mutants Retroviral Mouse Models Lead to a Myeloproliferative Neoplasm Mimicking an Essential Thrombocythemia Progressing to a Myelofibrosis. <i>Blood</i> , 2014 , 124, 157-157	2.2	8
77	Clonal architecture of chronic myelomonocytic leukemias. <i>Blood</i> , 2013 , 121, 2186-98	2.2	189
76	Prognostic score including gene mutations in chronic myelomonocytic leukemia. <i>Journal of Clinical Oncology</i> , 2013 , 31, 2428-36	2.2	373
75	Combination treatment for myeloproliferative neoplasms using JAK and pan-class I PI3K inhibitors. <i>Journal of Cellular and Molecular Medicine</i> , 2013 , 17, 1397-409	5.6	43
74	Concomitant germ-line RUNX1 and acquired ASXL1 mutations in a T-cell acute lymphoblastic leukemia. <i>European Journal of Haematology</i> , 2013 , 91, 277-9	3.8	21
73	JAK2V617F expression in mice amplifies early hematopoietic cells and gives them a competitive advantage that is hampered by IFN γ . <i>Blood</i> , 2013 , 122, 1464-77	2.2	95
72	Heterozygous and homozygous JAK2(V617F) states modeled by induced pluripotent stem cells from myeloproliferative neoplasm patients. <i>PLoS ONE</i> , 2013 , 8, e74257	3.7	23
71	TET2, a tumor suppressor in hematological disorders. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2012 , 1825, 173-7	11.2	13
70	Thrombopoietin receptor down-modulation by JAK2 V617F: restoration of receptor levels by inhibitors of pathologic JAK2 signaling and of proteasomes. <i>Blood</i> , 2012 , 119, 4625-35	2.2	37
69	Dysmegakaryopoiesis of FPD/AML pedigrees with constitutional RUNX1 mutations is linked to myosin II deregulated expression. <i>Blood</i> , 2012 , 120, 2708-18	2.2	75
68	MYH10 protein expression in platelets as a biomarker of RUNX1 and FLI1 alterations. <i>Blood</i> , 2012 , 120, 2719-22	2.2	61
67	RUNX1-induced silencing of non-muscle myosin heavy chain IIB contributes to megakaryocyte polyploidization. <i>Nature Communications</i> , 2012 , 3, 717	17.4	107
66	Presence of a defect in karyokinesis during megakaryocyte endomitosis. <i>Cell Cycle</i> , 2012 , 11, 4385-9	4.7	18

65	Down-regulation of the RUNX1-target gene NR4A3 contributes to hematopoiesis deregulation in familial platelet disorder/acute myelogenous leukemia. <i>Blood</i> , 2011 , 118, 6310-20	2.2	43
64	Monocytic cells derived from human embryonic stem cells and fetal liver share common differentiation pathways and homeostatic functions. <i>Blood</i> , 2011 , 117, 3065-75	2.2	41
63	Thrombocytopenia resulting from mutations in filamin A can be expressed as an isolated syndrome. <i>Blood</i> , 2011 , 118, 5928-37	2.2	128
62	TET2 inactivation results in pleiotropic hematopoietic abnormalities in mouse and is a recurrent event during human lymphomagenesis. <i>Cancer Cell</i> , 2011 , 20, 25-38	24.3	653
61	Myeloproliferative neoplasms: molecular pathophysiology, essential clinical understanding, and treatment strategies. <i>Journal of Clinical Oncology</i> , 2011 , 29, 573-82	2.2	238
60	New mutations and pathogenesis of myeloproliferative neoplasms. <i>Blood</i> , 2011 , 118, 1723-35	2.2	316
59	FLT3-mediated p38-MAPK activation participates in the control of megakaryopoiesis in primary myelofibrosis. <i>Cancer Research</i> , 2011 , 71, 2901-15	10.1	39
58	Inhibition of TET2-mediated conversion of 5-methylcytosine to 5-hydroxymethylcytosine disturbs erythroid and granulomonocytic differentiation of human hematopoietic progenitors. <i>Blood</i> , 2011 , 118, 2551-5	2.2	139
57	Orientation-specific signalling by thrombopoietin receptor dimers. <i>EMBO Journal</i> , 2011 , 30, 4398-413	13	57
56	A senescence-like cell-cycle arrest occurs during megakaryocytic maturation: implications for physiological and pathological megakaryocytic proliferation. <i>PLoS Biology</i> , 2010 , 8, e1000476	9.7	73
55	Induction of myeloproliferative disorder and myelofibrosis by thrombopoietin receptor W515 mutants is mediated by cytosolic tyrosine 112 of the receptor. <i>Blood</i> , 2010 , 115, 1037-48	2.2	56
54	A major role of TGF-beta1 in the homing capacities of murine hematopoietic stem cell/progenitors. <i>Blood</i> , 2010 , 116, 1244-53	2.2	27
