

Rhett P Ketterling

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.

462 papers	14,515 citations	57 h-index	110 g-index
465 ext. papers	16,985 ext. citations	4.5 avg, IF	6 L-index

#	Paper	IF	Citations
462	A simple additive staging system for newly diagnosed multiple myeloma.. <i>Blood Cancer Journal</i> , 2022 , 12, 21	7	4
461	Myelodysplastic/myeloproliferative neoplasms with ring sideroblasts and thrombocytosis (MDS/MPN-RS-T): Mayo-Moffitt collaborative study of 158 patients.. <i>Blood Cancer Journal</i> , 2022 , 12, 26	7	1
460	-mutant myelodysplastic syndrome/myeloproliferative neoplasms: a unique molecular and prognostic entity.. <i>Haematologica</i> , 2022 ,	6.6	1
459	Clonal compositions involving epigenetic regulator and splicing mutations in CHIP, CCUS, MDS, and CMML.. <i>Leukemia Research</i> , 2022 , 106818	2.7	0
458	Cytogenetic abnormalities in essential thrombocythemia: Clinical and molecular correlates and prognostic relevance in 809 informative cases.. <i>Blood Cancer Journal</i> , 2022 , 12, 44	7	1
457	Identification of EWSR1 rearrangements in patients with immature hematopoietic neoplasms: A case series and review of literature.. <i>Annals of Diagnostic Pathology</i> , 2022 , 58, 151942	2.2	0
456	Apparent coexistence of and fusions due to a nonproductive rearrangement in B-ALL.. <i>Leukemia and Lymphoma</i> , 2022 , 1-4	1.9	0
455	RNAseq identification of FISH-cryptic BCL6::TP63 rearrangement in ALK-negative anaplastic large cell lymphoma.. <i>Histopathology</i> , 2022 ,	7.3	0
454	Lymphocytopenia predicts shortened survival in myelodysplastic syndrome with ring sideroblasts (MDS-RS) but not in MDS/MPN-RS-T.. <i>American Journal of Hematology</i> , 2021 ,	7.1	2
453	Utilizing next-generation sequencing to characterize a case of acute myeloid leukemia with t(4;12)(q12;p13) in the absence of ETV6/CHIC2 and ETV6/PDGFRα gene fusions. <i>Cancer Genetics</i> , 2021 , 260-261, 1-5	2.3	
452	MYC break-apart FISH probe set reveals frequent unbalanced patterns of uncertain significance when evaluating aggressive B-cell lymphoma. <i>Blood Cancer Journal</i> , 2021 , 11, 184	7	0
451	Anthracycline Choices for Induction Chemotherapy Among 797 Consecutive Adult Patients with Acute Myeloid Leukemia: Daunorubicin-60 Vs Idarubicin-12 Vs Daunorubicin-90. <i>Blood</i> , 2021 , 138, 1267-1267	12.2	
450	Cytogenetics in Essential Thrombocythemia: Phenotype and Molecular Correlates and Prognostic Relevance in 818 Informative Cases. <i>Blood</i> , 2021 , 138, 3629-3629	2.2	
449	Characterization of Atypical t(11;14) CCND1/IGH Translocations in Multiple Myeloma. <i>Blood</i> , 2021 , 138, 3771-3771	2.2	0
448	Acute Myeloid Leukemia in the Context of Previous History of Cancer with or without Exposure to Chemotherapy or Radiotherapy. <i>Blood</i> , 2021 , 138, 3368-3368	2.2	0
447	Cryptic t(6;11) KMT2A rearrangement in a pediatric acute myeloid leukemia patient detected by next-generation sequencing and dual-fusion FISH analysis. <i>Pediatric Blood and Cancer</i> , 2021 , e29428	3	
446	False-Negative Centromere 15 Probe Results in Association with African Ancestry in Plasma Cell Dyscrasias. <i>Blood</i> , 2021 , 138, 4101-4101	2.2	

445	Conventional Cytogenetic Analysis of Hematologic Neoplasms: A 20-Year Review of Proficiency Test Results From the College of American Pathologists/American College of Medical Genetics and Genomics Cytogenetics Committee. <i>Archives of Pathology and Laboratory Medicine</i> , 2021 , 145, 176-190	5	2
444	Striking Association of Lymphoid Enhancing Factor (LEF1) Overexpression and DUSP22 Rearrangements in Anaplastic Large Cell Lymphoma. <i>American Journal of Surgical Pathology</i> , 2021 , 45, 550-557	6.7	3
443	Acute myeloid leukemia after age 70 years: A retrospective comparison of survival following treatment with intensive versus HMA + venetoclax chemotherapy. <i>American Journal of Hematology</i> , 2021 , 96, E108-E111	7.1	3
442	Mayo Clinic experience with 1123 adults with acute myeloid leukemia. <i>Blood Cancer Journal</i> , 2021 , 11, 46	7	1
441	CSF3R T618I mutant chronic myelomonocytic leukemia (CMML) defines a proliferative CMML subtype enriched in ASXL1 mutations with adverse outcomes. <i>Blood Cancer Journal</i> , 2021 , 11, 54	7	1
440	Genomic stratification of myelodysplastic/myeloproliferative neoplasms, unclassifiable: Sorting through the unsorted. <i>Leukemia</i> , 2021 , 35, 3329-3333	10.7	1
439	Molecular classification improves risk assessment in adult BCR-ABL1-negative B-ALL. <i>Blood</i> , 2021 , 138, 948-958	2.2	7
438	Clinical, molecular, and prognostic comparisons between CCUS and lower-risk MDS: a study of 187 molecularly annotated patients. <i>Blood Advances</i> , 2021 , 5, 2272-2278	7.8	3
437	Increased complexity of t(11;14) rearrangements in plasma cell neoplasms compared with mantle cell lymphoma. <i>Genes Chromosomes and Cancer</i> , 2021 , 60, 678-686	5	1
436	Assessment of isochromosome 12p and 12p abnormalities in germ cell tumors using fluorescence in situ hybridization, single-nucleotide polymorphism arrays, and next-generation sequencing/mate-pair sequencing. <i>Human Pathology</i> , 2021 , 112, 20-34	3.7	2
435	Landscape of RAS pathway mutations in patients with myelodysplastic syndrome/myeloproliferative neoplasm overlap syndromes: a study of 461 molecularly annotated patients. <i>Leukemia</i> , 2021 , 35, 644-649	10.7	6
434	Identification of a novel KMT2A/GIMAP8 gene fusion in a pediatric patient with acute undifferentiated leukemia. <i>Genes Chromosomes and Cancer</i> , 2021 , 60, 108-111	5	1
433	Lymphoma-like double-hit genetic abnormalities (and) in a case of non-secretory multiple myeloma. <i>Leukemia and Lymphoma</i> , 2021 , 62, 243-246	1.9	
432	Clinical correlates and prognostic impact of clonal hematopoiesis in multiple myeloma patients receiving post-autologous stem cell transplantation lenalidomide maintenance therapy. <i>American Journal of Hematology</i> , 2021 , 96, E157-E162	7.1	2
431	Myeloid malignancies with 5q and 7q deletions are associated with extreme genomic complexity, biallelic TP53 variants, and very poor prognosis. <i>Blood Cancer Journal</i> , 2021 , 11, 18	7	3
430	The Prognostic Role of Structural Variants Identified by NGS and FISH in Multiple Myeloma. <i>Clinical Cancer Research</i> , 2021 ,	12.9	5
429	De novo isolated myeloid sarcoma: comparative analysis of survival in 19 consecutive cases. <i>British Journal of Haematology</i> , 2021 , 195, 413-416	4.5	1
428	Clinical utility of next generation sequencing to detect IGH/IL3 rearrangements [t(5;14)(q31.1;q32.1)] in B-lymphoblastic leukemia/lymphoma. <i>Annals of Diagnostic Pathology</i> , 2021 , 53, 151761	2.2	0

