

Rhett P Ketterling

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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	Prognostic relevance of integrated genetic profiling in acute myeloid leukemia. <i>New England Journal of Medicine</i> , 2012 , 366, 1079-89	59.2	1378
461	Anthracycline dose intensification in acute myeloid leukemia. <i>New England Journal of Medicine</i> , 2009 , 361, 1249-59	59.2	658
460	Long-term survival and blast transformation in molecularly annotated essential thrombocythemia, polycythemia vera, and myelofibrosis. <i>Blood</i> , 2014 , 124, 2507-13; quiz 2615	2.2	424
459	CHIC2 deletion, a surrogate for FIP1L1-PDGFR 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
458	Leukemic transformation in myelofibrosis with myeloid metaplasia: a single-institution experience with 91 cases. <i>Blood</i> , 2005 , 105, 973-7	2.2	305
457	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
456	ALK-negative anaplastic large cell lymphoma is a genetically heterogeneous disease with widely disparate clinical outcomes. <i>Blood</i> , 2014 , 124, 1473-80	2.2	294
455	T-cell acute lymphoblastic leukemia in adults: clinical features, immunophenotype, cytogenetics, and outcome from the large randomized prospective trial (UKALL XII/ECOG 2993). <i>Blood</i> , 2009 , 114, 5136-45	2.2	277
454	FIP1L1-PDGFR fusion: prevalence and clinicopathologic correlates in 89 consecutive patients with moderate to severe eosinophilia. <i>Blood</i> , 2004 , 104, 3038-45	2.2	248
453	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
452	Lenalidomide therapy in myelofibrosis with myeloid metaplasia. <i>Blood</i> , 2006 , 108, 1158-64	2.2	211
451	Prolonged administration of azacitidine with or without entinostat for myelodysplastic syndrome and acute myeloid leukemia with myelodysplasia-related changes: results of the US Leukemia Intergroup trial E1905. <i>Journal of Clinical Oncology</i> , 2014 , 32, 1242-8	2.2	187
450	ASXL1 and SETBP1 mutations and their prognostic contribution in chronic myelomonocytic leukemia: a two-center study of 466 patients. <i>Leukemia</i> , 2014 , 28, 2206-12	10.7	186
449	Trisomies in multiple myeloma: impact on survival in patients with high-risk cytogenetics. <i>Blood</i> , 2012 , 119, 2100-5	2.2	181
448	HER2 and chromosome 17 effect on patient outcome in the N9831 adjuvant trastuzumab trial. <i>Journal of Clinical Oncology</i> , 2010 , 28, 4307-15	2.2	175
447	SF3B1 mutations are prevalent in myelodysplastic syndromes with ring sideroblasts but do not hold independent prognostic value. <i>Blood</i> , 2012 , 119, 569-72	2.2	164
446	Genome-wide analysis reveals recurrent structural abnormalities of TP63 and other p53-related genes in peripheral T-cell lymphomas. <i>Blood</i> , 2012 , 120, 2280-9	2.2	164

445	Targeted deep sequencing in polycythemia vera and essential thrombocythemia. <i>Blood Advances</i> , 2016 , 1, 21-30	7.8	163
444	Zosuquidar, a novel modulator of P-glycoprotein, does not improve the outcome of older patients with newly diagnosed acute myeloid leukemia: a randomized, placebo-controlled trial of the Eastern Cooperative Oncology Group 3999. <i>Blood</i> , 2010 , 116, 4077-85	2.2	157
443	Mayo prognostic model for WHO-defined chronic myelomonocytic leukemia: ASXL1 and spliceosome component mutations and outcomes. <i>Leukemia</i> , 2013 , 27, 1504-10	10.7	151
442	Type 1 versus Type 2 calreticulin mutations in essential thrombocythemia: a collaborative study of 1027 patients. <i>American Journal of Hematology</i> , 2014 , 89, E121-4	7.1	145
441	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
440	One thousand patients with primary myelofibrosis: the mayo clinic experience. <i>Mayo Clinic Proceedings</i> , 2012 , 87, 25-33	6.4	137
439	Clonally related histiocytic/dendritic cell sarcoma and chronic lymphocytic leukemia/small lymphocytic lymphoma: a study of seven cases. <i>Modern Pathology</i> , 2011 , 24, 1421-32	9.8	135
