Mark J Levis

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.

12,168 184 109 57 h-index g-index citations papers 6.57 4.8 14,192 191 L-index avg, IF ext. citations ext. papers

#	Paper	IF	Citations
184	Follow-up of patients with R/R FLT3-mutation-positive AML treated with gilteritinib in the phase 3 ADMIRAL trial <i>Blood</i> , 2022 ,	2.2	3
183	The role of the atypical chemokine receptor CCRL2 in myelodysplastic syndrome and secondary acute myeloid leukemia <i>Science Advances</i> , 2022 , 8, eabl8952	14.3	1
182	The Presence of SETBP1, RUNX1 or EZH2 Mutation in MDS/MPN Is Associated with Absence of Response to Hypo-Methylating Agents. <i>Blood</i> , 2021 , 138, 1520-1520	2.2	
181	The impact of FLT3 mutation clearance and treatment response after gilteritinib therapy on overall survival in patients with FLT3 mutation-positive relapsed/refractory acute myeloid leukemia. <i>Cancer Medicine</i> , 2021 , 10, 797-805	4.8	2
180	Is there evidence for the use of FLT3 inhibitors as maintenance therapy in AML?. <i>Best Practice and Research in Clinical Haematology</i> , 2021 , 34, 101246	4.2	
179	Single-cell DNA sequencing reveals complex mechanisms of resistance to quizartinib. <i>Blood Advances</i> , 2021 , 5, 1437-1441	7.8	0
178	A phase I/II study of the combination of quizartinib with azacitidine or low-dose cytarabine for the treatment of patients with acute myeloid leukemia and myelodysplastic syndrome. <i>Haematologica</i> , 2021 , 106, 2121-2130	6.6	10
177	Deletions in FLT-3 juxtamembrane domain define a new class of pathogenic mutations: case report and systematic analysis. <i>Blood Advances</i> , 2021 , 5, 2285-2293	7.8	4
176	Arsenic and all-retinoic acid for acute promyelocytic leukemia: yes, it really is as good as it seems. <i>Haematologica</i> , 2021 , 106, 3031-3032	6.6	
175	FLT3 tyrosine kinase inhibitors synergize with BCL-2 inhibition to eliminate FLT3/ITD acute leukemia cells through BIM activation. <i>Signal Transduction and Targeted Therapy</i> , 2021 , 6, 186	21	7
174	Deep learning for diagnosis of acute promyelocytic leukemia via recognition of genomically imprinted morphologic features. <i>Npj Precision Oncology</i> , 2021 , 5, 38	9.8	4
173	Arsenic trioxide dose capping to decrease toxicity in the treatment of acute promyelocytic leukemia. <i>Journal of Oncology Pharmacy Practice</i> , 2021 , 10781552211024727	1.7	1
172	A method for overcoming plasma protein inhibition of tyrosine kinase inhibitors. <i>Blood Cancer Discovery</i> , 2021 , 2, 532-547	7	O
171	Potential targeting of FLT3 acute myeloid leukemia. <i>Haematologica</i> , 2021 , 106, 671-681	6.6	11
170	Clinical Outcomes in Patients with FLT3-ITD-Mutated Relapsed/Refractory Acute Myelogenous Leukemia Undergoing Hematopoietic Stem Cell Transplantation after Quizartinib or Salvage Chemotherapy in the QuANTUM-R Trial. <i>Transplantation and Cellular Therapy</i> , 2021 , 27, 153-162		5
169	Midostaurin after allogeneic stem cell transplant in patients with FLT3-internal tandem duplication-positive acute myeloid leukemia. <i>Bone Marrow Transplantation</i> , 2021 , 56, 1180-1189	4.4	22
168	Concentration-QTc analysis of quizartinib in patients with relapsed/refractory acute myeloid leukemia. <i>Cancer Chemotherapy and Pharmacology</i> , 2021 , 87, 513-523	3.5	1

(2020-2021)

167	Characteristics and outcome of patients with acute myeloid leukaemia and t(8;16)(p11;p13): results from an International Collaborative Study. <i>British Journal of Haematology</i> , 2021 , 192, 832-842	4.5	4