427	Spectrum of hematological malignancies, clonal evolution and outcomes in 144 Mayo Clinic patients with germline predisposition syndromes. <i>American Journal of Hematology</i> , 2021 , 96, 1450-1460	7.1	1
426	Identification of adult Philadelphia-like acute lymphoblastic leukemia using a FISH-based algorithm distinguishes prognostic groups and outcomes. <i>Blood Cancer Journal</i> , 2021 , 11, 156	7	3
425	Dual Primary IGH Translocations in Multiple Myeloma: A Novel Finding. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2021 , 21, e710-e713	2	
424	A rare case of atypical chronic myeloid leukemia associated with t(8;22)(p11.2;q11.2)/ BCR-FGFR1 rearrangement: A case report and literature review. <i>Cancer Genetics</i> , 2021 , 258-259, 69-73	2.3	0
423	Detection of cryptic CCND1 rearrangements in mantle cell lymphoma by next generation sequencing. <i>Annals of Diagnostic Pathology</i> , 2020 , 46, 151533	2.2	3
422	Anaplastic large-cell lymphoma (ALK-negative)-related heart failure and recurrence after heart transplantation. <i>Journal of Heart and Lung Transplantation</i> , 2020 , 39, 1156-1158	5.8	0
421	The significance of genetic mutations and their prognostic impact on patients with incidental finding of isolated del(20q) in bone marrow without morphologic evidence of a myeloid neoplasm. <i>Blood Cancer Journal</i> , 2020 , 10, 7	7	5
420	High level MYC amplification in B-cell lymphomas: is it a marker of aggressive disease?. <i>Blood Cancer Journal</i> , 2020 , 10, 5	7	10
419	Genetic Factors in Acute Myeloid Leukemia With Myelodysplasia-Related Changes. <i>American Journal of Clinical Pathology</i> , 2020 , 153, 656-663	1.9	6
418	Mutation-enhanced international prognostic systems for essential thrombocythaemia and polycythaemia vera. <i>British Journal of Haematology</i> , 2020 , 189, 291-302	4.5	58
417	Phenotypic correlates and prognostic outcomes of TET2 mutations in myelodysplastic syndrome/myeloproliferative neoplasm overlap syndromes: A comprehensive study of 504 adult patients. <i>American Journal of Hematology</i> , 2020 , 95, E86-E89	7.1	2
416	Characterizing false-positive fluorescence in situ hybridization results by mate-pair sequencing in a patient with chronic myeloid leukemia and progression to myeloid blast crisis. <i>Cancer Genetics</i> , 2020 , 243, 48-51	2.3	2
415	Myeloid Sarcoma With CBFB-MYH11 Fusion (inv(16) or t(16;16)) Prevails in the Abdomen. <i>American Journal of Clinical Pathology</i> , 2020 , 153, 333-341	1.9	3
414	The Prognostic Significance of Acquired 1q22 Gain in Multiple Myeloma. <i>Blood</i> , 2020 , 136, 9-10	2.2	
413	Heterogeneity of MYC Abnormalities in Multiple Myeloma. <i>Blood</i> , 2020 , 136, 2-3	2.2	
412	Clinical Value of Next Generation Sequencing in the Detection of Recurring Structural Rearrangements and Copy Number Abnormalities in Acute Myeloid Leukemia. <i>Blood</i> , 2020 , 136, 21-22	2.2	
411	Striking Association of Lymphoid Enhancing Factor (LEF1) Overexpression and DUSP22 rearrangements in Anaplastic Large Cell Lymphoma. <i>Blood</i> , 2020 , 136, 22-23	2.2	
410	Metaphase cytogenetics and plasma cell proliferation index for risk stratification in newly diagnosed multiple myeloma. <i>Blood Advances</i> , 2020 , 4, 2236-2244	7.8	7

