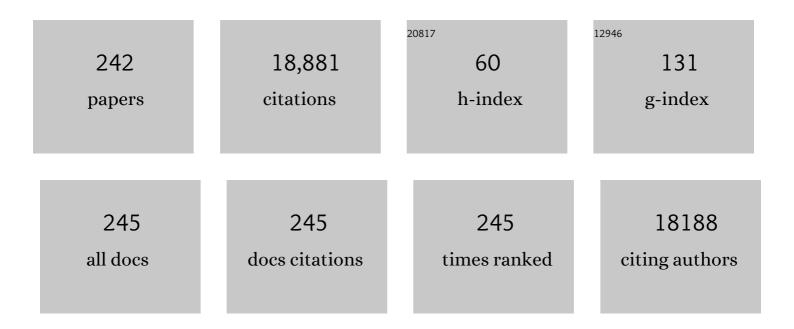
Carlo Gambacorti-Passerini

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Hematologic and Cytogenetic Responses to Imatinib Mesylate in Chronic Myelogenous Leukemia. New England Journal of Medicine, 2002, 346, 645-652.	27.0	1,899
2	Clinical and biological implications of driver mutations in myelodysplastic syndromes. Blood, 2013, 122, 3616-3627.	1.4	1,562
3	Imatinib induces durable hematologic and cytogenetic responses in patients with accelerated phase chronic myeloid leukemia: results of a phase 2 study. Blood, 2002, 99, 1928-1937.	1.4	943
4	Induction of resistance to the Abelson inhibitor STI571 in human leukemic cells through gene amplification. Blood, 2000, 95, 1758-1766.	1.4	454
5	Selective cytotoxicity of betulinic acid on tumor cell lines, but not on normal cells. Cancer Letters, 2002, 175, 17-25.	7.2	441
6	Clinical characteristics and risk factors associated with COVID-19 severity in patients with haematological malignancies in Italy: a retrospective, multicentre, cohort study. Lancet Haematology,the, 2020, 7, e737-e745.	4.6	430
7	Safety and efficacy of bosutinib (SKI-606) in chronic phase Philadelphia chromosome–positive chronic myeloid leukemia patients with resistance or intolerance to imatinib. Blood, 2011, 118, 4567-4576.	1.4	406
8	Bosutinib Versus Imatinib in Newly Diagnosed Chronic-Phase Chronic Myeloid Leukemia: Results From the BELA Trial. Journal of Clinical Oncology, 2012, 30, 3486-3492.	1.6	404
9	Ponatinib efficacy and safety in Philadelphia chromosome–positive leukemia: final 5-year results of the phase 2 PACE trial. Blood, 2018, 132, 393-404.	1.4	392
10	Activity of Bosutinib, Dasatinib, and Nilotinib Against 18 Imatinib-Resistant BCR/ABL Mutants. Journal of Clinical Oncology, 2009, 27, 469-471.	1.6	365
11	Multicenter Independent Assessment of Outcomes in Chronic Myeloid Leukemia Patients Treated With Imatinib. Journal of the National Cancer Institute, 2011, 103, 553-561.	6.3	362
12	Recurrent SETBP1 mutations in atypical chronic myeloid leukemia. Nature Genetics, 2013, 45, 18-24.	21.4	359
13	Bosutinib Versus Imatinib for Newly Diagnosed Chronic Myeloid Leukemia: Results From the Randomized BFORE Trial. Journal of Clinical Oncology, 2018, 36, 231-237.	1.6	356
14	In vitro and In vivo Activity of SKI-606, a Novel Src-Abl Inhibitor, against Imatinib-Resistant Bcr-Abl+ Neoplastic Cells. Cancer Research, 2006, 66, 11314-11322.	0.9	352
15	Molecular mechanisms of resistance to imatinib in Philadelphia-chromosome-positive leukaemias. Lancet Oncology, The, 2003, 4, 75-85.	10.7	349
16	In Vivo Eradication of Human BCR/ABL-Positive Leukemia Cells With an ABL Kinase Inhibitor. Journal of the National Cancer Institute, 1999, 91, 163-168.	6.3	341
17	FTY720, a new alternative for treating blast crisis chronic myelogenous leukemia and Philadelphia chromosome–positive acute lymphocytic leukemia. Journal of Clinical Investigation, 2007, 117, 2408-2421.	8.2	308
18	BCR-ABL suppresses C/EBPα expression through inhibitory action of hnRNP E2. Nature Genetics, 2002, 30, 48-58.	21.4	301

#	Article	IF	CITATIONS
19	Bosutinib is active in chronic phase chronic myeloid leukemia after imatinib and dasatinib and/or nilotinib therapy failure. Blood, 2012, 119, 3403-3412.	1.4	281
20	Inhibition of the ABL Kinase Activity Blocks the Proliferation of BCR/ABL+Leukemic Cells and Induces Apoptosis. Blood Cells, Molecules, and Diseases, 1997, 23, 380-394.	1.4	273
21	Crizotinib in Anaplastic Large-Cell Lymphoma. New England Journal of Medicine, 2011, 364, 775-776.	27.0	256