438	Clinical correlates of JAK2V617F allele burden in essential thrombocythemia. <i>Cancer</i> , 2007 , 109, 2279-84	6.4	132
437	SRSF2 mutations in primary myelofibrosis: significant clustering with IDH mutations and independent association with inferior overall and leukemia-free survival. <i>Blood</i> , 2012 , 120, 4168-71	2.2	128
436	Chromosomal rearrangements of 6p25.3 define a new subtype of lymphomatoid papulosis. <i>American Journal of Surgical Pathology</i> , 2013 , 37, 1173-81	6.7	125
435	Overexpression of IL-1 receptor accessory protein in stem and progenitor cells and outcome correlation in AML and MDS. <i>Blood</i> , 2012 , 120, 1290-8	2.2	123
434	Targeted deep sequencing in primary myelofibrosis. <i>Blood Advances</i> , 2016 , 1, 105-111	7.8	120
433	GIPSS: genetically inspired prognostic scoring system for primary myelofibrosis. <i>Leukemia</i> , 2018 , 32, 1631-42	10.7	117
432	Spliceosome mutations involving SRSF2, SF3B1, and U2AF35 in chronic myelomonocytic leukemia: prevalence, clinical correlates, and prognostic relevance. <i>American Journal of Hematology</i> , 2013 , 88, 201-6	7.1	112
431	Differential prognostic effect of IDH1 versus IDH2 mutations in myelodysplastic syndromes: a Mayo Clinic study of 277 patients. <i>Leukemia</i> , 2012 , 26, 101-5	10.7	112
430	FIP1L1-PDGFRα in eosinophilic disorders: prevalence in routine clinical practice, long-term experience with imatinib therapy, and a critical review of the literature. <i>Leukemia Research</i> , 2006 , 30, 965-70	2.7	110
429	The prognostic advantage of calreticulin mutations in myelofibrosis might be confined to type 1 or type 1-like CALR variants. <i>Blood</i> , 2014 , 124, 2465-6	2.2	105
428	Molecular and prognostic correlates of cytogenetic abnormalities in chronic myelomonocytic leukemia: a Mayo Clinic-French Consortium Study. <i>American Journal of Hematology</i> , 2014 , 89, 1111-5	7.1	104

427	Relationship of patient survival and chromosome anomalies detected in metaphase and/or interphase cells at diagnosis of myeloma. <i>Blood</i> , 2005 , 106, 3553-8	2.2	100
426	Prognostic interaction between ASXL1 and TET2 mutations in chronic myelomonocytic leukemia. <i>Blood Cancer Journal</i> , 2016 , 6, e385	7	83
425	Utility of ALK-1 protein expression and ALK rearrangements in distinguishing inflammatory myofibroblastic tumor from malignant spindle cell lesions of the urinary bladder. <i>Modern Pathology</i> , 2007 , 20, 592-603	9.8	83
424	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
423	Preclinical validation of fluorescence in situ hybridization assays for clinical practice. <i>Genetics in Medicine</i> , 2006 , 8, 16-23	8.1	81
422	Benefit of high-dose daunorubicin in AML induction extends across cytogenetic and molecular groups. <i>Blood</i> , 2016 , 127, 1551-8	2.2	81
421	Translocation t(11;14) and survival of patients with light chain (AL) amyloidosis. <i>Haematologica</i> , 2009 , 94, 380-6	6.6	80
420	CD25 expression status improves prognostic risk classification in AML independent of established biomarkers: ECOG phase 3 trial, E1900. <i>Blood</i> , 2012 , 120, 2297-306	2.2	78
419	SETBP1 mutations in 415 patients with primary myelofibrosis or chronic myelomonocytic leukemia: independent prognostic impact in CMML. <i>Leukemia</i> , 2013 , 27, 2100-2	10.7	75
418	Analysis of Intratumoral Heterogeneity and Amplification Status in Breast Carcinomas With Equivocal (2+) HER-2 Immunostaining. <i>American Journal of Clinical Pathology</i> , 2005 , 124, 273-281	1.9	73
417	WHO-defined 'myelodysplastic syndrome with isolated del(5q)' in 88 consecutive patients: survival data, leukemic transformation rates and prevalence of JAK2, MPL and IDH mutations. <i>Leukemia</i> , 2010 , 24, 1283-9	10.7	72