53	Myeloproliferative neoplasm induced by constitutive expression of JAK2V617F in knock-in mice. <i>Blood</i> , 2010 , 116, 783-7	2.2	129
52	Aurora B is dispensable for megakaryocyte polyploidization, but contributes to the endomitotic process. <i>Blood</i> , 2010 , 116, 2345-55	2.2	34
51	Two routes to leukemic transformation after a JAK2 mutation-positive myeloproliferative neoplasm. <i>Blood</i> , 2010 , 115, 2891-900	2.2	224
50	Incidence and prognostic value of TET2 alterations in de novo acute myeloid leukemia achieving complete remission. <i>Blood</i> , 2010 , 116, 1132-5	2.2	105
49	TET2 gene mutation is a frequent and adverse event in chronic myelomonocytic leukemia. <i>Haematologica</i> , 2009 , 94, 1676-81	6.6	198
48	An activating mutation in the CSF3R gene induces a hereditary chronic neutrophilia. <i>Journal of Experimental Medicine</i> , 2009 , 206, 1701-7	16.6	66

47	Mutation in TET2 in myeloid cancers. <i>New England Journal of Medicine</i> , 2009 , 360, 2289-301	59.2	1376
46	Analysis of the ten-eleven translocation 2 (TET2) gene in familial myeloproliferative neoplasms. <i>Blood</i> , 2009 , 114, 1628-32	2.2	84
45	Molecular and genetic bases of myeloproliferative disorders: questions and perspectives. <i>Clinical Lymphoma and Myeloma</i> , 2009 , 9 Suppl 3, S329-39		11
44	Selective reduction of JAK2V617F-dependent cell growth by siRNA/shRNA and its reversal by cytokines. <i>Blood</i> , 2009 , 114, 1842-51	2.2	21
43	A common bipotent progenitor generates the erythroid and megakaryocyte lineages in embryonic stem cell-derived primitive hematopoiesis. <i>Blood</i> , 2009 , 114, 1506-17	2.2	123
42	MAL/SRF complex is involved in platelet formation and megakaryocyte migration by regulating MYL9 (MLC2) and MMP9. <i>Blood</i> , 2009 , 114, 4221-32	2.2	63
41	TET2 mutation is an independent favorable prognostic factor in myelodysplastic syndromes (MDSs). <i>Blood</i> , 2009 , 114, 3285-91	2.2	231
40	The OTT-MAL fusion oncogene activates RBPJ-mediated transcription and induces acute megakaryoblastic leukemia in a knockin mouse model. <i>Journal of Clinical Investigation</i> , 2009 , 119, 852-64 ¹⁵⁻⁹		75
39	JAKs in pathology: role of Janus kinases in hematopoietic malignancies and immunodeficiencies. <i>Seminars in Cell and Developmental Biology</i> , 2008 , 19, 385-93	7.5	137
38	A nonsynonymous SNP in the ITGB3 gene disrupts the conserved membrane-proximal cytoplasmic salt bridge in the alphaIIb beta3 integrin and cosegregates dominantly with abnormal proplatelet formation and macrothrombocytopenia. <i>Blood</i> , 2008 , 111, 3407-14	2.2	81
37	P19INK4D links endomitotic arrest and megakaryocyte maturation and is regulated by AML-1. <i>Blood</i> , 2008 , 111, 4081-91	2.2	44
36	JAK2 stimulates homologous recombination and genetic instability: potential implication in the heterogeneity of myeloproliferative disorders. <i>Blood</i> , 2008 , 112, 1402-12	2.2	140
35	Activating mutations in human acute megakaryoblastic leukemia. <i>Blood</i> , 2008 , 112, 4220-6	2.2	121
34	The hematopoietic stem cell compartment of JAK2V617F-positive myeloproliferative disorders is a reflection of disease heterogeneity. <i>Blood</i> , 2008 , 112, 2429-38	2.2	90
33	Megakaryocyte endomitosis is a failure of late cytokinesis related to defects in the contractile ring and Rho/Rock signaling. <i>Blood</i> , 2008 , 112, 3164-74	2.2	144
32	Evidence for MPL W515L/K mutations in hematopoietic stem cells in primitive myelofibrosis. <i>Blood</i> , 2007 , 110, 3735-43	2.2	84
31	Evidence that the JAK2 G1849T (V617F) mutation occurs in a lymphomyeloid progenitor in polycythemia vera and idiopathic myelofibrosis. <i>Blood</i> , 2007 , 109, 71-7	2.2	135
30	Proplatelet formation is regulated by the Rho/ROCK pathway. <i>Blood</i> , 2007 , 109, 4229-36	2.2	139

29	Interrelation between polyploidization and megakaryocyte differentiation: a gene profiling approach. <i>Blood</i> , 2007 , 109, 3225-34	2.2	85
28	The JAK2 617V>F mutation triggers erythropoietin hypersensitivity and terminal erythroid amplification in primary cells from patients with polycythemia vera. <i>Blood</i> , 2007 , 110, 1013-21	2.2	152
