

Jaroslav P Maciejewski

List of Publications by Citations

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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.

314 papers	7,694 citations	37 h-index	85 g-index
328 ext. papers	9,848 ext. citations	5 avg, IF	5.59 L-index

#	Paper	IF	Citations
314	Revised international prognostic scoring system for myelodysplastic syndromes. <i>Blood</i> , 2012 , 120, 2454-65	65	1799
313	Somatic STAT3 mutations in large granular lymphocytic leukemia. <i>New England Journal of Medicine</i> , 2012 , 366, 1905-13	59.2	535
312	The pathophysiology of acquired aplastic anemia. <i>New England Journal of Medicine</i> , 1997 , 336, 1365-72	59.2	417
311	Somatic Mutations and Clonal Hematopoiesis in Aplastic Anemia. <i>New England Journal of Medicine</i> , 2015 , 373, 35-47	59.2	361
310	STAT3 mutations unify the pathogenesis of chronic lymphoproliferative disorders of NK cells and T-cell large granular lymphocyte leukemia. <i>Blood</i> , 2012 , 120, 3048-57	2.2	285
309	Dynamics of clonal evolution in myelodysplastic syndromes. <i>Nature Genetics</i> , 2017 , 49, 204-212	36.3	228
308	Inherited and Somatic Defects in DDX41 in Myeloid Neoplasms. <i>Cancer Cell</i> , 2015 , 27, 658-70	24.3	228
307	In-vivo dominant immune responses in aplastic anaemia: molecular tracking of putatively pathogenetic T-cell clones by TCR beta-CDR3 sequencing. <i>Lancet, The</i> , 2004 , 364, 355-64	40	192
306	Genetic abnormalities in myelodysplasia and secondary acute myeloid leukemia: impact on outcome of stem cell transplantation. <i>Blood</i> , 2017 , 129, 2347-2358	2.2	184
305	Genetic alterations of the cohesin complex genes in myeloid malignancies. <i>Blood</i> , 2014 , 124, 1790-8	2.2	151
304	Wild-type and mutated IDH1/2 enzymes and therapy responses. <i>Oncogene</i> , 2018 , 37, 1949-1960	9.2	127
303	Application of array-based whole genome scanning technologies as a cytogenetic tool in haematological malignancies. <i>British Journal of Haematology</i> , 2009 , 146, 479-88	4.5	118
302	Whole genome scanning as a cytogenetic tool in hematologic malignancies. <i>Blood</i> , 2008 , 112, 965-74	2.2	116
301	Radioprotection of IDH1-Mutated Cancer Cells by the IDH1-Mutant Inhibitor AGI-5198. <i>Cancer Research</i> , 2015 , 75, 4790-802	10.1	108
300	Dose-dependent role of the cohesin complex in normal and malignant hematopoiesis. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1819-32	16.6	106
299	The driver and passenger effects of isocitrate dehydrogenase 1 and 2 mutations in oncogenesis and survival prolongation. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2014 , 1846, 326-41	11.2	93
298	SNP array-based karyotyping: differences and similarities between aplastic anemia and hypocellular myelodysplastic syndromes. <i>Blood</i> , 2011 , 117, 6876-84	2.2	88