416	Integrative epigenomic analysis identifies biomarkers and therapeutic targets in adult B-acute lymphoblastic leukemia. <i>Cancer Discovery</i> , 2012 , 2, 1004-23	24.4	70
415	Morphologic Features of ALK-negative Anaplastic Large Cell Lymphomas With DUSP22 Rearrangements. <i>American Journal of Surgical Pathology</i> , 2016 , 40, 36-43	6.7	70
414	and rearrangements predict outcome of ALK-negative anaplastic large cell lymphoma: a Danish cohort study. <i>Blood</i> , 2017 , 130, 554-557	2.2	68
413	Blast phase myeloproliferative neoplasm: Mayo-AGIMM study of 410 patients from two separate cohorts. <i>Leukemia</i> , 2018 , 32, 1200-1210	10.7	68
412	Change in Pattern of HER2 Fluorescent in Situ Hybridization (FISH) Results in Breast Cancers Submitted for FISH Testing: Experience of a Reference Laboratory Using US Food and Drug Administration Criteria and American Society of Clinical Oncology and College of American Pathologists Guidelines. <i>Journal of Clinical Oncology</i> , 2016 , 34, 3502-3510	2.2	68
411	Revised cytogenetic risk stratification in primary myelofibrosis: analysis based on 1002 informative patients. <i>Leukemia</i> , 2018 , 32, 1189-1199	10.7	65
410	Genetic subtyping of breast implant-associated anaplastic large cell lymphoma. <i>Blood</i> , 2018 , 132, 544-547	7.2	60

409	Clonally related follicular lymphomas and Langerhans cell neoplasms: expanding the spectrum of transdifferentiation. <i>American Journal of Surgical Pathology</i> , 2013 , 37, 978-86	6.7	60
408	Molecular diagnosis of Ewing's sarcoma/primitive neuroectodermal tumor in formalin-fixed paraffin-embedded tissues by RT-PCR and fluorescence in situ hybridization. <i>Diagnostic Molecular Pathology</i> , 2005 , 14, 23-8		60
407	Prognostic irrelevance of ring sideroblast percentage in World Health Organization-defined myelodysplastic syndromes without excess blasts. <i>Blood</i> , 2012 , 119, 5674-7	2.2	59
406	Mutation-enhanced international prognostic systems for essential thrombocythaemia and polycythaemia vera. <i>British Journal of Haematology</i> , 2020 , 189, 291-302	4.5	58
405	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
404	Rosette-forming glioneuronal tumor: report of a chiasmal-optic nerve example in neurofibromatosis type 1: special pathology report. <i>Neurosurgery</i> , 2009 , 64, E771-2; discussion E772	3.2	56
403	Cytogenetic studies at diagnosis in polycythemia vera: clinical and JAK2V617F allele burden correlates. <i>European Journal of Haematology</i> , 2008 , 80, 197-200	3.8	56
402	Extending Jak2V617F and MplW515 mutation analysis to single hematopoietic colonies and B and T lymphocytes. <i>Stem Cells</i> , 2007 , 25, 2358-62	5.8	55
401	Predictors of survival in refractory anemia with ring sideroblasts and thrombocytosis (RARS-T) and the role of next-generation sequencing. <i>American Journal of Hematology</i> , 2016 , 91, 492-8	7.1	55
400	Targeted next-generation sequencing in blast phase myeloproliferative neoplasms. <i>Blood Advances</i> , 2018 , 2, 370-380	7.8	55
399	Biologic and genetic characterization of the novel amyloidogenic lambda light chain-secreting human cell lines, ALMC-1 and ALMC-2. <i>Blood</i> , 2008 , 112, 1931-41	2.2	54
398	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
397	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
396	Autologous transplantation gives encouraging results for young adults with favorable-risk acute myeloid leukemia, but is not improved with gemtuzumab ozogamicin. <i>Blood</i> , 2011 , 117, 5306-13	2.2	53
395	Cytogenetic abnormalities in essential thrombocythemia: prevalence and prognostic significance. <i>European Journal of Haematology</i> , 2009 , 83, 17-21	3.8	53
394	Evidence for cytogenetic and fluorescence in situ hybridization risk stratification of newly diagnosed multiple myeloma in the era of novel therapies. <i>Mayo Clinic Proceedings</i> , 2010 , 85, 532-7	6.4	52