166	Differentiation syndrome with lower-intensity treatments for acute myeloid leukemia. <i>American Journal of Hematology</i> , 2021 , 96, 735-746	7.1	2
165	Updates on targeted therapies for acute myeloid leukaemia. British Journal of Haematology, 2021,	4.5	10
164	Phase II Trial of Pembrolizumab after High-Dose Cytarabine in Relapsed/Refractory Acute Myeloid Leukemia. <i>Blood Cancer Discovery</i> , 2021 , 2, 616-629	7	8
163	Phase 1 study of the histone deacetylase inhibitor entinostat plus clofarabine for poor-risk Philadelphia chromosome-negative (newly diagnosed older adults or adults with relapsed refractory disease) acute lymphoblastic leukemia or biphenotypic leukemia. <i>Leukemia Research</i> ,	2.7	1
162	2021 , 110, 106707 A novel combination regimen of BET and FLT3 inhibition for FLT3-ITD acute myeloid leukemia. Haematologica, 2021 , 106, 1022-1033	6.6	1
161	FLT3 inhibitors added to induction therapy induce deeper remissions. <i>Blood</i> , 2020 , 135, 75-78	2.2	13
160	Population Pharmacokinetic Analysis of Quizartinib in Healthy Volunteers and Patients With Relapsed/Refractory Acute Myeloid Leukemia. <i>Journal of Clinical Pharmacology</i> , 2020 , 60, 1629-1641	2.9	1
159	Outcome of older (🛮 0 years) APL patients frontline treated with or without arsenic trioxide-an International Collaborative Study. <i>Leukemia</i> , 2020 , 34, 2333-2341	10.7	13
158	A phase 1/2 study of the oral FLT3 inhibitor pexidartinib in relapsed/refractory FLT3-ITD-mutant acute myeloid leukemia. <i>Blood Advances</i> , 2020 , 4, 1711-1721	7.8	12
157	Pharmacokinetic Profile of Gilteritinib: A Novel FLT-3 Tyrosine Kinase Inhibitor. <i>Clinical Pharmacokinetics</i> , 2020 , 59, 1273-1290	6.2	15
156	Quizartinib, a selective FLT3 inhibitor, maintains antileukemic activity in preclinical models of RAS-mediated midostaurin-resistant acute myeloid leukemia cells. <i>Oncotarget</i> , 2020 , 11, 943-955	3.3	10
155	A Prospective Study of Peritransplant Sorafenib for Patients with FLT3-ITD Acute Myeloid Leukemia Undergoing Allogeneic Transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2020 , 26, 300-306	4.7	24
154	A Phase II Study of Midostaurin and 5-Azacitidine for Untreated Elderly and Unfit Patients With FLT3 Wild-type Acute Myelogenous Leukemia. <i>Clinical Lymphoma, Myeloma and Leukemia</i> , 2020 , 20, 226	5-233.e	17
153	Immunomodulation with pomalidomide at early lymphocyte recovery after induction chemotherapy in newly diagnosed AML and high-risk MDS. <i>Leukemia</i> , 2020 , 34, 1563-1576	10.7	7
152	Allogeneic transplantation for Ph+ acute lymphoblastic leukemia with posttransplantation cyclophosphamide. <i>Blood Advances</i> , 2020 , 4, 5078-5088	7.8	6
151	Allogeneic hematopoietic cell transplantation improves outcome of adults with t(6;9) acute myeloid leukemia: results from an international collaborative study. <i>Haematologica</i> , 2020 , 105, 161-169	6.6	8
150	Gilteritinib: potent targeting of FLT3 mutations in AML. <i>Blood Advances</i> , 2020 , 4, 1178-1191	7.8	35

149	The K666N mutation in SF3B1 is associated with increased progression of MDS and distinct RNA splicing. <i>Blood Advances</i> , 2020 , 4, 1192-1196	7.8	18
148	Gilteritinib or Chemotherapy for Relapsed or Refractory -Mutated AML. <i>New England Journal of Medicine</i> , 2019 , 381, 1728-1740	59.2	413
147	Quizartinib versus salvage chemotherapy in relapsed or refractory FLT3-ITD acute myeloid leukaemia (QuANTUM-R): a multicentre, randomised, controlled, open-label, phase 3 trial. <i>Lancet Oncology, The</i> , 2019 , 20, 984-997	21.7	182