297	STAT3 mutations indicate the presence of subclinical T-cell clones in a subset of aplastic anemia and myelodysplastic syndrome patients. <i>Blood</i> , 2013 , 122, 2453-9	2.2	87
296	Deep sequencing reveals stepwise mutation acquisition in paroxysmal nocturnal hemoglobinuria. <i>Journal of Clinical Investigation</i> , 2014 , 124, 4529-38	15.9	87
295	Somatic Mutations in MDS Patients Are Associated with Clinical Features and Predict Prognosis Independent of the IPSS-R: Analysis of Combined Datasets from the International Working Group for Prognosis in MDS-Molecular Committee. <i>Blood</i> , 2015 , 126, 907-907	2.2	73
294	SF3B1-mutant MDS as a distinct disease subtype: a proposal from the International Working Group for the Prognosis of MDS. <i>Blood</i> , 2020 , 136, 157-170	2.2	72
293	Targeting the MALAT1/PARP1/LIG3 complex induces DNA damage and apoptosis in multiple myeloma. <i>Leukemia</i> , 2018 , 32, 2250-2262	10.7	70
292	Loss of Tifab, a del(5q) MDS gene, alters hematopoiesis through derepression of Toll-like receptor-TRAF6 signaling. <i>Journal of Experimental Medicine</i> , 2015 , 212, 1967-85	16.6	65
291	Evolution of clonal cytogenetic abnormalities in aplastic anemia. <i>Leukemia and Lymphoma</i> , 2004 , 45, 433-40	1.9	63
290	Genetic and environmental effects in paroxysmal nocturnal hemoglobinuria: this little PIG-A goes "Why? Why? Why?". <i>Journal of Clinical Investigation</i> , 2000 , 106, 637-41	15.9	63
289	Tet2 loss leads to hypermutagenicity in haematopoietic stem/progenitor cells. <i>Nature Communications</i> , 2017 , 8, 15102	17.4	61
288	The analysis of clonal diversity and therapy responses using STAT3 mutations as a molecular marker in large granular lymphocytic leukemia. <i>Haematologica</i> , 2015 , 100, 91-9	6.6	61
287	Leukemogenic nucleophosmin mutation disrupts the transcription factor hub that regulates granulomonocytic fates. <i>Journal of Clinical Investigation</i> , 2018 , 128, 4260-4279	15.9	61
286	Ubiquitination of hnRNPA1 by TRAF6 links chronic innate immune signaling with myelodysplasia. <i>Nature Immunology</i> , 2017 , 18, 236-245	19.1	60
285	Mutations in G protein β subunits promote transformation and kinase inhibitor resistance. <i>Nature Medicine</i> , 2015 , 21, 71-5	50.5	60
284	Mutations Sensitize Acute Myeloid Leukemia to PARP Inhibition and This Is Reversed by IDH1/2-Mutant Inhibitors. <i>Clinical Cancer Research</i> , 2018 , 24, 1705-1715	12.9	53
283	High incidence of activating STAT5B mutations in CD4-positive T-cell large granular lymphocyte leukemia. <i>Blood</i> , 2016 , 128, 2465-2468	2.2	51
282	Deep sequencing of the T-cell receptor repertoire in CD8+ T-large granular lymphocyte leukemia identifies signature landscapes. <i>Blood</i> , 2013 , 122, 4077-85	2.2	50
281	Clonal drift demonstrates unexpected dynamics of the T-cell repertoire in T-large granular lymphocyte leukemia. <i>Blood</i> , 2011 , 118, 4384-93	2.2	50
280	Myeloid malignancies with chromosome 5q deletions acquire a dependency on an intrachromosomal NF- κ B gene network. <i>Cell Reports</i> , 2014 , 8, 1328-38	10.6	46

279	Clinical features and treatment outcomes in large granular lymphocytic leukemia (LGLL). <i>Leukemia and Lymphoma</i> , 2018 , 59, 416-422	1.9	42
278	mutations define a specific subgroup of MDS and MDS/MPN patients with favorable outcomes with intensive chemotherapy. <i>Blood Advances</i> , 2019 , 3, 922-933	7.8	39
277	Molecular landscape and clonal architecture of adult myelodysplastic/myeloproliferative neoplasms. <i>Blood</i> , 2020 , 136, 1851-1862	2.2	34
276	Hematopoietic stem cells in aplastic anemia. <i>Archives of Medical Research</i> , 2003 , 34, 520-7	6.6	34
275	Transfer of glycosylphosphatidylinositol-anchored proteins to deficient cells after erythrocyte transfusion in paroxysmal nocturnal hemoglobinuria. <i>Blood</i> , 2004 , 104, 3782-8	2.2	34
274	Immune-mediated bone marrow failure syndromes of progenitor and stem cells: molecular analysis of cytotoxic T cell clones. <i>Folia Histochemica Et Cytobiologica</i> , 2007 , 45, 5-14	1.4	34
273	Origins of myelodysplastic syndromes after aplastic anemia. <i>Blood</i> , 2017 , 130, 1953-1957	2.2	32
272	Therapeutic implications of variable expression of CD52 on clonal cytotoxic T cells in CD8+ large granular lymphocyte leukemia. <i>Haematologica</i> , 2009 , 94, 1407-14	6.6	32
271	The efficacy of current prognostic models in predicting outcome of patients with myelodysplastic syndromes at the time of hypomethylating agent failure. <i>Haematologica</i> , 2016 , 101, e224-7	6.6	30
270	Consequences of mutant TET2 on clonality and subclonal hierarchy. <i>Leukemia</i> , 2018 , 32, 1751-1761	10.7	30
269	Invariant patterns of clonal succession determine specific clinical features of myelodysplastic syndromes. <i>Nature Communications</i> , 2019 , 10, 5386	17.4	29
268	Rational management approach to pure red cell aplasia. <i>Haematologica</i> , 2018 , 103, 221-230	6.6	29
267	DDX41-related myeloid neoplasia. <i>Seminars in Hematology</i> , 2017 , 54, 94-97	4	28
266	Baseline clinical characteristics and disease burden in patients with paroxysmal nocturnal hemoglobinuria (PNH): updated analysis from the International PNH Registry. <i>Annals of Hematology</i> , 2020 , 99, 1505-1514	3	27
265	GATA4 loss of function in liver cancer impedes precursor to hepatocyte transition. <i>Journal of Clinical Investigation</i> , 2017 , 127, 3527-3542	15.9	26
264	Mutational landscape of myelodysplastic/myeloproliferative neoplasm-unclassifiable. <i>Blood</i> , 2018 , 132, 2100-2103	2.2	26
263	Novel therapeutic strategies to target leukemic cells that hijack compartmentalized continuous hematopoietic stem cell niches. <i>Biochimica Et Biophysica Acta: Reviews on Cancer</i> , 2017 , 1868, 183-198	11.2	25
262	Phenotypic differences between healthy effector CTL and leukemic LGL cells support the notion of antigen-triggered clonal transformation in T-LGL leukemia. <i>Journal of Leukocyte Biology</i> , 2008 , 83, 589-601	6.5	25