22	Dasatinib and low-intensity chemotherapy in elderly patients with Philadelphia chromosome–positive ALL. Blood, 2016, 128, 774-782.	1.4	243
23	Crizotinib in Advanced, Chemoresistant Anaplastic Lymphoma Kinase–Positive Lymphoma Patients. Journal of the National Cancer Institute, 2014, 106, djt378.	6.3	207
24	Bcr-Abl stabilizes β-catenin in chronic myeloid leukemia through its tyrosine phosphorylation. EMBO Journal, 2007, 26, 1456-1466.	7.8	204
25	Favorable long-term follow-up results over 6 years for response, survival, and safety with imatinib mesylate therapy in chronic-phase chronic myeloid leukemia after failure of interferon-α treatment. Blood, 2008, 111, 1039-1043.	1.4	195
26	Age and d <scp>PCR</scp> can predict relapse in <scp>CML</scp> patients who discontinued imatinib: The <scp>ISAV</scp> study. American Journal of Hematology, 2015, 90, 910-914.	4.1	181
27	BRAF Silencing by Short Hairpin RNA or Chemical Blockade by PLX4032 Leads to Different Responses in Melanoma and Thyroid Carcinoma Cells. Molecular Cancer Research, 2008, 6, 751-759.	3.4	178
28	Bosutinib <i>versus</i> imatinib in newly diagnosed chronicâ€phase chronic myeloid leukaemia: results from the 24â€month followâ€up of the BELA trial. British Journal of Haematology, 2015, 168, 69-81.	2.5	177
29	Alpha1 acid glycoprotein binds to imatinib (STI571) and substantially alters its pharmacokinetics in chronic myeloid leukemia patients. Clinical Cancer Research, 2003, 9, 625-32.	7.0	159
30	Mutation-Independent Anaplastic Lymphoma Kinase Overexpression in Poor Prognosis Neuroblastoma Patients. Cancer Research, 2009, 69, 7338-7346.	0.9	157
31	Bosutinib safety and management of toxicity in leukemia patients with resistance or intolerance to imatinib and other tyrosine kinase inhibitors. Blood, 2014, 123, 1309-1318.	1.4	124
32	The prognosis for patients with chronic myeloid leukemia who have clonal cytogenetic abnormalities in philadelphia chromosomeâ€negative cells. Cancer, 2007, 110, 1509-1519.	4.1	121
33	SKI-606 Decreases Growth and Motility of Colorectal Cancer Cells by Preventing pp60(c-Src)–Dependent Tyrosine Phosphorylation of β-Catenin and Its Nuclear Signaling. Cancer Research, 2006, 66, 2279-2286.	0.9	117
34	hnRNP A1 Nucleocytoplasmic Shuttling Activity Is Required for Normal Myelopoiesis and BCR/ABL Leukemogenesis. Molecular and Cellular Biology, 2002, 22, 2255-2266.	2.3	115
35	Recurrent ETNK1 mutations in atypical chronic myeloid leukemia. Blood, 2015, 125, 499-503.	1.4	115
36	Epigenetic silencing of BIM in glucocorticoid poor-responsive pediatric acute lymphoblastic leukemia, and its reversal by histone deacetylase inhibition. Blood, 2010, 116, 3013-3022.	1.4	110

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37	Bosutinib efficacy and safety in chronic phase chronic myeloid leukemia after imatinib resistance or intolerance: Minimum 24â€month followâ€up. American Journal of Hematology, 2014, 89, 732-742.	4.1	102
38	Validation of PDGFRÎ ² and c-Src tyrosine kinases as tumor/vessel targets in patients with multiple myeloma: preclinical efficacy of the novel, orally available inhibitor dasatinib. Blood, 2008, 112, 1346-1356.	1.4	99
39	Safety of bosutinib versus imatinib in the phase 3 BELA trial in newly diagnosed chronic phase chronic myeloid leukemia. American Journal of Hematology, 2014, 89, 947-953.	4.1	98
40	Laying the foundation for genomically-based risk assessment in chronic myeloid leukemia. Leukemia, 2019, 33, 1835-1850.	7.2	97
41	Three novel patientâ€derived BCR/ABL mutants show different sensitivity to second and third generation tyrosine kinase inhibitors. American Journal of Hematology, 2012, 87, E125-8.	4.1	93