409	SF3B1-mutant CMML defines a predominantly dysplastic CMML subtype with a superior acute leukemia-free survival. <i>Blood Advances</i> , 2020 , 4, 5716-5721	7.8	5
408	Clinical utility of fluorescence in situ hybridization-based diagnosis of BCR-ABL1 like (Philadelphia chromosome like) B-acute lymphoblastic leukemia. <i>American Journal of Hematology</i> , 2020 , 95, E68-E72	7.1	2
407	Secondary acquisition of BCR-ABL1 fusion in de novo GATA2-MECOM positive acute myeloid leukemia with subsequent emergence of a rare KMT2A-ASXL2 fusion. <i>Cancer Genetics</i> , 2020 , 241, 67-71	2.3	1
406	Molecular and phenotypic characterization of an early T-cell precursor acute lymphoblastic lymphoma harboring PICALM-MLLT10 fusion with aberrant expression of B-cell antigens. <i>Cancer Genetics</i> , 2020 , 240, 40-44	2.3	3
405	Characterization of a cryptic fusion by mate-pair sequencing in a case of acute promyelocytic leukemia with a normal karyotype and negative FISH studies. <i>Leukemia and Lymphoma</i> , 2020 , 61, 975-978	1.9	4
404	Implications of MYC Rearrangements in Newly Diagnosed Multiple Myeloma. <i>Clinical Cancer Research</i> , 2020 , 26, 6581-6588	12.9	9
403	Siblings with ETV6/RUNX1-positive B-lymphoblastic leukemia: A single site experience and review of the literature. <i>Annals of Diagnostic Pathology</i> , 2020 , 48, 151588	2.2	0
402	Clinical characteristics and treatment outcomes of newly diagnosed multiple myeloma with chromosome 1q abnormalities. <i>Blood Advances</i> , 2020 , 4, 3509-3519	7.8	27
401	Cytogenetic abnormalities in multiple myeloma: association with disease characteristics and treatment response. <i>Blood Cancer Journal</i> , 2020 , 10, 82	7	17
400	Fluorescence in-situ hybridisation for TP63 rearrangements in T cell lymphomas: single-site experience of 470 patients and implications for clinical testing. <i>Histopathology</i> , 2020 , 76, 481-485	7.3	3
399	Clinicopathologic characteristics, prognostication and treatment outcomes for myelodysplastic/myeloproliferative neoplasm, unclassifiable (MDS/MPN-U): Mayo Clinic-Moffitt Cancer Center study of 135 consecutive patients. <i>Leukemia</i> , 2020 , 34, 656-661	10.7	17
398	Cryptic and atypical KMT2A-USP2 and KMT2A-USP8 rearrangements identified by mate pair sequencing in infant and childhood leukemia. <i>Genes Chromosomes and Cancer</i> , 2020 , 59, 422-427	5	4
397	Combined Tumors in Hematolymphoid Neoplasms: Case Series of Histiocytic and Langerhans Cell Sarcomas Arising From Low-Grade B-Cell Lymphoma. <i>BMC Clinical Pathology</i> , 2019 , 12, 2632010X19878410	1.3	5
396	Leukemic transformation among 1306 patients with primary myelofibrosis: risk factors and development of a predictive model. <i>Blood Cancer Journal</i> , 2019 , 9, 12	7	28
395	Tetraploidy is associated with poor prognosis at diagnosis in multiple myeloma. <i>American Journal of Hematology</i> , 2019 , 94, E117-E120	7.1	6
394	Characterization of a t(1;2)(p36;p21) involving the PRDM16 gene region by mate-pair sequencing (MPseq) in a patient with newly diagnosed acute myeloid leukemia with myelodysplasia-related changes. <i>Journal of Hematopathology</i> , 2019 , 12, 85-90	0.4	
393	Characterization of a cryptic KMT2A/AFF1 gene fusion by mate-pair sequencing (MPseq) in a young adult with newly diagnosed B-lymphoblastic leukemia. <i>Journal of Hematopathology</i> , 2019 , 12, 99-104	0.4	0
392	Elucidating a false-negative break-apart fluorescence in situ hybridization probe study by next-generation sequencing in a patient with high-grade B-cell lymphoma with and rearrangements. <i>Journal of Physical Education and Sports Management</i> , 2019 , 5,	2.8	8

391	Recurrent mutations in ALK-negative anaplastic large cell lymphoma. <i>Blood</i> , 2019 , 133, 2776-2789	2.2	23
390	Cytogenetic clonal evolution in myeloproliferative neoplasms: contexts and prognostic impact among 648 patients with serial bone marrow biopsies. <i>Leukemia</i> , 2019 , 33, 2522-2553	10.7	1
389	ADDENDUM: Section E9 of the American College of Medical Genetics Technical Standards and Guidelines: Fluorescence in situ hybridization. <i>Genetics in Medicine</i> , 2019 , 21, 2405	8.1	1
388	Natural history of multiple myeloma with de novo del(17p). <i>Blood Cancer Journal</i> , 2019 , 9, 32	7	22
387	Hyperhaploid plasma cell myeloma characterized by poor outcome and monosomy 17 with frequently co-occurring TP53 mutations. <i>Blood Cancer Journal</i> , 2019 , 9, 20	7	6
386	3023 Mayo Clinic Patients With Myeloproliferative Neoplasms: Risk-Stratified Comparison of Survival and Outcomes Data Among Disease Subgroups. <i>Mayo Clinic Proceedings</i> , 2019 , 94, 599-610	6.4	50
385	Suboptimal response rates to hypomethylating agent therapy in chronic myelomonocytic leukemia; a single institutional study of 121 patients. <i>American Journal of Hematology</i> , 2019 , 94, 767-779	7.1	27
384	A near-haploid clone harboring a BCR/ABL1 gene fusion in an adult patient with newly diagnosed B-lymphoblastic leukemia. <i>Genes Chromosomes and Cancer</i> , 2019 , 58, 665-668	5	1
383	Substratification of patients with newly diagnosed standard-risk multiple myeloma. <i>British Journal of Haematology</i> , 2019 , 185, 254-260	4.5	8
382	A prospective evaluation of vitamin B1 (thiamine) level in myeloproliferative neoplasms: clinical correlations and impact of JAK2 inhibitor therapy. <i>Blood Cancer Journal</i> , 2019 , 9, 11	7	8
381	Whole Genome Mate-pair Sequencing of Plasma Cell Neoplasm as a Novel Diagnostic Strategy: A Case of Unrecognized t(2;11) Structural Variation. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2019 , 19, 598-602	2	2
380	Characterization of a rarely reported STAT5B/RARA gene fusion in a young adult with newly diagnosed acute promyelocytic leukemia with resistance to ATRA therapy. <i>Cancer Genetics</i> , 2019 , 237, 51-54	2.3	3
379	Characterization of TCF3 rearrangements in pediatric B-lymphoblastic leukemia/lymphoma by mate-pair sequencing (MPseq) identifies complex genomic rearrangements and a novel TCF3/TEF gene fusion. <i>Blood Cancer Journal</i> , 2019 , 9, 81	7	9
378	Metaphase Cytogenetics for Risk Stratification in Newly Diagnosed Multiple Myeloma. <i>Blood</i> , 2019 , 134, 4396-4396	2.2	
377	Recurrent Chromosomal Abnormalities in Tumoral Lesions of Small Lymphocytic Lymphoma/Chronic Lymphocytic Leukemia: A Large-Scale Fluorescent in-Situ Hybridization Study on Tissue Biopsy Sections. <i>Blood</i> , 2019 , 134, 4282-4282	2.2	
376	A Novel Approach to Risk Stratification in Multiple Myeloma Using ISS Stage and FISH. <i>Blood</i> , 2019 , 134, 1800-1800	2.2	1
375	Differentiating between Hyperdiploidy and Pseudo-Hyperdiploidy in B-Lymphoblastic Leukemia Utilizing Low-Coverage Mate-Pair Sequencing. <i>Blood</i> , 2019 , 134, 5212-5212	2.2	
374	Acute leukemias harboring KMT2A/MLLT10 fusion: a 10-year experience from a single genomics laboratory. <i>Genes Chromosomes and Cancer</i> , 2019 , 58, 567-577	5	11