261	Recurrent genetic defects on chromosome 5q in myeloid neoplasms. <i>Oncotarget</i> , 2017 , 8, 6483-6495	3.3	24
260	Germline loss-of-function and alterations in adult myelodysplastic syndromes. <i>Blood</i> , 2018 , 132, 2309-2313	2.3	23
259	Mutations in DNMT3A, U2AF1, and EZH2 identify intermediate-risk acute myeloid leukemia patients with poor outcome after CR1. <i>Blood Cancer Journal</i> , 2018 , 8, 4	7	21
258	Machine learning demonstrates that somatic mutations imprint invariant morphologic features in myelodysplastic syndromes. <i>Blood</i> , 2020 , 136, 2249-2262	2.2	21
257	Immune pathophysiology of aplastic anemia. <i>International Journal of Hematology</i> , 2002 , 76 Suppl 1, 207-14	14	20
256	The mutational burden of therapy-related myeloid neoplasms is similar to primary myelodysplastic syndrome but has a distinctive distribution. <i>Leukemia</i> , 2019 , 33, 2842-2853	10.7	19
255	The Revised International Prognostic Scoring System (IPSS-R) is not predictive of survival in patients with secondary myelodysplastic syndromes. <i>Leukemia and Lymphoma</i> , 2015 , 56, 3437-9	1.9	19
254	Targeting of CD38 by the Tumor Suppressor miR-26a Serves as a Novel Potential Therapeutic Agent in Multiple Myeloma. <i>Cancer Research</i> , 2020 , 80, 2031-2044	10.1	19
253	Decitabine- and 5-azacytidine resistance emerges from adaptive responses of the pyrimidine metabolism network. <i>Leukemia</i> , 2021 , 35, 1023-1036	10.7	19
252	Tet2 Regulates Osteoclast Differentiation by Interacting with Runx1 and Maintaining Genomic 5-Hydroxymethylcytosine (5hmC). <i>Genomics, Proteomics and Bioinformatics</i> , 2018 , 16, 172-186	6.5	18
251	Genomic patterns associated with hypoplastic compared to hyperplastic myelodysplastic syndromes. <i>Haematologica</i> , 2015 , 100, e434-7	6.6	18
250	Aberrant DNA Methylation Is Associated with a Poor Outcome in Juvenile Myelomonocytic Leukemia. <i>PLoS ONE</i> , 2015 , 10, e0145394	3.7	18
249	Invariant phenotype and molecular association of biallelic mutant myeloid neoplasia. <i>Blood Advances</i> , 2019 , 3, 339-349	7.8	18
248	Molecular pathogenesis of disease progression in MLL-rearranged AML. <i>Leukemia</i> , 2019 , 33, 612-624	10.7	18
247	Mutation clonal burden and allogeneic hematopoietic cell transplantation outcomes in acute myeloid leukemia and myelodysplastic syndromes. <i>Bone Marrow Transplantation</i> , 2019 , 54, 1281-1286	4.4	17
246	Human erythroleukemia genetics and transcriptomes identify master transcription factors as functional disease drivers. <i>Blood</i> , 2020 , 136, 698-714	2.2	16
245	ETV6 and signaling gene mutations are associated with secondary transformation of myelodysplastic syndromes to chronic myelomonocytic leukemia. <i>Blood</i> , 2014 , 123, 3675-7	2.2	16
244	Myb expression is critical for myeloid leukemia development induced by Setbp1 activation. <i>Oncotarget</i> , 2016 , 7, 86300-86312	3.3	16