393	Section E9 of the American College of Medical Genetics technical standards and guidelines: fluorescence in situ hybridization. <i>Genetics in Medicine</i> , 2011 , 13, 667-75	8.1	52
392	Reflex fluorescent in situ hybridization testing for unsuccessful product of conception cultures: a retrospective analysis of 5555 samples attempted by conventional cytogenetics and fluorescent in situ hybridization. <i>Genetics in Medicine</i> , 2011 , 13, 545-52	8.1	52

391	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
390	Pituitary blastoma. <i>Acta Neuropathologica</i> , 2008 , 116, 657-66	14.3	51
389	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
388	Targeted next-generation sequencing in myelodysplastic syndromes and prognostic interaction between mutations and IPSS-R. <i>American Journal of Hematology</i> , 2017 , 92, 1311-1317	7.1	50
387	Azacitidine with or without Entinostat for the treatment of therapy-related myeloid neoplasm: further results of the E1905 North American Leukemia Intergroup study. <i>British Journal of Haematology</i> , 2016 , 172, 384-91	4.5	49
386	Histiocytoid Sweet syndrome may indicate leukemia cutis: a novel application of fluorescence in situ hybridization. <i>Journal of the American Academy of Dermatology</i> , 2014 , 70, 1021-7	4.5	49
385	C-MYC alterations and association with patient outcome in early-stage HER2-positive breast cancer from the north central cancer treatment group N9831 adjuvant trastuzumab trial. <i>Journal of Clinical Oncology</i> , 2011 , 29, 651-9	2.2	49
384	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
383	DNMT3A mutations are associated with inferior overall and leukemia-free survival in chronic myelomonocytic leukemia. <i>American Journal of Hematology</i> , 2017 , 92, 56-61	7.1	48
382	U2AF1 mutation types in primary myelofibrosis: phenotypic and prognostic distinctions. <i>Leukemia</i> , 2018 , 32, 2274-2278	10.7	47
381	The oncogenic transcription factor IRF4 is regulated by a novel CD30/NF- κ B positive feedback loop in peripheral T-cell lymphoma. <i>Blood</i> , 2015 , 125, 3118-27	2.2	47
380	Karyotype complements the International Prognostic Scoring System for primary myelofibrosis. <i>European Journal of Haematology</i> , 2009 , 82, 255-9	3.8	47
379	Next-generation sequencing in systemic mastocytosis: Derivation of a mutation-augmented clinical prognostic model for survival. <i>American Journal of Hematology</i> , 2016 , 91, 888-93	7.1	47
378	Treatment-influenced associations of PML-RAR α mutations, FLT3 mutations, and additional chromosome abnormalities in relapsed acute promyelocytic leukemia. <i>Blood</i> , 2012 , 120, 2098-108	2.2	46
377	Recurrent fusions in indolent T-cell lymphoproliferative disorder of the gastrointestinal tract. <i>Blood</i> , 2018 , 131, 2262-2266	2.2	45
376	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
375	Blast transformation in chronic myelomonocytic leukemia: Risk factors, genetic features, survival, and treatment outcome. <i>American Journal of Hematology</i> , 2015 , 90, 411-6	7.1	45
374	Pediatric histiocytic sarcoma clonally related to precursor B-cell acute lymphoblastic leukemia with homozygous deletion of CDKN2A encoding p16INK4A. <i>Pediatric Blood and Cancer</i> , 2011 , 56, 307-10	3	45

373	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
372	Molecular profiling reveals immunogenic cues in anaplastic large cell lymphomas with rearrangements. <i>Blood</i> , 2018 , 132, 1386-1398	2.2	44
371	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
370	Recurrent IDH mutations in high-risk myelodysplastic syndrome or acute myeloid leukemia with isolated del(5q). <i>Leukemia</i> , 2010 , 24, 1370-2	10.7	42
369	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
368	Mayo alliance prognostic system for mastocytosis: clinical and hybrid clinical-molecular models. <i>Blood Advances</i> , 2018 , 2, 2964-2972	7.8	40