42	Chronic myeloid leukemia: reminiscences and dreams. Haematologica, 2016, 101, 541-558.	3.5	92
43	Sorafenib Functions to Potently Suppress RET Tyrosine Kinase Activity by Direct Enzymatic Inhibition and Promoting RET Lysosomal Degradation Independent of Proteasomal Targeting. Journal of Biological Chemistry, 2007, 282, 29230-29240.	3.4	90
44	Longâ€ŧerm bosutinib for chronic phase chronic myeloid leukemia after failure of imatinib plus dasatinib and/or nilotinib. American Journal of Hematology, 2016, 91, 1206-1214.	4.1	90
45	ALK as a novel lymphoma-associated tumor antigen: identification of 2 HLA-A2.1–restricted CD8+ T-cell epitopes. Blood, 2002, 99, 2100-2106.	1.4	89
46	Gynaecomastia in men with chronic myeloid leukaemia after imatinib. Lancet, The, 2003, 361, 1954-1956.	13.7	88
47	Ponatinib is a potent inhibitor of wild-type and drug-resistant gatekeeper mutant RET kinase. Molecular and Cellular Endocrinology, 2013, 377, 1-6.	3.2	81
48	Crizotinib-Resistant NPM-ALK Mutants Confer Differential Sensitivity to Unrelated Alk Inhibitors. Molecular Cancer Research, 2013, 11, 122-132.	3.4	79
49	Longâ€term evaluation of cardiac and vascular toxicity in patients with Philadelphia chromosomeâ€positive leukemias treated with bosutinib. American Journal of Hematology, 2016, 91, 606-616.	4.1	76
50	Longâ€ŧerm effects of crizotinib in ALKâ€positive tumors (excluding NSCLC): A phase 1b openâ€label study. American Journal of Hematology, 2018, 93, 607-614.	4.1	75
51	Imatinib discontinuation in chronic myeloid leukaemia patients with undetectable BCR-ABL transcript level: AÂsystematic review and a meta-analysis. European Journal of Cancer, 2017, 77, 48-56.	2.8	74
52	Tumor Resistance against ALK Targeted Therapy-Where It Comes From and Where It Goes. Cancers, 2018, 10, 62.	3.7	73
53	Activity of secondâ€generation ALK inhibitors against crizotinibâ€resistant mutants in an NPMâ€ALK model compared to EML4â€ALK. Cancer Medicine, 2015, 4, 953-965.	2.8	72
54	Longâ€ŧerm efficacy and safety of bosutinib in patients with advanced leukemia following resistance/intolerance to imatinib and other tyrosine kinase inhibitors. American Journal of Hematology, 2015, 90, 755-768.	4.1	72

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55	Lorlatinib Treatment Elicits Multiple On- and Off-Target Mechanisms of Resistance in ALK-Driven Cancer. Cancer Research, 2018, 78, 6866-6880.	0.9	69
56	Inhibition of RET tyrosine kinase by SU5416. Journal of Molecular Endocrinology, 2006, 37, 199-212.	2.5	68
57	Abrupt Relapse of <i>ALK</i> -Positive Lymphoma after Discontinuation of Crizotinib. New England Journal of Medicine, 2016, 374, 95-96.	27.0	67
58	Panniculitis during Dasatinib Therapy for Imatinib-Resistant Chronic Myelogenous Leukemia. New England Journal of Medicine, 2006, 354, 2623-2624.	27.0	66
59	SETBP1 induces transcription of a network of development genes by acting as an epigenetic hub. Nature Communications, 2018, 9, 2192.	12.8	66
60	Synthesis, structure–activity relationship and crystallographic studies of 3-substituted indolin-2-one RET inhibitors. Bioorganic and Medicinal Chemistry, 2010, 18, 1482-1496.	3.0	64
61	Bosutinib: A review of preclinical studies in chronic myelogenous leukaemia. European Journal of Cancer, 2010, 46, 1781-1789.	2.8	62
62	Characterization of Some Molecular Mechanisms Governing Autoactivation of the Catalytic Domain of the Anaplastic Lymphoma Kinase. Journal of Biological Chemistry, 2008, 283, 3743-3750.	3.4	61
63	Constitutive activation of Jak2 contributes to proliferation and resistance to apoptosis in NPM/ALK-transformed cells. Experimental Hematology, 2003, 31, 309-315.	0.4	59