243	Therapy-related acute lymphoblastic leukemia is a distinct entity with adverse genetic features and clinical outcomes. <i>Blood Advances</i> , 2019 , 3, 4228-4237	7.8	16
242	Large granular lymphocytic leukemia coexists with myeloid clones and myelodysplastic syndrome. <i>Leukemia</i> , 2020 , 34, 957-962	10.7	16
241	Impact of SNP array karyotyping on the diagnosis and the outcome of chronic myelomonocytic leukemia with low risk cytogenetic features or no metaphases. <i>American Journal of Hematology</i> , 2016 , 91, 185-92	7.1	15
240	Synergistic effect of major histocompatibility complex class I-related chain a and human leukocyte antigen-DPB1 mismatches in association with acute graft-versus-host disease after unrelated donor hematopoietic stem cell transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2014 , 20, 1835-40	4.7	14
239	Personalized Prediction Model to Risk Stratify Patients With Myelodysplastic Syndromes. <i>Journal of Clinical Oncology</i> , 2021 , 39, 3737-3746	2.2	14
238	Molecular features of early onset adult myelodysplastic syndrome. <i>Haematologica</i> , 2017 , 102, 1028-1034	6.6	13
237	Context dependent effects of ascorbic acid treatment in TET2 mutant myeloid neoplasia. <i>Communications Biology</i> , 2020 , 3, 493	6.7	13
236	Fanconi Anemia germline variants as susceptibility factors in aplastic anemia, MDS and AML. <i>Oncotarget</i> , 2018 , 9, 2050-2057	3.3	12
235	Vacuolization of hematopoietic precursors: an enigma with multiple etiologies. <i>Blood</i> , 2021 , 137, 3685-3689	6.9	12
234	Subclonal STAT3 mutations solidify clonal dominance. <i>Blood Advances</i> , 2019 , 3, 917-921	7.8	12
233	Distinct clinical and biological implications of in myeloid neoplasms. <i>Blood Advances</i> , 2019 , 3, 2164-2178	7.8	12
232	Effectiveness of eculizumab in patients with paroxysmal nocturnal hemoglobinuria (PNH) with or without aplastic anemia in the International PNH Registry. <i>American Journal of Hematology</i> , 2019 , 94, E37-E41	7.1	12
231	Complex landscape of alternative splicing in myeloid neoplasms. <i>Leukemia</i> , 2021 , 35, 1108-1120	10.7	12
230	The functional mechanisms of mutations in myelodysplastic syndrome. <i>Leukemia</i> , 2019 , 33, 2779-2794	10.7	11
229	Impact of allogeneic hematopoietic cell transplant in patients with myeloid neoplasms carrying spliceosomal mutations. <i>American Journal of Hematology</i> , 2016 , 91, 406-9	7.1	11
228	Clinical implications of somatic mutations in aplastic anemia and myelodysplastic syndrome in genomic age. <i>Hematology American Society of Hematology Education Program</i> , 2017 , 2017, 66-72	3.1	10
227	Molecular pathogenesis of myelodysplastic syndromes. <i>Translational Medicine @ UniSa</i> , 2014 , 8, 19-30	0.5	10
226	Genomics of therapy-related myeloid neoplasms. <i>Haematologica</i> , 2020 , 105, e98-e101	6.6	10