367	ALK-1 protein expression and ALK gene rearrangements aid in the diagnosis of inflammatory myofibroblastic tumors of the female genital tract. <i>Archives of Pathology and Laboratory Medicine</i> , 2012 , 136, 623-6	5	39
366	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
365	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
364	Calreticulin variant stratified driver mutational status and prognosis in essential thrombocythemia. <i>American Journal of Hematology</i> , 2016 , 91, 503-6	7.1	37
363	Development of acute megakaryoblastic leukemia in Down syndrome is associated with sequential epigenetic changes. <i>Blood</i> , 2013 , 122, e33-43	2.2	36
362	Establishment and characterization of a novel Waldenstrom macroglobulinemia cell line, MWCL-1. <i>Blood</i> , 2011 , 117, e190-7	2.2	36
361	Impact of American Society of Clinical Oncology/College of American Pathologists guideline recommendations on HER2 interpretation in breast cancer. <i>Human Pathology</i> , 2010 , 41, 103-6	3.7	36
360	Mutations and thrombosis in essential thrombocythemia: prognostic interaction with age and thrombosis history. <i>European Journal of Haematology</i> , 2015 , 94, 31-6	3.8	35
359	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
358	Novel recurrent mutations in ethanolamine kinase 1 (ETNK1) gene in systemic mastocytosis with eosinophilia and chronic myelomonocytic leukemia. <i>Blood Cancer Journal</i> , 2015 , 5, e275	7	35
357	Analysis of genetic abnormalities provides insights into genetic evolution of hyperdiploid myeloma. <i>Genes Chromosomes and Cancer</i> , 2006 , 45, 1111-20	5	35
356	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

355	Intralymphatic cutaneous anaplastic large cell lymphoma/lymphomatoid papulosis: expanding the spectrum of CD30-positive lymphoproliferative disorders. <i>American Journal of Surgical Pathology</i> , 2014 , 38, 1203-11	6.7	34
354	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
353	Expression of p63 protein in anaplastic large cell lymphoma: implications for genetic subtyping. <i>Human Pathology</i> , 2017 , 64, 19-27	3.7	31
352	Monocytosis in polycythemia vera: Clinical and molecular correlates. <i>American Journal of Hematology</i> , 2017 , 92, 640-645	7.1	31
351	Inferior survival in high-grade B-cell lymphoma with and and/or rearrangements is not associated with gene rearrangements. <i>Haematologica</i> , 2018 , 103, 1899-1907	6.6	31
350	Chromosome 8p11.2 translocations: prevalence, FISH analysis for FGFR1 and MYST3, and clinicopathologic correlates in a consecutive cohort of 13 cases from a single institution. <i>American Journal of Hematology</i> , 2010 , 85, 238-42	7.1	31
349	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
348	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
347	Chronic lymphocytic leukemia with t(14;19)(q32;q13) is characterized by atypical morphologic and immunophenotypic features and distinctive genetic features. <i>American Journal of Clinical Pathology</i> , 2011 , 135, 686-96	1.9	30
346	Evaluation of revised IPSS cytogenetic risk stratification and prognostic impact of monosomal karyotype in 783 patients with primary myelodysplastic syndromes. <i>American Journal of Hematology</i> , 2013 , 88, 690-3	7.1	29
345	t(8;9)(p22;p24)/PCM1-JAK2 activates SOCS2 and SOCS3 via STAT5. <i>PLoS ONE</i> , 2013 , 8, e53767	3.7	29
344	Rearrangements and amplification of IER3 (IEX-1) represent a novel and recurrent molecular abnormality in myelodysplastic syndromes. <i>Cancer Research</i> , 2009 , 69, 7518-23	10.1	29
343	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
342	Desmoplastic small round cell tumor of the central nervous system: report of two cases and review of the literature. <i>Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin</i> , 2009 , 454, 431-9	5.1	28