27	Novel activating JAK2 mutation in a patient with Down syndrome and B-cell precursor acute lymphoblastic leukemia. <i>Blood</i> , 2007 , 109, 2202-4	2.2	103
26	New insights into the pathogenesis of JAK2 V617F-positive myeloproliferative disorders and consequences for the management of patients. <i>Seminars in Thrombosis and Hemostasis</i> , 2006 , 32, 341-51 ⁵⁻³		30
25	Deficiency in the Wiskott-Aldrich protein induces premature proplatelet formation and platelet production in the bone marrow compartment. <i>Blood</i> , 2006 , 108, 134-40	2.2	155
24	Monocyte/macrophage dysfunctions do not impair the promotion of myelofibrosis by high levels of thrombopoietin. <i>Journal of Immunology</i> , 2006 , 176, 6425-33	5-3	20
23	Genetic and clinical implications of the Val617Phe JAK2 mutation in 72 families with myeloproliferative disorders. <i>Blood</i> , 2006 , 108, 346-52	2.2	195
22	Reduced retention of radioprotective hematopoietic cells within the bone marrow microenvironment in CXCR4 ^{-/-} chimeric mice. <i>Blood</i> , 2006 , 107, 2243-51	2.2	95
21	An amphipathic motif at the transmembrane-cytoplasmic junction prevents autonomous activation of the thrombopoietin receptor. <i>Blood</i> , 2006 , 107, 1864-71	2.2	115
20	Mammalian target of rapamycin (mTOR) regulates both proliferation of megakaryocyte progenitors and late stages of megakaryocyte differentiation. <i>Blood</i> , 2006 , 107, 2303-10	2.2	76
19	The SCL relative LYL-1 is required for fetal and adult hematopoietic stem cell function and B-cell differentiation. <i>Blood</i> , 2006 , 107, 4678-86	2.2	63
18	JAK2V617F expression in murine hematopoietic cells leads to MPD mimicking human PV with secondary myelofibrosis. <i>Blood</i> , 2006 , 108, 1652-60	2.2	362
17	High molecular response rate of polycythemia vera patients treated with pegylated interferon alpha-2a. <i>Blood</i> , 2006 , 108, 2037-40	2.2	200
16	RGS16 is a negative regulator of SDF-1-CXCR4 signaling in megakaryocytes. <i>Blood</i> , 2005 , 106, 2962-8	2.2	80
15	A unique clonal JAK2 mutation leading to constitutive signalling causes polycythaemia vera. <i>Nature</i> , 2005 , 434, 1144-8	50.4	2769
14	JAK1 and Tyk2 activation by the homologous polycythemia vera JAK2 V617F mutation: cross-talk with IGF1 receptor. <i>Journal of Biological Chemistry</i> , 2005 , 280, 41893-9	5-4	128
13	Mechanisms of WASp-mediated hematologic and immunologic disease. <i>Blood</i> , 2004 , 104, 3454-62	2.2	122
12	Differential regulation of actin stress fiber assembly and proplatelet formation by alpha2beta1 integrin and GPVI in human megakaryocytes. <i>Blood</i> , 2004 , 104, 3117-25	2.2	83

11	FLI1 monoallelic expression combined with its hemizygous loss underlies Paris-Trousseau/Jacobsen thrombopenia. <i>Journal of Clinical Investigation</i> , 2004 , 114, 77-84	15.9	133
10	Megakaryocyte polyploidization is associated with a functional gene amplification. <i>Blood</i> , 2003 , 101, 541-4	2.2	70
9	Prominent role of TGF-beta 1 in thrombopoietin-induced myelofibrosis in mice. <i>Blood</i> , 2002 , 100, 3495-503	5.3	192
8	Distinct effects of thrombopoietin depending on a threshold level of activated Mpl in BaF-3 cells. <i>Journal of Cell Science</i> , 2002 , 115, 2329-2337	5.3	14
7	Asymmetrical segregation of chromosomes with a normal metaphase/anaphase checkpoint in polyploid megakaryocytes. <i>Blood</i> , 2001 , 97, 2238-47	2.2	45
6	Role of p21(Cip1/Waf1) in cell-cycle exit of endomitotic megakaryocytes. <i>Blood</i> , 2001 , 98, 3274-82	2.2	59
5	Phenotypic and Functional Evidence for the Expression of CXCR4 Receptor During Megakaryocytopoiesis. <i>Blood</i> , 1999 , 93, 1511-1523	2.2	105
4	The Thrombocytopenia of Wiskott Aldrich Syndrome Is Not Related to a Defect in Proplatelet Formation. <i>Blood</i> , 1999 , 94, 509-518	2.2	79
3	Effects of Cytokines on Platelet Production From Blood and Marrow CD34+ Cells. <i>Blood</i> , 1998 , 91, 830-843	2.2	114
2	Endomitosis of Human Megakaryocytes Are Due to Abortive Mitosis. <i>Blood</i> , 1998 , 91, 3711-3723	2.2	147
1	High Thrombopoietin Production by Hematopoietic Cells Induces a Fatal Myeloproliferative Syndrome in Mice. <i>Blood</i> , 1997 , 90, 4369-4383	2.2	196