225	Somatic mutations in lymphocytes in patients with immune-mediated aplastic anemia. <i>Leukemia</i> , 2021 , 35, 1365-1379	10.7	10
224	Connect MDS/AML: design of the myelodysplastic syndromes and acute myeloid leukemia disease registry, a prospective observational cohort study. <i>BMC Cancer</i> , 2016 , 16, 652	4.8	10
223	Whole-exome sequencing enhances prognostic classification of myeloid malignancies. <i>Journal of Biomedical Informatics</i> , 2015 , 58, 104-113	10.2	9
222	Idiopathic aplastic anemia vs hypocellular myelodysplastic syndrome. <i>Hematology American Society of Hematology Education Program</i> , 2019 , 2019, 97-104	3.1	9
221	and mutations in myelodysplastic syndromes (MDS): clonal architecture and impact on outcomes. <i>Leukemia and Lymphoma</i> , 2019 , 60, 1587-1590	1.9	9
220	The evolution of paroxysmal nocturnal haemoglobinuria depends on intensity of immunosuppressive therapy. <i>British Journal of Haematology</i> , 2018 , 182, 730-733	4.5	9
219	A Personalized Prediction Model to Risk Stratify Patients with Acute Myeloid Leukemia (AML) Using Artificial Intelligence. <i>Blood</i> , 2019 , 134, 2091-2091	2.2	8
218	Extended experience with a non-cytotoxic DNMT1-targeting regimen of decitabine to treat myeloid malignancies. <i>British Journal of Haematology</i> , 2020 , 188, 924-929	4.5	8
217	From Bench to Bedside and Beyond: Therapeutic Scenario in Acute Myeloid Leukemia. <i>Cancers</i> , 2020 , 12,	6.6	7
216	TP53 Mutations and Outcome in Patients with Myelodysplastic Syndromes (MDS). <i>Blood</i> , 2016 , 128, 4336-4336	4.3	7
215	Gene-centric functional dissection of human genetic variation uncovers regulators of hematopoiesis. <i>ELife</i> , 2019 , 8,	8.9	7
214	How I manage acquired pure red cell aplasia in adults. <i>Blood</i> , 2021 , 137, 2001-2009	2.2	7
213	New drugs for pharmacological extension of replicative life span in normal and progeroid cells. <i>Npj Aging and Mechanisms of Disease</i> , 2019 , 5, 2	5.5	7
212	Phase 2 study of danicopan in patients with paroxysmal nocturnal hemoglobinuria with an inadequate response to eculizumab. <i>Blood</i> , 2021 , 138, 1928-1938	2.2	7
211	Promoter Methylation Is Linked to Defective Homologous Recombination Repair and Elevated to Disrupt Myeloid Differentiation in Myeloid Malignancies. <i>Clinical Cancer Research</i> , 2019 , 25, 2513-2522	12.9	6
210	Therapeutic outcomes using subcutaneous low dose alemtuzumab for acquired bone marrow failure conditions. <i>British Journal of Haematology</i> , 2018 , 183, 133-136	4.5	6
209	Clonal PIGA mosaicism and dynamics in paroxysmal nocturnal hemoglobinuria. <i>Leukemia</i> , 2018 , 32, 2507-2511	12.1	6
208	Germline DDX41 mutations cause ineffective hematopoiesis and myelodysplasia. <i>Cell Stem Cell</i> , 2021 , 28, 1966-1981.e6	18	6