341	CCND1 rearrangements and cyclin D1 overexpression in renal oncocytomas: frequency, clinicopathologic features, and utility in differentiation from chromophobe renal cell carcinoma. <i>Human Pathology</i> , 2009 , 40, 1296-303	3.7	28
340	Functioning paraganglioma and gastrointestinal stromal tumor of the jejunum in three women: syndrome or coincidence. <i>American Journal of Surgical Pathology</i> , 2006 , 30, 42-9	6.7	28
339	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
338	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

337	Momelotinib therapy for myelofibrosis: a 7-year follow-up. <i>Blood Cancer Journal</i> , 2018 , 8, 29	7	27
336	Chronic lymphocytic leukemia With t(2;14)(p16;q32) involves the BCL11A and IgH genes and is associated with atypical morphologic features and unmutated IgVH genes. <i>American Journal of Clinical Pathology</i> , 2009 , 131, 663-70	1.9	27
335	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
334	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
333	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
332	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
331	A compendium of cytogenetic abnormalities in myelofibrosis: molecular and phenotypic correlates in 826 patients. <i>British Journal of Haematology</i> , 2015 , 169, 71-6	4.5	26
330	Clonal relationship between precursor B-cell acute lymphoblastic leukemia and histiocytic sarcoma: a case report and discussion in the context of similar cases. <i>Leukemia Research</i> , 2010 , 34, e71-3	2.7	25
329	Comparison of fluorescence in situ hybridization, p57 immunostaining, flow cytometry, and digital image analysis for diagnosing molar and nonmolar products of conception. <i>American Journal of Clinical Pathology</i> , 2010 , 133, 196-204	1.9	24
328	Image analysis of HER2 immunohistochemical staining. Reproducibility and concordance with fluorescence in situ hybridization of a laboratory-validated scoring technique. <i>American Journal of Clinical Pathology</i> , 2012 , 137, 270-6	1.9	24
327	Isolated trisomy 15: a clonal chromosome abnormality in bone marrow with doubtful hematologic significance. <i>American Journal of Clinical Pathology</i> , 2008 , 129, 478-85	1.9	24
326	Recurrent mutations in ALK-negative anaplastic large cell lymphoma. <i>Blood</i> , 2019 , 133, 2776-2789	2.2	23
325	Epidemiology of adult acute myeloid leukemia: Impact of exposures on clinical phenotypes and outcomes after therapy. <i>Cancer Epidemiology</i> , 2015 , 39, 1084-92	2.8	23
324	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
323	An MLL-SEPT9 fusion and t(11;17)(q23;q25) associated with de novo myelodysplastic syndrome. <i>Leukemia Research</i> , 2007 , 31, 1145-8	2.7	23
322	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
321	Natural history of multiple myeloma with de novo del(17p). <i>Blood Cancer Journal</i> , 2019 , 9, 32	7	22
320	Polyclonal immunoglobulin free light chain levels predict survival in myeloid neoplasms. <i>Journal of Clinical Oncology</i> , 2012 , 30, 1087-94	2.2	22

319	Splenic small B-cell lymphoma with IGH/BCL3 translocation. <i>Human Pathology</i> , 2006 , 37, 218-30	3.7	22
318	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
317	Isochromosome 12p and polysomy 12 in primary central nervous system germ cell tumors: frequency and association with clinicopathologic features. <i>Human Pathology</i> , 2010 , 41, 232-8	3.7	21
316	Chromosome 1 abnormalities in myeloid malignancies: a literature survey and karyotype-phenotype associations. <i>European Journal of Haematology</i> , 2010 , 84, 191-200	3.8	21
315	Chromosome 9p24 abnormalities: prevalence, description of novel JAK2 translocations, JAK2V617F mutation analysis and clinicopathologic correlates. <i>European Journal of Haematology</i> , 2010 , 84, 518-24	3.8	21
314	Deletions with inversions: report of a mutation and review of the literature. <i>Human Mutation</i> , 1993 , 2, 53-7	4.7	21
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