373	Impact of acquired del(17p) in multiple myeloma. <i>Blood Advances</i> , 2019 , 3, 1930-1938	7.8	20
372	Characterization of a cryptic rearrangement in a case of mantle cell lymphoma with negative FISH studies. <i>Blood Advances</i> , 2019 , 3, 1298-1302	7.8	11
371	Functional evaluation of isocitrate dehydrogenase 1 and 2 variants of unclear significance in chronic myeloid neoplasms. <i>Leukemia Research</i> , 2019 , 87, 106264	2.7	
370	Mate pair sequencing outperforms fluorescence in situ hybridization in the genomic characterization of multiple myeloma. <i>Blood Cancer Journal</i> , 2019 , 9, 103	7	18
369	Rapid assessment of hyperdiploidy in plasma cell disorders using a novel multi-parametric flow cytometry method. <i>American Journal of Hematology</i> , 2019 , 94, 424-430	7.1	5
368	Extramedullary acute myeloid leukemia presenting in young adults demonstrates sensitivity to high-dose anthracycline: a subset analysis from ECOG-ACRIN 1900. <i>Haematologica</i> , 2019 , 104, e147-e150	6.6	4
367	Calculator-free point-of-care prognostication in myelodysplastic syndromes. <i>American Journal of Hematology</i> , 2019 , 94, E99-E101	7.1	
366	Clinical correlates, prognostic impact and survival outcomes in chronic myelomonocytic leukemia patients with the V617F mutation. <i>Haematologica</i> , 2019 , 104, e236-e239	6.6	13
365	A randomized trial of three novel regimens for recurrent acute myeloid leukemia demonstrates the continuing challenge of treating this difficult disease. <i>American Journal of Hematology</i> , 2019 , 94, 111-117	7.1	19
364	Determinants of long-term outcome in type 1 calreticulin-mutated myelofibrosis. <i>Leukemia</i> , 2019 , 33, 780-785	10.7	2
363	The germline JAK2 GGCC (46/1) haplotype and survival among 414 molecularly-annotated patients with primary myelofibrosis. <i>American Journal of Hematology</i> , 2019 , 94, 299-305	7.1	5
362	20+ Years and alive with primary myelofibrosis: Phenotypic signature of very long-lived patients. <i>American Journal of Hematology</i> , 2019 , 94, 286-290	7.1	3
361	Constitutional chromosome rearrangements that mimic the 2017 world health organization "acute myeloid leukemia with recurrent genetic abnormalities": A study of three cases and review of the literature. <i>Cancer Genetics</i> , 2019 , 230, 37-46	2.3	4
360	Decreased survival and increased rate of fibrotic progression in essential thrombocythemia chronicled after the FDA approval date of anagrelide. <i>American Journal of Hematology</i> , 2019 , 94, 5-9	7.1	3
359	Use of mate-pair sequencing to characterize a complex cryptic BCR/ABL1 rearrangement observed in a newly diagnosed case of chronic myeloid leukemia. <i>Human Pathology</i> , 2019 , 89, 109-114	3.7	6
358	Mate pair sequencing improves detection of genomic abnormalities in acute myeloid leukemia. <i>European Journal of Haematology</i> , 2019 , 102, 87-96	3.8	23
357	U2AF1 mutation types in primary myelofibrosis: phenotypic and prognostic distinctions. <i>Leukemia</i> , 2018 , 32, 2274-2278	10.7	47
356	The impact of sex on disease phenotype and prognostic thresholds of anemia in myelodysplastic syndromes. <i>American Journal of Hematology</i> , 2018 , 93, E164-E167	7.1	1