207	Impact of germline CTC1 alterations on telomere length in acquired bone marrow failure. <i>British Journal of Haematology</i> , 2019 , 185, 935-939	4.5	5
206	A Study Comparing Dosing Regimens and Efficacy of Subcutaneous to Intravenous Azacitidine (AZA) for the Treatment of Myelodysplastic Syndromes (MDS).. <i>Blood</i> , 2009 , 114, 3797-3797	2.2	5
205	Clonal Mutational Landscape of Childhood Myelodysplastic Syndromes. <i>Blood</i> , 2015 , 126, 1662-1662	2.2	5
204	Distinctive and common features of moderate aplastic anaemia. <i>British Journal of Haematology</i> , 2020 , 189, 967-975	4.5	4
203	Genetic and Epigenetic Defects in DNA Repair Lead to Synthetic Lethality of Poly (ADP-Ribose) Polymerase (PARP) Inhibitors in Aggressive Myeloproliferative Disorders. <i>Blood</i> , 2011 , 118, 400-400	2.2	4
202	The Mechanism By Which Mutant Nucleophosmin (NPM1) Creates Leukemic Self-Renewal Is Readily Reversed. <i>Blood</i> , 2016 , 128, 444-444	2.2	4
201	Functional analyses of human LUC7-like proteins involved in splicing regulation and myeloid neoplasms. <i>Cell Reports</i> , 2021 , 35, 108989	10.6	4
200	Novel invariant features of Good syndrome. <i>Leukemia</i> , 2021 , 35, 1792-1796	10.7	4
199	Transcriptomic rationale for synthetic lethality-targeting ERCC1 and CDKN1A in chronic myelomonocytic leukaemia. <i>British Journal of Haematology</i> , 2018 , 182, 373-383	4.5	4
198	TET-dioxygenase deficiency in oncogenesis and its targeting for tumor-selective therapeutics. <i>Seminars in Hematology</i> , 2021 , 58, 27-34	4	4
197	Large granular lymphocytic leukaemia after solid organ and haematopoietic stem cell transplantation. <i>British Journal of Haematology</i> , 2020 , 189, 318-322	4.5	3
196	Effects of the Therapeutic Armamentarium on Survival and Time to Next Treatment in CMML Subtypes: An International Analysis of 950 Cases Coordinated By the AGMT Study Group. <i>Blood</i> , 2019 , 134, 844-844	2.2	3
195	Combined Treatment with Lenalidomide and Epoetin Alfa Leads to Durable Responses in Patients with Epo-Refractory, Lower Risk Non-Deletion 5q [Del(5q)] MDS: Final Results of the E2905 Intergroup Phase III Study - an ECOG-ACRIN Cancer Research Group Study, Grant CA180820, and the National Cancer Institute of the National Institutes of Health. <i>Blood</i> , 2019 , 134, 842-842	2.2	3
194	High Density SNP Arrays Reveal That Distinct Clonal Lesions Including Uniparental Disomy Can Be Detected in a Proportion of Patients with Aplastic Anemia with Normal Metaphase Cytogenetics.. <i>Blood</i> , 2006 , 108, 125-125	2.2	3
193	Non-Cytotoxic Differentiation Therapy Based On Mechanism of Disease Produces Complete Remission in Myelodysplastic Syndromes (MDS) with High Risk Cytogenetics. <i>Blood</i> , 2012 , 120, 1696-1696	2.2	3
192	The Role of LUC7L2 in Splicing and MDS. <i>Blood</i> , 2016 , 128, 5504-5504	2.2	3
191	Rare germline variant contributions to myeloid malignancy susceptibility. <i>Leukemia</i> , 2020 , 34, 1675-1678	10.7	3
190	Machine learning integrates genomic signatures for subclassification beyond primary and secondary acute myeloid leukemia. <i>Blood</i> , 2021 , 138, 1885-1895	2.2	3

189	Frequency and perturbations of various peripheral blood cell populations before and after eculizumab treatment in paroxysmal nocturnal hemoglobinuria. <i>Blood Cells, Molecules, and Diseases</i> , 2021 , 87, 102528	2.1	3
188	Implication of PIGA genotype on erythrocytes phenotype in Paroxysmal Nocturnal Hemoglobinuria. <i>Leukemia</i> , 2021 , 35, 2431-2434	10.7	3
187	Introduction: molecular pathogenesis of hematologic malignancies. <i>Seminars in Oncology</i> , 2012 , 39, 9-12	5.5	2
186	A case of mistaken identity: When lupus masquerades as primary myelofibrosis. <i>SAGE Open Medical Case Reports</i> , 2013 , 1, 2050313X13498709	0.7	2
185	Geno-Clinical Model for the Diagnosis of Bone Marrow Myeloid Neoplasms. <i>Blood</i> , 2019 , 134, 4238-4238	2.2	2
184	TET Dioxygenase Inhibition As a Therapeutic Strategy in TET2 Mutant Myeloid Neoplasia. <i>Blood</i> , 2019 , 134, 880-880	2.2	2
183	A Personalized Clinical-Decision Tool to Improve the Diagnostic Accuracy of Myelodysplastic Syndromes. <i>Blood</i> , 2020 , 136, 33-35	2.2	2
182	Clinical Impacts of Germline DDX41 Mutations on Myeloid Neoplasms. <i>Blood</i> , 2020 , 136, 38-40	2.2	2
181	A Decision Analysis To Determine the Appropriate Treatment for Low-Risk Myelodysplastic Syndromes.. <i>Blood</i> , 2005 , 106, 2533-2533	2.2	2
180	SNP Array-Based Analysis of Chromosome 17 Reveals Biallelic TP53 Mutations Due to Uniparental Disomy 17p in Advanced MDS and AML with Cooperating Deletions of Chromosomes 5 and 7. <i>Blood</i> , 2008 , 112, 2521-2521	2.2	2
179	Association of SF3B1 with Ring Sideroblasts in patients, In Vivo, and In Vitro models of Spliceosomal Dysfunction. <i>Blood</i> , 2011 , 118, 457-457	2.2	2
178	Survival Outcomes of Leukemias and Myelodysplastic Syndromes Occurring As Second Cancers in the United States: A SEER Registry-Based Population Analysis. <i>Blood</i> , 2015 , 126, 2507-2507	2.2	2
177	Genetic and Epigenetic Defects in the Autophagy Machinery in Myelodysplastic Syndromes. <i>Blood</i> , 2016 , 128, 4301-4301	2.2	2
176	The Similarity of Class II HLA Genotypes Defines Patterns of Autoreactivity in Idiopathic Bone Marrow Failure Disorders. <i>Blood</i> , 2021 ,	2.2	2
175	Is nature truly healing itself? Spontaneous remissions in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood Cancer Journal</i> , 2021 , 11, 187	7	2
174	Extent and Clinical Implications of Subclonal Diversity in Paroxysmal Nocturnal Hemoglobinuria. <i>Blood</i> , 2017 , 130, 779-779	2.2	2
173	Expression of MICA by Granulocytes in Neutropenia Due to Large Granular Lymphocyte Leukemia Points towards Cytotoxicity Exerted Via NKG2D on Clonal Cytotoxic T Cells.. <i>Blood</i> , 2008 , 112, 1262-1262	2.2	2
172	Therapeutic Targeting of Protein Disulfide Isomerase PDIA1 in Multiple Myeloma. <i>Cancers</i> , 2021 , 13,	6.6	2