64	Observational study of chronic myeloid leukemia Italian patients who discontinued tyrosine kinase inhibitors in clinical practice. Haematologica, 2019, 104, 1589-1596.	3.5	58
65	Adoptive immunotherapy of advanced melanoma patients with interleukin-2 (IL-2) and tumor-infiltrating lymphocytes selected in vitro with low doses of IL-2. Cancer Immunology, Immunotherapy, 1993, 36, 315-322.	4.2	57
66	Adherence and future discontinuation of tyrosine kinase inhibitors in chronic phase chronic myeloid leukemia. A patient-based survey on 1133 patients. Leukemia Research, 2015, 39, 1055-1059.	0.8	57
67	Wiskott–Aldrich syndrome protein (WASP) is a tumor suppressor in T cell lymphoma. Nature Medicine, 2019, 25, 130-140.	30.7	57
68	COVIDâ€19 elicits an impaired antibody response against SARS oVâ€2 in patients with haematological malignancies. British Journal of Haematology, 2021, 195, 371-377.	2.5	56
69	c-MYC Generates Repair Errors via Increased Transcription of Alternative-NHEJ Factors, LIG3 and PARP1, in Tyrosine Kinase–Activated Leukemias. Molecular Cancer Research, 2015, 13, 699-712.	3.4	55
70	In reply to 'Cardiotoxicity of the cancer therapeutic agent imatinib mesylate'. Nature Medicine, 2007, 13, 13-14.	30.7	54
71	Unique Substrate Specificity of Anaplastic Lymphoma Kinase (ALK):  Development of Phosphoacceptor Peptides for the Assay of ALK Activity. Biochemistry, 2005, 44, 8533-8542.	2.5	53
72	BCR-ABL nuclear entrapment kills human CML cells: ex vivo study on 35 patients with the combination of imatinib mesylate and leptomycin B. Blood, 2006, 107, 1591-1598.	1.4	53

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73	Determination of α-1 Acid Glycoprotein in Patients with Ph+ Chronic Myeloid Leukemia during the First 13 Weeks of Therapy with STI571. Blood Cells, Molecules, and Diseases, 2002, 28, 75-85.	1.4	52
74	Treatment Efficacy and Resistance Mechanisms Using the Second-Generation ALK Inhibitor AP26113 in Human NPM-ALK–Positive Anaplastic Large Cell Lymphoma. Molecular Cancer Research, 2015, 13, 775-783.	3.4	52
75	Management of adverse events associated with bosutinib treatment of chronic-phase chronic myeloid leukemia: expert panel review. Journal of Hematology and Oncology, 2018, 11, 143.	17.0	52
76	In Vitro Transcriptional and Translational Block of the bcl-2 Gene Operated by Peptide Nucleic Acid. Biochemical and Biophysical Research Communications, 1999, 264, 537-543.	2.1	51
77	Part I: Milestones in personalised medicine—imatinib. Lancet Oncology, The, 2008, 9, 600.	10.7	51
78	Killer immunoglobulin-like receptors can predict TKI treatment-free remission in chronic myeloid leukemia patients. Experimental Hematology, 2015, 43, 1015-1018.e1.	0.4	51
79	Increased <scp>sFLTâ€I </scp> / <scp>PIGF</scp> ratio in <scp>COVID</scp> â€19: A novel link to angiotensin <scp>II</scp> â€mediated endothelial dysfunction. American Journal of Hematology, 2020, 95, E188-E191.	4.1	51
80	Focal Adhesion Kinase (FAK) Binds RET Kinase via Its FERM Domain, Priming a Direct and Reciprocal RET-FAK Transactivation Mechanism. Journal of Biological Chemistry, 2011, 286, 17292-17302.	3.4	50
81	Safety and efficacy of second-line bosutinib for chronic phase chronic myeloid leukemia over a five-year period: final results of a phase I/II study. Haematologica, 2018, 103, 1298-1307.	3.5	49
82	Inhibitors of the RET tyrosine kinase based on a 2-(alkylsulfanyl)-4-(3-thienyl)nicotinonitrile scaffold. European Journal of Medicinal Chemistry, 2010, 45, 2919-2927.	5.5	47