146	FLT3 Inhibitor Maintenance After Allogeneic Transplantation: Is a Placebo-Controlled, Randomized Trial Ethical?. <i>Journal of Clinical Oncology</i> , 2019 , 37, 1604-1607	2.2	19
145	Arsenic and old. <i>Blood</i> , 2019 , 133, 1392-1393	2.2	
144	Serum Flt3 ligand is a biomarker of progenitor cell mass and prognosis in acute myeloid leukemia. <i>Blood Advances</i> , 2019 , 3, 3052-3061	7.8	8
143	Role of CYP3A4 in bone marrow microenvironment-mediated protection of FLT3/ITD AML from tyrosine kinase inhibitors. <i>Blood Advances</i> , 2019 , 3, 908-916	7.8	25
142	Hedgehog/GLI1 activation leads to leukemic transformation of myelodysplastic syndrome in vivo and GLI1 inhibition results in antitumor activity. <i>Oncogene</i> , 2019 , 38, 687-698	9.2	11
141	Clinical implications of molecular markers in acute myeloid leukemia. <i>European Journal of Haematology</i> , 2019 , 102, 20-35	3.8	26
140	Targeting FLT3 mutations in AML: review of current knowledge and evidence. <i>Leukemia</i> , 2019 , 33, 299	-3 1 2. ₇	324
140	Targeting FLT3 mutations in AML: review of current knowledge and evidence. <i>Leukemia</i> , 2019 , 33, 299. Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325	-31 2 . ₇	324
	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in</i>		•
139	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325 Emerging molecular predictive and prognostic factors in acute myeloid leukemia. <i>Leukemia and</i>	0.6	1
139	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325 Emerging molecular predictive and prognostic factors in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2018 , 59, 2021-2039 Cabozantinib is well tolerated in acute myeloid leukemia and effectively inhibits the	0.6	7
139 138 137	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325 Emerging molecular predictive and prognostic factors in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2018 , 59, 2021-2039 Cabozantinib is well tolerated in acute myeloid leukemia and effectively inhibits the resistance-conferring FLT3/tyrosine kinase domain/F691 mutation. <i>Cancer</i> , 2018 , 124, 306-314 A next-generation sequencing-based assay for minimal residual disease assessment in AML patients	0.6	1 7 17
139 138 137	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325 Emerging molecular predictive and prognostic factors in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2018 , 59, 2021-2039 Cabozantinib is well tolerated in acute myeloid leukemia and effectively inhibits the resistance-conferring FLT3/tyrosine kinase domain/F691 mutation. <i>Cancer</i> , 2018 , 124, 306-314 A next-generation sequencing-based assay for minimal residual disease assessment in AML patients with -ITD mutations. <i>Blood Advances</i> , 2018 , 2, 825-831 Sorafenib Combined with 5-azacytidine in Older Patients with Untreated FLT3-ITD Mutated Acute	0.61.96.47.8	1 7 17 69
139 138 137 136	Midostaurin for patients with acute myeloid leukemia and FLT3 mutations. <i>Clinical Advances in Hematology and Oncology</i> , 2019 , 17, 323-325 Emerging molecular predictive and prognostic factors in acute myeloid leukemia. <i>Leukemia and Lymphoma</i> , 2018 , 59, 2021-2039 Cabozantinib is well tolerated in acute myeloid leukemia and effectively inhibits the resistance-conferring FLT3/tyrosine kinase domain/F691 mutation. <i>Cancer</i> , 2018 , 124, 306-314 A next-generation sequencing-based assay for minimal residual disease assessment in AML patients with -ITD mutations. <i>Blood Advances</i> , 2018 , 2, 825-831 Sorafenib Combined with 5-azacytidine in Older Patients with Untreated FLT3-ITD Mutated Acute Myeloid Leukemia. <i>American Journal of Hematology</i> , 2018 , 93, 1136-1141	0.61.96.47.87.1	1 7 17 69 54