355	GIPSS: genetically inspired prognostic scoring system for primary myelofibrosis. <i>Leukemia</i> , 2018 , 32, 1631-1642	10.7	17
354	Evaluation of Revised International Staging System (R-ISS) for transplant-eligible multiple myeloma patients. <i>Annals of Hematology</i> , 2018 , 97, 1453-1462	3	18
353	Prognostic significance of interphase FISH in monoclonal gammopathy of undetermined significance. <i>Leukemia</i> , 2018 , 32, 1811-1815	10.7	18
352	Sex and degree of severity influence the prognostic impact of anemia in primary myelofibrosis: analysis based on 1109 consecutive patients. <i>Leukemia</i> , 2018 , 32, 1254-1258	10.7	26
351	Blast phase myeloproliferative neoplasm: Mayo-AGIMM study of 410 patients from two separate cohorts. <i>Leukemia</i> , 2018 , 32, 1200-1210	10.7	68
350	Prognostic interaction between bone marrow morphology and SF3B1 and ASXL1 mutations in myelodysplastic syndromes with ring sideroblasts. <i>Blood Cancer Journal</i> , 2018 , 8, 18	7	13
349	FGFR1 rearranged hematological neoplasms - molecularly defined and clinically heterogeneous. <i>Leukemia and Lymphoma</i> , 2018 , 59, 1520-1522	1.9	5
348	Allogeneic hematopoietic stem cell transplant overcomes the adverse survival effect of very high risk and unfavorable karyotype in myelofibrosis. <i>American Journal of Hematology</i> , 2018 , 93, 649-654	7.1	26
347	Revised cytogenetic risk stratification in primary myelofibrosis: analysis based on 1002 informative patients. <i>Leukemia</i> , 2018 , 32, 1189-1199	10.7	65
346	Mutations and prognosis in myelodysplastic syndromes: karyotype-adjusted analysis of targeted sequencing in 300 consecutive cases and development of a genetic risk model. <i>American Journal of Hematology</i> , 2018 , 93, 691-697	7.1	34
345	Normal karyotype in myelofibrosis: is prognostic integrity affected by the number of metaphases analyzed?. <i>Blood Cancer Journal</i> , 2018 , 8, 8	7	0
344	EZH2 mutations in chronic myelomonocytic leukemia cluster with ASXL1 mutations and their co-occurrence is prognostically detrimental. <i>Blood Cancer Journal</i> , 2018 , 8, 12	7	30
343	Monocytosis is a powerful and independent predictor of inferior survival in primary myelofibrosis. <i>British Journal of Haematology</i> , 2018 , 183, 835-838	4.5	23
342	Myeloid neoplasm with eosinophilia associated with isolated extramedullary FIP1L1/PDGFRΑ rearrangement. <i>Cancer Genetics</i> , 2018 , 220, 13-18	2.3	3
341	Cytogenetic findings in WHO-defined polycythaemia vera and their prognostic relevance. <i>British Journal of Haematology</i> , 2018 , 182, 437-440	4.5	14
340	A comparison of clinical and molecular characteristics of patients with systemic mastocytosis with chronic myelomonocytic leukemia to CMML alone. <i>Leukemia</i> , 2018 , 32, 1850-1856	10.7	19
339	Recurrent fusions in indolent T-cell lymphoproliferative disorder of the gastrointestinal tract. <i>Blood</i> , 2018 , 131, 2262-2266	2.2	45
338	Nonhepatosplenic extramedullary manifestations of chronic myelomonocytic leukemia: clinical, molecular and prognostic correlates. <i>Leukemia and Lymphoma</i> , 2018 , 59, 2998-3001	1.9	5

337	Infrequent occurrence of TET1, TET3, and ASXL2 mutations in myelodysplastic/myeloproliferative neoplasms. <i>Blood Cancer Journal</i> , 2018 , 8, 32	7	9
336	Momelotinib therapy for myelofibrosis: a 7-year follow-up. <i>Blood Cancer Journal</i> , 2018 , 8, 29	7	27
335	Prefibrotic versus overtly fibrotic primary myelofibrosis: clinical, cytogenetic, molecular and prognostic comparisons. <i>British Journal of Haematology</i> , 2018 , 182, 594-597	4.5	18
334	Therapy related-chronic myelomonocytic leukemia (CMML): Molecular, cytogenetic, and clinical distinctions from de novo CMML. <i>American Journal of Hematology</i> , 2018 , 93, 65-73	7.1	37
333	Revisiting the need for bone marrow examination in chronic myeloid leukemia. <i>American Journal of Hematology</i> , 2018 , 93, 5-7	7.1	1
332	Validation of the WHO-defined 20% circulating blasts threshold for diagnosis of leukemic transformation in primary myelofibrosis. <i>Blood Cancer Journal</i> , 2018 , 8, 57	7	17
331	A Test Utilization Approach to the Diagnostic Workup of Isolated Eosinophilia in Otherwise Morphologically Unremarkable Bone Marrow: A Single Institutional Experience. <i>American Journal of Clinical Pathology</i> , 2018 , 150, 421-431	1.9	6
330	The Utilization of Chromosomal Microarray Technologies for Hematologic Neoplasms: An ACLPS Critical Review. <i>American Journal of Clinical Pathology</i> , 2018 , 150, 375-384	1.9	9
329	Early thrombotic events and preemptive systemic anticoagulation following splenectomy for myelofibrosis. <i>American Journal of Hematology</i> , 2018 , 93, E235-E238	7.1	7
328	Prognostic impact of ASXL1 mutations in patients with myelodysplastic syndromes and multilineage dysplasia with or without ring sideroblasts. <i>Leukemia Research</i> , 2018 , 71, 60-62	2.7	11
327	Blast phase chronic myelomonocytic leukemia: Mayo-MDACC collaborative study of 171 cases. <i>Leukemia</i> , 2018 , 32, 2512-2518	10.7	19
326	Development of a prognostically relevant cachexia index in primary myelofibrosis using serum albumin and cholesterol levels. <i>Blood Advances</i> , 2018 , 2, 1980-1984	7.8	8
325	Cytogenetic abnormalities in systemic mastocytosis: WHO subcategory-specific incidence and prognostic impact among 348 informative cases. <i>American Journal of Hematology</i> , 2018 , 93, 1461-1466	7.1	17
324	Splenectomy in patients with chronic myelomonocytic leukemia: Indications, histopathological findings and clinical outcomes in a single institutional series of thirty-nine patients. <i>American Journal of Hematology</i> , 2018 , 93, 1347-1357	7.1	7
323	Mayo Alliance Prognostic Model for Myelodysplastic Syndromes: Integration of Genetic and Clinical Information. <i>Mayo Clinic Proceedings</i> , 2018 , 93, 1363-1374	6.4	14
322	Inferior survival in high-grade B-cell lymphoma with and/or rearrangements is not associated with gene rearrangements. <i>Haematologica</i> , 2018 , 103, 1899-1907	6.6	31
321	Very poor long-term survival in past and more recent studies for relapsed AML patients: The ECOG-ACRIN experience. <i>American Journal of Hematology</i> , 2018 , 93, 1074-1081	7.1	45
320	Genetic subtyping of breast implant-associated anaplastic large cell lymphoma. <i>Blood</i> , 2018 , 132, 544-547.	7.2	60