83	Bosutinib for pretreated patients with chronic phase chronic myeloid leukemia: primary results of the phase 4 BYOND study. Leukemia, 2020, 34, 2125-2137.	7.2	47
84	Identification of novel posttranscriptional targets of the BCR/ABL oncoprotein by ribonomics: requirement of E2F3 for BCR/ABL leukemogenesis. Blood, 2008, 111, 816-828.	1.4	44
85	Epigenetic Silencing of the Proapoptotic Gene BIM in Anaplastic Large Cell Lymphoma through an MeCP2/SIN3a Deacetylating Complex. Neoplasia, 2013, 15, 511-IN17.	5.3	44
86	Bosutinib versus imatinib for newly diagnosed chronic phase chronic myeloid leukemia: final results from the BFORE trial. Leukemia, 2022, 36, 1825-1833.	7.2	43
87	Phenotypic and functional analysis of lymphocytes infiltrating paediatric tumours, with a characterization of the tumour phenotype. Cancer Immunology, Immunotherapy, 1992, 34, 241-251.	4.2	42
88	Effects of Bosutinib Treatment on Renal Function in Patients With Philadelphia Chromosome-Positive Leukemias. Clinical Lymphoma, Myeloma and Leukemia, 2017, 17, 684-695.e6.	0.4	42
89	Factors influencing longâ€ŧerm efficacy and tolerability of bosutinib in chronic phase chronic myeloid leukaemia resistant or intolerant to imatinib. British Journal of Haematology, 2016, 172, 97-110.	2.5	41
90	Gene expression signature of non-involved lung tissue associated with survival in lung adenocarcinoma patients. Carcinogenesis, 2013, 34, 2767-2773.	2.8	40

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91	Whole Exome Sequencing reveals NOTCH1 mutations in anaplastic large cell lymphoma and points to Notch both as a key pathway and a potential therapeutic target. Haematologica, 2021, 106, 1693-1704.	3.5	40
92	Are Chronic Myeloid Leukemia Patients More at Risk for Second Malignancies? A Population-based Study. American Journal of Epidemiology, 2010, 172, 1028-1033.	3.4	39
93	Synergistic Effects of Combined Wnt/KRAS Inhibition in Colorectal Cancer Cells. PLoS ONE, 2012, 7, e51449.	2.5	39
94	BCR and BCR-ABL regulation during myeloid differentiation in healthy donors and in chronic phase/blast crisis CML patients. Leukemia, 2010, 24, 1445-1449.	7.2	37
95	Sphingosine kinase 1 overexpression is regulated by signaling through PI3K, AKT2, and mTOR in imatinib-resistant chronic myeloid leukemia cells. Experimental Hematology, 2011, 39, 653-665.e6.	0.4	37
96	Excess of NPM-ALK oncogenic signaling promotes cellular apoptosis and drug dependency. Oncogene, 2016, 35, 3854-3865.	5.9	37
97	Locking Src/Abl Tyrosine Kinase Activities Regulate Cell Differentiation and Invasion of Human Cervical Cancer Cells Expressing E6/E7 Oncoproteins of High-Risk HPV. Journal of Oncology, 2010, 2010, 1-10.	1.3	36
98	Firstâ€line treatment selection and early monitoring patterns in chronic phaseâ€chronic myeloid leukemia in routine clinical practice: SIMPLICITY. American Journal of Hematology, 2017, 92, 1214-1223.	4.1	36
99	BCR/ABL1 and BCR are under the transcriptional control of the MYC oncogene. Molecular Cancer, 2015, 14, 132.	19.2	35
100	Binding of imatinib by α1-acid glycoprotein. Blood, 2002, 100, 367-369.	1.4	34
101	NPM/ALK binds and phosphorylates the RNA/DNA-binding protein PSF in anaplastic large-cell lymphoma. Blood, 2007, 110, 2600-2609.	1.4	34
102	Structural Insights into the ATP Binding Pocket of the Anaplastic Lymphoma Kinase by Site-Directed Mutagenesis, Inhibitor Binding Analysis, and Homology Modeling. Journal of Medicinal Chemistry, 2006, 49, 5759-5768.	6.4	33
103	Imatinib-loaded polyelectrolyte microcapsules for sustained targeting of BCR-ABL ⁺ leukemia stem cells. Nanomedicine, 2010, 5, 419-431.	3.3	33