(2017-2018)

131	Combination of ATO with FLT3 TKIs eliminates FLT3/ITD+ leukemia cells through reduced expression of FLT3. <i>Oncotarget</i> , 2018 , 9, 32885-32899	3.3	12
130	Advances in targeted therapy for acute myeloid leukaemia. <i>British Journal of Haematology</i> , 2018 , 180, 484-500	4.5	59
129	Quizartinib, an FLT3 inhibitor, as monotherapy in patients with relapsed or refractory acute myeloid leukaemia: an open-label, multicentre, single-arm, phase 2 trial. <i>Lancet Oncology, The</i> , 2018 , 19, 889-903	21.7	145
128	FLT3 as a marker of minimal residual disease: Time to re-think?. <i>American Journal of Hematology</i> , 2017 , 92, 329-330	7.1	4
127	Comparable composite endpoints after HLA-matched and HLA-haploidentical transplantation with post-transplantation cyclophosphamide. <i>Haematologica</i> , 2017 , 102, 391-400	6.6	119
126	A randomized assessment of adding the kinase inhibitor lestaurtinib to first-line chemotherapy for FLT3-mutated AML. <i>Blood</i> , 2017 , 129, 1143-1154	2.2	99
125	The Future of Targeting FLT3 Activation in AML. Current Hematologic Malignancy Reports, 2017, 12, 153-	-4647	32
124	Heterogeneous resistance to quizartinib in acute myeloid leukemia revealed by single-cell analysis. <i>Blood</i> , 2017 , 130, 48-58	2.2	100
123	Selective inhibition of FLT3 by gilteritinib in relapsed or refractory acute myeloid leukaemia: a multicentre, first-in-human, open-label, phase 1-2 study. <i>Lancet Oncology, The</i> , 2017 , 18, 1061-1075	21.7	305
122	Mechanisms of Resistance to FLT3 Inhibitors and the Role of the Bone Marrow Microenvironment. Hematology/Oncology Clinics of North America, 2017 , 31, 681-692	3.1	31
121	Preclinical studies of gilteritinib, a next-generation FLT3 inhibitor. <i>Blood</i> , 2017 , 129, 257-260	2.2	144
120	How I treat FLT3-mutated AML. <i>Blood</i> , 2017 , 129, 565-571	2.2	52
119	Midostaurin treatment in FLT3-mutated acute myeloid leukemia and systemic mastocytosis. <i>Expert Review of Clinical Pharmacology</i> , 2017 , 10, 1177-1189	3.8	15
118	Adaptation to TKI Treatment Reactivates ERK Signaling in Tyrosine Kinase-Driven Leukemias and Other Malignancies. <i>Cancer Research</i> , 2017 , 77, 5554-5563	10.1	22
117	Timed sequential therapy for acute myelogenous leukemia: Results of a retrospective study of 301 patients and review of the literature. <i>Leukemia Research</i> , 2017 , 61, 25-32	2.7	12
116	FLT3 dancing on the stem cell. <i>Journal of Experimental Medicine</i> , 2017 , 214, 1857-1859	16.6	2
115	Midostaurin approved for FLT3-mutated AML. <i>Blood</i> , 2017 , 129, 3403-3406	2.2	145
114	A Phase 1 Study of the PARP Inhibitor Veliparib in Combination with Temozolomide in Acute Myeloid Leukemia. <i>Clinical Cancer Research</i> , 2017 , 23, 697-706	12.9	40

113	FLT3 activating mutations display differential sensitivity to multiple tyrosine kinase inhibitors. <i>Oncotarget</i> , 2017 , 8, 10931-10944	3.3	22
112	Prospective study of nonmyeloablative, HLA-mismatched unrelated BMT with high-dose posttransplantation cyclophosphamide. <i>Blood Advances</i> , 2017 , 1, 288-292	7.8	58