319	Mutation-Enhanced International Prognostic Systems for Essential Thrombocythemia (MIPSS-ET) and Polycythemia Vera (MIPSS-PV). <i>Blood</i> , 2018 , 132, 578-578	2.2	5
318	20+ Years and Alive with Primary Myelofibrosis: Phenotypic Signature of Very Long-Lived Patients. <i>Blood</i> , 2018 , 132, 4301-4301	2.2	0
317	3,023 Mayo Clinic Patients with Myeloproliferative Neoplasms: Risk-Stratified Comparison of Survival and Outcomes Data Among Disease Subgroups. <i>Blood</i> , 2018 , 132, 3035-3035	2.2	1
316	Mutations and Thrombosis in Essential Thrombocythemia and Polycythemia Vera: Mayo-Careggi Alliance Study. <i>Blood</i> , 2018 , 132, 3040-3040	2.2	1
315	High Level MYC Amplification in Aggressive B-Cell Lymphomas: Is It a Marker of Aggressive Disease?. <i>Blood</i> , 2018 , 132, 1693-1693	2.2	1
314	Impact of Acquired Del(17p) in Patients with Multiple Myeloma. <i>Blood</i> , 2018 , 132, 4449-4449	2.2	
313	Predictors of Spleen and Anemia Response to Specific Drugs in Primary Myelofibrosis. <i>Blood</i> , 2018 , 132, 4300-4300	2.2	
312	Association of Clinical Epidemiologic Exposures and Overall Survival with Genome-Wide DNA Methylation Profiles in Acute Myeloid Leukemia: Analysis of the Mayo Clinic AML Epidemiology Cohort. <i>Blood</i> , 2018 , 132, 3987-3987	2.2	
311	Serum Erythropoietin Levels in Essential Thrombocythemia: Phenotypic and Prognostic Correlates. <i>Blood</i> , 2018 , 132, 3034-3034	2.2	
310	The Germline JAK2 GGCC (46/1) Haplotype and Survival Among 414 Molecularly-Annotated Patients with Primary Myelofibrosis. <i>Blood</i> , 2018 , 132, 1761-1761	2.2	
309	Decreased Survival and Increased Rate of Fibrotic Progression in Essential Thrombocythemia Chronicled after the FDA Approval Date of Anagrelide. <i>Blood</i> , 2018 , 132, 4287-4287	2.2	
308	Clinical and Molecular Models of Prognostication in Mastocytosis: Analysis Based on 580 Consecutive Cases. <i>Blood</i> , 2018 , 132, 582-582	2.2	
307	Determinants of Long-Term Outcome in Type 1/like Calreticulin-Mutated Myelofibrosis. <i>Blood</i> , 2018 , 132, 1767-1767	2.2	
306	Cytogenetic Abnormalities in Systemic Mastocytosis: Who Subcategory-Specific Incidence and Prognostic Impact Among 348 Informative Cases. <i>Blood</i> , 2018 , 132, 3050-3050	2.2	
305	Myeloproliferative Neoplasms in Young Patients: The Mayo Clinic Experience with 361 Cases Age 40 Years or Younger. <i>Blood</i> , 2018 , 132, 3033-3033	2.2	
304	Cytogenetic Clonal Evolution in Myeloproliferative Neoplasms: Contexts and Prognostic Impact Among 650 Patients with Serial Bone Marrow Biopsies. <i>Blood</i> , 2018 , 132, 4291-4291	2.2	
303	Frequency of Acquired Genetic Mutations and Their Prognostic Impact on Patients with Incidental Finding of Isolated 20q- in Bone Marrow without Morphologic Evidence of a Myeloid Neoplasm. <i>Blood</i> , 2018 , 132, 4382-4382	2.2	
302	MPL-Mutated Essential Thrombocythemia: A Morphologic Reappraisal. <i>Blood</i> , 2018 , 132, 3036-3036	2.2	

301	Clinical Correlates, Prognostic Impact and Survival Outcomes in Chronic Myelomonocytic Leukemia Patients with Myeloproliferative Neoplasm Associated-Driver Mutations. <i>Blood</i> , 2018 , 132, 3100-3100	2.2	
300	Efficacy of Mitoxantrone-Based Salvage Therapies in Relapsed or Refractory Acute Myeloid Leukemia in the Mayo Clinic Cancer Center: Analysis of Survival after CLAG-M Vs. MEC. <i>Blood</i> , 2018 , 132, 2678-2678	2.2	
299	1,123 Consecutive Adults with Non-APL Acute Myeloid Leukemia: The Mayo Clinic Experience. <i>Blood</i> , 2018 , 132, 2689-2689	2.2	
298	Risk Factors for Leukemic Transformation Among 1,306 Patients with Primary Myelofibrosis: Mutations Predict Early Events. <i>Blood</i> , 2018 , 132, 3044-3044	2.2	
297	A Prospective Evaluation of Vitamin B1 (thiamine) Level in Myeloproliferative Neoplasms: Clinical Correlations and Impact of JAK2 Inhibitor Therapy. <i>Blood</i> , 2018 , 132, 1771-1771	2.2	
296	Driver mutations and prognosis in primary myelofibrosis: Mayo-Careggi MPN alliance study of 1,095 patients. <i>American Journal of Hematology</i> , 2018 , 93, 348-355	7.1	54
295	Screening for ASXL1 and SRSF2 mutations is imperative for treatment decision-making in otherwise low or intermediate-1 risk patients with myelofibrosis. <i>British Journal of Haematology</i> , 2018 , 183, 678-681	4.5	10
294	Immunophenotypic and laboratory features of t(11;14)(q13;q32)-positive plasma cell neoplasms. <i>Leukemia and Lymphoma</i> , 2018 , 59, 1913-1919	1.9	2
293	MIPSS70: Mutation-Enhanced International Prognostic Score System for Transplantation-Age Patients With Primary Myelofibrosis. <i>Journal of Clinical Oncology</i> , 2018 , 36, 310-318	2.2	224
292	MPL-mutated essential thrombocythemia: a morphologic reappraisal. <i>Blood Cancer Journal</i> , 2018 , 8, 1217		13
291	Serum erythropoietin levels in essential thrombocythemia: phenotypic and prognostic correlates. <i>Blood Cancer Journal</i> , 2018 , 8, 118	7	3
290	Genetic predictors of response to specific drugs in primary myelofibrosis. <i>Blood Cancer Journal</i> , 2018 , 8, 120	7	0
289	Molecular profiling reveals immunogenic cues in anaplastic large cell lymphomas with rearrangements. <i>Blood</i> , 2018 , 132, 1386-1398	2.2	44
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285	Mutations and karyotype predict treatment response in myelodysplastic syndromes. <i>American Journal of Hematology</i> , 2018 , 93, 1420-1426	7.1	18
284	Practice-relevant demarcation of systemic mastocytosis associated with another hematologic neoplasm. <i>American Journal of Hematology</i> , 2018 , 93, E383-E386	7.1	2