171	Analysis of distinct hotspot mutations in relation to clinical phenotypes and response to therapy in myeloid neoplasia. <i>Leukemia and Lymphoma</i> , 2021 , 62, 735-738	1.9	2
170	A Phase II Trial of Imatinib Mesylate as Maintenance Therapy for Patients With Newly Diagnosed C-kit-positive Acute Myeloid Leukemia. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021 , 21, 113-118	2	2
169	Clinical and basic implications of dynamic T cell receptor clonotyping in hematopoietic cell transplantation. <i>JCI Insight</i> , 2021 , 6,	9.9	2
168	A geno-clinical decision model for the diagnosis of myelodysplastic syndromes. <i>Blood Advances</i> , 2021 , 5, 4361-4369	7.8	2
167	Large Granular Lymphocytic Leukemia: From Immunopathogenesis to Treatment of Refractory Disease. <i>Cancers</i> , 2021 , 13,	6.6	2
166	Circulating microbial content in myeloid malignancy patients is associated with disease subtypes and patient outcomes.. <i>Nature Communications</i> , 2022 , 13, 1038	17.4	2
165	Single-cell characterization of leukemic and non-leukemic immune repertoires in CD8 T-cell large granular lymphocytic leukemia.. <i>Nature Communications</i> , 2022 , 13, 1981	17.4	2
164	Chronic myeloid leukemia: Two mysteries. <i>Leukemia Research</i> , 2019 , 79, 3-5	2.7	1
163	5-formylcytosine and 5-hydroxymethyluracil as surrogate markers of TET2 and SF3B1 mutations in myelodysplastic syndrome, respectively. <i>Haematologica</i> , 2020 , 105, e213-e215	6.6	1
162	Distinct mutational pattern of myelodysplastic syndromes with and without 5q- treated with lenalidomide. <i>British Journal of Haematology</i> , 2020 , 189, e133-e137	4.5	1
161	Genotype-Phenotype Relationships and Therapeutic Targets in Acute Erythroid Leukemia. <i>Blood</i> , 2020 , 136, 17-18	2.2	1
160	A Phase I/II Trial of CPX-351 + Palbociclib in Patients with Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 13-14	2.2	1
159	Aberrant Telomere Length and Composition Are Recurrent Features of Myeloid Disorders. <i>Blood</i> , 2020 , 136, 29-30	2.2	1
158	Eltrombopag inhibits TET dioxygenase to contribute to hematopoietic stem cell expansion in aplastic anemia.. <i>Journal of Clinical Investigation</i> , 2022 ,	15.9	1
157	RORA Is a Potential Prognostic Biomarker and Therapeutic Target for Patients with Acute Myeloid Leukemia. <i>Blood</i> , 2019 , 134, 2696-2696	2.2	1
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