111	A phase 2 study incorporating sorafenib into the chemotherapy for older adults with -mutated acute myeloid leukemia: CALGB 11001. <i>Blood Advances</i> , 2017 , 1, 331-340	7.8	42
110	The current therapeutic landscape of FLT3 inhibitors. <i>Blood Advances</i> , 2017 , 1, 1944	7.8	1
109	Donor cell leukemia arising from clonal hematopoiesis after bone marrow transplantation. <i>Leukemia</i> , 2016 , 30, 1916-1920	10.7	59
108	Signaling Adaptation to TKI Treatment Reactivates ERK Signaling in FLT3/ITD Leukemia. <i>Blood</i> , 2016 , 128, 33-33	2.2	2
107	Laboratory and Clinical Investigations to Identify the Optimal Dosing Strategy for Quizartinib (AC220) Monotherapy in FLT3-Itd-Positive (+) Relapsed/Refractory (R/R) Acute Myeloid Leukemia (AML). <i>Blood</i> , 2016 , 128, 4042-4042	2.2	14
106	Crenolanib besylate, a type I pan-FLT3 inhibitor, to demonstrate clinical activity in multiply relapsed FLT3-ITD and D835 AML <i>Journal of Clinical Oncology</i> , 2016 , 34, 7008-7008	2.2	55
105	Pre-Clinical Activity of Novel Hypoxia-Activated FLT3 Inhibitors in FLT3-Mutated AML. <i>Blood</i> , 2016 , 128, 5210-5210	2.2	1
104	Carotidynia Heralding the Onset of Acute Leukemia. <i>American Journal of Medicine</i> , 2016 , 129, e43-5	2.4	2
103	Integration of Hedgehog and mutant FLT3 signaling in myeloid leukemia. <i>Science Translational Medicine</i> , 2015 , 7, 291ra96	17.5	37
102	FLT3 Tyrosine Kinase Inhibition as a Paradigm for Targeted Drug Development in Acute Myeloid Leukemia. <i>Seminars in Hematology</i> , 2015 , 52, 193-9	4	32
101	Inhibition of c-Kit by tyrosine kinase inhibitors. <i>Haematologica</i> , 2015 , 100, e77-9	6.6	76
100	Outcomes of Nonmyeloablative HLA-Haploidentical Blood or Marrow Transplantation With High-Dose Post-Transplantation Cyclophosphamide in Older Adults. <i>Journal of Clinical Oncology</i> , 2015 , 33, 3152-61	2.2	165
99	A Novel Tandem Duplication Assay to Detect Minimal Residual Disease in FLT3/ITD AML. <i>Molecular Diagnosis and Therapy</i> , 2015 , 19, 409-17	4.5	8
98	Targeting FLT3 to treat leukemia. Expert Opinion on Therapeutic Targets, 2015, 19, 37-54	6.4	60
97	Phase I/II trial of the combination of midostaurin (PKC412) and 5-azacytidine for patients with acute myeloid leukemia and myelodysplastic syndrome. <i>American Journal of Hematology</i> , 2015 , 90, 276-	·871 ¹	114
96	Risk-stratified outcomes of nonmyeloablative HLA-haploidentical BMT with high-dose posttransplantation cyclophosphamide. <i>Blood</i> , 2015 , 125, 3024-31	2.2	212

(2014-2015)

95	with High-Dose Post-Transplant Cyclophosphmade (PTCy) for Acute Myeloid Leukemia (AML): Donor Age Impacts Outcome. <i>Blood</i> , 2015 , 126, 151-151	2.2	4
94	Addition of Sorafenib to Chemotherapy Improves the Overall Survival of Older Adults with FLT3-ITD Mutated Acute Myeloid Leukemia (AML) (Alliance C11001). <i>Blood</i> , 2015 , 126, 319-319	2.2	9
93	Pharmacokinetic Profile and Pharmacodynamic Effects of ASP2215, a Selective, Potent Inhibitor of FLT3/AXL, in Patients with Relapsed or Refractory Acute Myeloid Leukemia: Results from a First-in-Human Phase 1/2 Study. <i>Blood</i> , 2015 , 126, 4836-4836	2.2	4
92	Liberal Vs. Restrictive Transfusion Thresholds in Leukemia Patients: A Feasibility Pilot Study. <i>Blood</i> , 2015 , 126, 771-771	2.2	2
91	Results of a first-in-human, phase I/II trial of ASP2215, a selective, potent inhibitor of FLT3/Axl in patients with relapsed or refractory (R/R) acute myeloid leukemia (AML) <i>Journal of Clinical Oncology</i> , 2015 , 33, 7003-7003	2.2	27