283	Myeloproliferative neoplasms in the young: Mayo Clinic experience with 361 patients age 40 years or younger. <i>American Journal of Hematology</i> , 2018 , 93, 1474-1484	7.1	31
282	Targeted next-generation sequencing in blast phase myeloproliferative neoplasms. <i>Blood Advances</i> , 2018 , 2, 370-380	7.8	55
281	Mayo alliance prognostic system for mastocytosis: clinical and hybrid clinical-molecular models. <i>Blood Advances</i> , 2018 , 2, 2964-2972	7.8	40
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277	Mantle cell lymphoma with a novel t(11;12)(q13;p11.2): a proposed alternative mechanism of CCND1 up-regulation. <i>Human Pathology</i> , 2017 , 64, 207-212	3.7	5
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272	Targeted next generation sequencing and identification of risk factors in World Health Organization defined atypical chronic myeloid leukemia. <i>American Journal of Hematology</i> , 2017 , 92, 542-548	7.1	41
271	Diagnosis and Management of Waldenström Macroglobulinemia: Mayo Stratification of Macroglobulinemia and Risk-Adapted Therapy (mSMART) Guidelines 2016. <i>JAMA Oncology</i> , 2017 , 3, 1257-1265	13.4	82
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265	Spectrum of autoimmune diseases and systemic inflammatory syndromes in patients with chronic myelomonocytic leukemia. <i>Leukemia and Lymphoma</i> , 2017 , 58, 1488-1493	1.9	35
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250	Abnormal Karyotype and Prognosis in Polycythemia Vera: A Single Center Experience in 239 Informative Cases. <i>Blood</i> , 2016 , 128, 3115-3115	2.2	1
249	Prefibrotic Versus Overtly Fibrotic Primary Myelofibrosis: Clinical, Cytogenetic, Molecular and Prognostic Comparisons. <i>Blood</i> , 2016 , 128, 4247-4247	2.2	2
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245	Risk Factors for Arterial Versus Venous Thrombosis in Polycythemia Vera: Single Center Experience in 587 Patients. <i>Blood</i> , 2016 , 128, 948-948	2.2	1
244	"Proliferative" Versus "Dysplastic" Chronic Myelomonocytic Leukemia: Molecular and Prognostic Correlates. <i>Blood</i> , 2016 , 128, 1987-1987	2.2	0
243	Unique Clinical Epidemiologic Risk Factors Are Associated with Distinct Methylation Subgroups in Newly-Diagnosed Acute Myeloid Leukemia (AML). <i>Blood</i> , 2016 , 128, 1719-1719	2.2	0
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240	Prognostic Implications of Multiple Cytogenetic High-Risk Abnormalities in Patients with Newly Diagnosed Multiple Myeloma. <i>Blood</i> , 2016 , 128, 5615-5615	2.2	
239	Monocytosis in Polycythemia Vera: Clinical and Molecular Correlates. <i>Blood</i> , 2016 , 128, 4259-4259	2.2	
238	Utilization of Mate-Pair Sequencing to Characterize Complex and Novel TCF3 Translocations. <i>Blood</i> , 2016 , 128, 4086-4086	2.2	
237	Gene Expression Profiling Identifies Distinct Signatures for Dysplastic and Proliferative Chronic Myelomonocytic Leukemia. <i>Blood</i> , 2016 , 128, 110-110	2.2	
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234	Subnormal Lymphocyte Count Predicts Inferior Survival in Myelodysplastic Syndromes: A Single Center Experience in 889 Patients. <i>Blood</i> , 2016 , 128, 5534-5534	2.2	0
233	Predicting Poor Overall Survival in Patients with Newly Diagnosed Multiple Myeloma and Standard-Risk Cytogenetics Treated with Novel Agents. <i>Blood</i> , 2016 , 128, 3255-3255	2.2	
232	DNTM3A Mutations and Prognosis in Chronic Myelomonocytic Leukemia. <i>Blood</i> , 2016 , 128, 1988-1988	2.2	
231	Evaluation of Revised International Staging System for Transplant-Eligible Multiple Myeloma Patients. <i>Blood</i> , 2016 , 128, 3452-3452	2.2	
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218	Calreticulin variant stratified driver mutational status and prognosis in essential thrombocythemia. <i>American Journal of Hematology</i> , 2016 , 91, 503-6	7.1	37
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216	Chronic myelomonocytic leukemia in younger patients: molecular and cytogenetic predictors of survival and treatment outcome. <i>Blood Cancer Journal</i> , 2015 , 5, e270	7	21
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199	Targeted Next-Generation Sequencing in Polycythemia Vera and Essential Thrombocythemia. <i>Blood</i> , 2015 , 126, 354-354	2.2	9
198	ASXL1 and CBL Mutations Are Independently Predictive of Inferior Survival in Advanced Systemic Mastocytosis. <i>Blood</i> , 2015 , 126, 828-828	2.2	2
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28	FIP1L1-PDGFRα fusion: prevalence and clinicopathologic correlates in 89 consecutive patients with moderate to severe eosinophilia. <i>Blood</i> , 2004 , 104, 3038-45	2.2	248
27	Familial 22q11.2 deletions in DiGeorge/velocardiofacial syndrome are predominantly smaller than the commonly observed 3Mb. <i>Genetics in Medicine</i> , 2004 , 6, 517-20	8.1	17
26	Bone Marrow Histologic and Cytogenetic Changes during Leukemic Transformation of Myelofibrosis with Myeloid Metaplasia.. <i>Blood</i> , 2004 , 104, 4750-4750	2.2	
25	Discovery of 6 Novel Translocations Involving the Imatinib Responsive Genes PDGFRB and PDGFRB from Screening 29,047 Abnormal Bone Marrow Specimens.. <i>Blood</i> , 2004 , 104, 2902-2902	2.2	2
24	CHIC2 deletion, a surrogate for FIP1L1-PDGFRα fusion, occurs in systemic mastocytosis associated with eosinophilia and predicts response to imatinib mesylate therapy. <i>Blood</i> , 2003 , 102, 3093-6	2.2	327
23	New highly sensitive fluorescence in situ hybridization method to detect PML/RARA fusion in acute promyelocytic leukemia. <i>Cancer Genetics and Cytogenetics</i> , 2003 , 145, 144-51		52
22	Concomitant myelodysplastic syndrome and chronic myeloid leukaemia: treatment outcomes with imatinib mesylate. <i>British Journal of Haematology</i> , 2003 , 123, 366-7	4.5	2
21	Microduplication 22q11.2, an emerging syndrome: clinical, cytogenetic, and molecular analysis of thirteen patients. <i>American Journal of Human Genetics</i> , 2003 , 73, 1027-40	11	298
20	Utility of subtelomeric fluorescent DNA probes for detection of chromosome anomalies in 425 patients. <i>Genetics in Medicine</i> , 2003 , 5, 28-34	8.1	49
19	Histologic and Immunohistochemical Study of Bone Marrow Monocytic Nodules in 21 Cases With Myelodysplasia. <i>American Journal of Clinical Pathology</i> , 2003 , 120, 874-881	1.9	1
18	Primary myelodysplastic syndrome with normal cytogenetics: utility of 'FISH panel testing' and M-FISH. <i>Leukemia Research</i> , 2002 , 26, 235-40	2.7	57
17	Inherited interstitial deletion of chromosomes 5p and 16q without apparent phenotypic effect: further confirmation. <i>Prenatal Diagnosis</i> , 2000 , 20, 144-8; discussion 149-51	3.2	27
16	Reported in vivo splice-site mutations in the factor IX gene: severity of splicing defects and a hypothesis for predicting deleterious splice donor mutations. <i>Human Mutation</i> , 1999 , 13, 221-31	4.7	45
15	Germline mutations in Peruvian patients with hemophilia B: pattern of mutation in AmerIndians is similar to the putative endogenous germline pattern. <i>Human Mutation</i> , 1998 , 11, 372-6	4.7	10
14	The factor IX gene as a model for analysis of human germline mutations: an update. <i>Human Molecular Genetics</i> , 1996 , 5 Spec No, 1505-14	5.6	43