90	TTT-3002 is a novel FLT3 tyrosine kinase inhibitor with activity against FLT3-associated leukemias in vitro and in vivo. <i>Blood</i> , 2014 , 123, 1525-34	2.2	18
89	Phase I trial of maintenance sorafenib after allogeneic hematopoietic stem cell transplantation for fms-like tyrosine kinase 3 internal tandem duplication acute myeloid leukemia. <i>Biology of Blood and Marrow Transplantation</i> , 2014 , 20, 2042-8	4.7	178
88	Improved FLT3 internal tandem duplication PCR assay predicts outcome after allogeneic transplant for acute myeloid leukemia. <i>Biology of Blood and Marrow Transplantation</i> , 2014 , 20, 1989-95	4.7	25
87	Bone marrow stroma-mediated resistance to FLT3 inhibitors in FLT3-ITD AML is mediated by persistent activation of extracellular regulated kinase. <i>British Journal of Haematology</i> , 2014 , 164, 61-72	4.5	72
86	Is targeted therapy feasible in acute myelogenous leukemia?. <i>Current Hematologic Malignancy Reports</i> , 2014 , 9, 118-27	4.4	8
85	FLT3 tyrosine kinase inhibitors in acute myeloid leukemia: clinical implications and limitations.		70
	Leukemia and Lymphoma, 2014 , 55, 243-55	1.9	, -
84	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants. Blood, 2014, 123, 94-100	2.2	175
	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants.		
84	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants. <i>Blood</i> , 2014 , 123, 94-100 Single-agent GVHD prophylaxis with posttransplantation cyclophosphamide after myeloablative, HLA-matched BMT for AML, ALL, and MDS. <i>Blood</i> , 2014 , 124, 3817-27	2.2	175
84	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants. <i>Blood</i> , 2014 , 123, 94-100 Single-agent GVHD prophylaxis with posttransplantation cyclophosphamide after myeloablative, HLA-matched BMT for AML, ALL, and MDS. <i>Blood</i> , 2014 , 124, 3817-27 FLT3 kinase inhibitor TTT-3002 overcomes both activating and drug resistance mutations in FLT3 in acute myeloid leukemia. <i>Cancer Research</i> , 2014 , 74, 5206-17	2.2	175
84 83 82	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants. <i>Blood</i> , 2014 , 123, 94-100 Single-agent GVHD prophylaxis with posttransplantation cyclophosphamide after myeloablative, HLA-matched BMT for AML, ALL, and MDS. <i>Blood</i> , 2014 , 124, 3817-27 FLT3 kinase inhibitor TTT-3002 overcomes both activating and drug resistance mutations in FLT3 in acute myeloid leukemia. <i>Cancer Research</i> , 2014 , 74, 5206-17	2.2	175 128 18
84 83 82 81	Crenolanib is a potent inhibitor of FLT3 with activity against resistance-conferring point mutants. <i>Blood</i> , 2014 , 123, 94-100 Single-agent GVHD prophylaxis with posttransplantation cyclophosphamide after myeloablative, HLA-matched BMT for AML, ALL, and MDS. <i>Blood</i> , 2014 , 124, 3817-27 FLT3 kinase inhibitor TTT-3002 overcomes both activating and drug resistance mutations in FLT3 in acute myeloid leukemia. <i>Cancer Research</i> , 2014 , 74, 5206-17 Quizartinib for the treatment of FLT3/ITD acute myeloid leukemia. <i>Future Oncology</i> , 2014 , 10, 1571-9 The evolving role of FLT3 inhibitors in acute myeloid leukemia: quizartinib and beyond. <i>Therapeutic</i>	2.2 2.2 10.1 3.6	175 128 18 38

77	Advances in treating acute myeloid leukemia. F1000prime Reports, 2014, 6, 96		25
76	FLT3 inhibitors for acute myeloid leukemia: a review of their efficacy and mechanisms of resistance. <i>International Journal of Hematology</i> , 2013 , 97, 683-94	2.3	127
75	FLT3 mutations in acute myeloid leukemia: what is the best approach in 2013?. Hematology American Society of Hematology Education Program, 2013, 2013, 220-6	3.1	164
74	Phase I study of quizartinib administered daily to patients with relapsed or refractory acute myeloid leukemia irrespective of FMS-like tyrosine kinase 3-internal tandem duplication status. <i>Journal of Clinical Oncology</i> , 2013 , 31, 3681-7	2.2	278
73	FLT3 inhibitor-induced neutrophilic dermatosis. <i>Blood</i> , 2013 , 122, 239-42	2.2	35
72	Phase 2 study of azacytidine plus sorafenib in patients with acute myeloid leukemia and FLT-3 internal tandem duplication mutation. <i>Blood</i> , 2013 , 121, 4655-62	2.2	296
71	Are FLT3 inhibitors likely to improve FLT3-mutated acute myeloid leukemia in the foreseeable future?. <i>International Journal of Hematologic Oncology</i> , 2013 , 2, 9-11	1	1
70	Results Of a Phase 2 Randomized, Open-Label, Study Of Lower Doses Of Quizartinib (AC220; ASP2689) In Subjects With FLT3-ITD Positive Relapsed Or Refractory Acute Myeloid Leukemia (AML). <i>Blood</i> , 2013 , 122, 494-494	2.2	31
69	Effect of quizartinib (AC220) on response rates and long-term survival in elderly patients with FLT3-ITD positive or negative relapsed/refractory acute myeloid leukemia <i>Journal of Clinical Oncology</i> , 2013 , 31, 7021-7021	2.2	4
68	Efficacy and safety of quizartinib (AC220) in patients age I70 years with FLT3-ITD positive or negative relapsed/refractory acute myeloid leukemia (AML) <i>Journal of Clinical Oncology</i> , 2013 , 31, 702	23 -7 02:	3 3
68 67	Efficacy and safety of quizartinib (AC220) in patients age I70 years with FLT3-ITD positive or negative relapsed/refractory acute myeloid leukemia (AML) <i>Journal of Clinical Oncology</i> , 2013 , 31, 7026-7026 Variations in FLT3 ligand levels during the course of AML treatment <i>Journal of Clinical Oncology</i> , 2013 , 31, 7026-7026	23 ² 7022 2.2	3 3
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67	negative relapsed/refractory acute myeloid leukemia (AML) <i>Journal of Clinical Oncology</i> , 2013 , 31, 702 Variations in FLT3 ligand levels during the course of AML treatment <i>Journal of Clinical Oncology</i> , 2013 , 31, 7026-7026		
6 ₇	negative relapsed/refractory acute myeloid leukemia (AML) Journal of Clinical Oncology, 2013, 31, 702 Variations in FLT3 ligand levels during the course of AML treatment Journal of Clinical Oncology, 2013, 31, 7026-7026 Quizartinib in acute myeloid leukemia. Clinical Advances in Hematology and Oncology, 2013, 11, 586-8	0.6	9
67 66 65	Nariations in FLT3 ligand levels during the course of AML treatment <i>Journal of Clinical Oncology</i> , 2013 , 31, 7026-7026 Quizartinib in acute myeloid leukemia. <i>Clinical Advances in Hematology and Oncology</i> , 2013 , 11, 586-8 A potential therapeutic target for FLT3-ITD AML: PIM1 kinase. <i>Leukemia Research</i> , 2012 , 36, 224-31 Phase 1 dose-escalation trial of clofarabine followed by escalating dose of fractionated cyclophosphamide in adults with relapsed or refractory acute leukaemias. <i>British Journal of</i>	2.2 0.6 2.7	9 45
67 66 65 64	negative relapsed/refractory acute myeloid leukemia (AML) Journal of Clinical Oncology, 2013, 31, 702 Variations in FLT3 ligand levels during the course of AML treatment Journal of Clinical Oncology, 2013, 31, 7026-7026 Quizartinib in acute myeloid leukemia. Clinical Advances in Hematology and Oncology, 2013, 11, 586-8 A potential therapeutic target for FLT3-ITD AML: PIM1 kinase. Leukemia Research, 2012, 36, 224-31 Phase 1 dose-escalation trial of clofarabine followed by escalating dose of fractionated cyclophosphamide in adults with relapsed or refractory acute leukaemias. British Journal of Haematology, 2012, 158, 198-207 Terminal myeloid differentiation in vivo is induced by FLT3 inhibition in FLT3/ITD AML. Blood, 2012,	2.20.62.74.5	9456