13	Absence of somatic mosaicism in 17 families with hemophilia B: an analysis with a sensitivity 10- to 1000-fold greater than that of sequencing gels. <i>Human Genetics</i> , 1996 , 98, 539-45	6.3	10
12	Two novel factor IX promoter mutations: incremental progress towards 'saturation in vivo mutagenesis' of a human promoter region. <i>Human Molecular Genetics</i> , 1995 , 4, 769-70	5.6	11
11	Molecular Epidemiology of Factor IX Germline Mutations in Mexican Hispanics: Pattern of Mutation and Potential Founder Effects. <i>Thrombosis and Haemostasis</i> , 1995 , 74, 1416-1422	7	15
10	Factor VIII gene inversions causing severe hemophilia A originate almost exclusively in male germ cells. <i>Human Molecular Genetics</i> , 1994 , 3, 1035-9	5.6	144
9	"Cryptic" dinucleotide polymorphism in the 3' region of the factor IX gene shows substantial variation among different populations. <i>Human Genetics</i> , 1994 , 93, 357-8	6.3	10
8	How precisely can data from transgenic mouse mutation-detection systems be extrapolated to humans?: lesions from the human factor IX gene. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 1994 , 307, 517-31	3.3	15
7	A mutation in the 3' untranslated region of the factor IX gene in four families with hemophilia B. <i>Human Molecular Genetics</i> , 1993 , 2, 1309-10	5.6	18
6	Germline mutations in the factor IX gene: a comparison of the pattern in Caucasians and non-Caucasians. <i>Human Molecular Genetics</i> , 1993 , 2, 293-8	5.6	28
5	Deletions with inversions: report of a mutation and review of the literature. <i>Human Mutation</i> , 1993 , 2, 53-7	4.7	21
4	The pattern of spontaneous germ-line mutation: relative rates of mutation at or near CpG dinucleotides in the factor IX gene. <i>Human Genetics</i> , 1993 , 91, 496-503	6.3	37
3	PRE: a novel element with the hallmarks of a retrotransposon derived from an unknown structural RNA. <i>Nucleic Acids Research</i> , 1992 , 20, 5233	20.1	2
2	Evidence that descendants of three founders constitute about 25% of hemophilia B in the United States. <i>Genomics</i> , 1991 , 10, 1093-6	4.3	54
1	A past mutation at isoleucine 397 is now a common cause of moderate/mild haemophilia B. <i>British Journal of Haematology</i> , 1990 , 75, 212-6	4.5	33