6766656463	Nariations in FLT3 ligand levels during the course of AML treatment <i>Journal of Clinical Oncology</i> , 2013 , 31, 7026-7026 Quizartinib in acute myeloid leukemia. <i>Clinical Advances in Hematology and Oncology</i> , 2013 , 11, 586-8 A potential therapeutic target for FLT3-ITD AML: PIM1 kinase. <i>Leukemia Research</i> , 2012 , 36, 224-31 Phase 1 dose-escalation trial of clofarabine followed by escalating dose of fractionated cyclophosphamide in adults with relapsed or refractory acute leukaemias. <i>British Journal of Haematology</i> , 2012 , 158, 198-207 Terminal myeloid differentiation in vivo is induced by FLT3 inhibition in FLT3/ITD AML. <i>Blood</i> , 2012 , 120, 4205-14 Validation of ITD mutations in FLT3 as a therapeutic target in human acute myeloid leukaemia.	2.2 0.6 2.7 4.5	19456122

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56	Final Results of a Phase 2 Open-Label, Monotherapy Efficacy and Safety Study of Quizartinib (AC220) in Patients 160 Years of Age with FLT3 ITD Positive or Negative Relapsed/Refractory Acute Myeloid Leukemia. <i>Blood</i> , 2012 , 120, 48-48	2.2	54
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12	FLT3 ligand causes autocrine signaling in acute myeloid leukemia cells. <i>Blood</i> , 2004 , 103, 267-74 Internal tandem duplication mutation of FLT3 blocks myeloid differentiation through suppression	2.2	166
12	FLT3 ligand causes autocrine signaling in acute myeloid leukemia cells. <i>Blood</i> , 2004 , 103, 267-74 Internal tandem duplication mutation of FLT3 blocks myeloid differentiation through suppression of C/EBPalpha expression. <i>Blood</i> , 2004 , 103, 1883-90 Single-agent CEP-701, a novel FLT3 inhibitor, shows biologic and clinical activity in patients with	2.2	166
12 11 10	FLT3 ligand causes autocrine signaling in acute myeloid leukemia cells. <i>Blood</i> , 2004 , 103, 267-74 Internal tandem duplication mutation of FLT3 blocks myeloid differentiation through suppression of C/EBPalpha expression. <i>Blood</i> , 2004 , 103, 1883-90 Single-agent CEP-701, a novel FLT3 inhibitor, shows biologic and clinical activity in patients with relapsed or refractory acute myeloid leukemia. <i>Blood</i> , 2004 , 103, 3669-76 In vitro studies of a FLT3 inhibitor combined with chemotherapy: sequence of administration is	2.2	166 132 542
12 11 10 9	FLT3 ligand causes autocrine signaling in acute myeloid leukemia cells. <i>Blood</i> , 2004 , 103, 267-74 Internal tandem duplication mutation of FLT3 blocks myeloid differentiation through suppression of C/EBPalpha expression. <i>Blood</i> , 2004 , 103, 1883-90 Single-agent CEP-701, a novel FLT3 inhibitor, shows biologic and clinical activity in patients with relapsed or refractory acute myeloid leukemia. <i>Blood</i> , 2004 , 103, 3669-76 In vitro studies of a FLT3 inhibitor combined with chemotherapy: sequence of administration is important to achieve synergistic cytotoxic effects. <i>Blood</i> , 2004 , 104, 1145-50 Pediatric AML primary samples with FLT3/ITD mutations are preferentially killed by FLT3 inhibition.	2.2 2.2 2.2	166 132 542 192

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1	Novel FLT3 tyrosine kinase inhibitors		2