104	Chronic myeloid leukemia: Secondâ€line drugs of choice. American Journal of Hematology, 2016, 91, 67-75.	4.1	33
105	Alterations in creatine kinase, phosphate and lipid values in patients with chronic myeloid leukemia during treatment with imatinib. Haematologica, 2008, 93, 317-318.	3.5	32
106	Tyrosine kinase inhibitor interruptions, discontinuations and switching in patients with chronicâ€phase chronic myeloid leukemia in routine clinical practice: SIMPLICITY. American Journal of Hematology, 2019, 94, 46-54.	4.1	32
107	Reversal of microRNA-150 silencing disadvantages crizotinib-resistant NPM-ALK(+) cell growth. Journal of Clinical Investigation, 2015, 125, 3505-3518.	8.2	32
108	OncoScore: a novel, Internet-based tool to assess the oncogenic potential of genes. Scientific Reports, 2017, 7, 46290.	3.3	31

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109	Bcl-XL down-regulation suppresses the tumorigenic potential of NPM/ALK in vitro and in vivo. Blood, 2004, 103, 2787-2794.	1.4	30
110	Expression, purification, and inhibition of human RET tyrosine kinase. Protein Expression and Purification, 2005, 41, 177-185.	1.3	30
111	Valproic acid enhances bosutinib cytotoxicity in colon cancer cells. International Journal of Cancer, 2009, 124, 1990-1996.	5.1	29
112	FusionAnalyser: a new graphical, event-driven tool for fusion rearrangements discovery. Nucleic Acids Research, 2012, 40, e123-e123.	14.5	29
113	STAT3 and TP53 mutations associate with poor prognosis in anaplastic large cell lymphoma. Leukemia, 2021, 35, 1500-1505.	7.2	29
114	NPM/ALK mutants resistant to ASP3026 display variable sensitivity to alternative ALK inhibitors but succumb to the novel compound PF-06463922. Oncotarget, 2015, 6, 5720-5734.	1.8	29
115	A prognostic model for patients with lymphoma and COVID-19: aÂmulticentre cohort study. Blood Advances, 2022, 6, 327-338.	5.2	28
116	Systemic administration of autologous, alloactivated helper-enriched lymphocytes to patients with metastatic melanoma of the lung. Cancer Immunology, Immunotherapy, 1986, 21, 148-55.	4.2	27
117	Oncogenic Fusion Tyrosine Kinases as Molecular Targets for Anti-Cancer Therapy. Anti-Cancer Agents in Medicinal Chemistry, 2007, 7, 594-611.	1.7	27
118	Bosutinib (BOS) Versus Imatinib for Newly Diagnosed Chronic Phase (CP) Chronic Myeloid Leukemia (CML): Final 5-Year Results from the Bfore Trial. Blood, 2020, 136, 41-42.	1.4	27
119	The role of bosutinib in the treatment of chronic myeloid leukemia. Future Oncology, 2020, 16, 4395-4408.	2.4	26
120	VERSO: A comprehensive framework for the inference of robust phylogenies and the quantification of intra-host genomic diversity of viral samples. Patterns, 2021, 2, 100212.	5.9	26
121	ERG Deregulation Induces PIM1 Over-Expression and Aneuploidy in Prostate Epithelial Cells. PLoS ONE, 2011, 6, e28162.	2.5	25
122	Synergistic activity of ALK and mTOR inhibitors for the treatment of NPM-ALK positive lymphoma. Oncotarget, 2016, 7, 72886-72897.	1.8	25
123	Firstâ€line treatment of 102 chronic myeloid leukemia patients with imatinib: A longâ€ŧerm single institution analysis. American Journal of Hematology, 2014, 89, E184-7.	4.1	24
124	ALK a Novel Lymphoma-associated Tumor Antigen for Vaccination Strategies. Leukemia and Lymphoma, 2003, 44, 1675-1681.	1.3	23
125	A Compound L1196M/G1202R ALK Mutation in a Patient with ALK-Positive Lung Cancer with Acquired Resistance to Brigatinib Also Confers Primary Resistance to Lorlatinib. Journal of Thoracic Oncology, 2019, 14, e257-e259.	1.1	23
126	Autologous cellular immune response to primary and metastatic human melanomas and its regulation by DR antigens expressed on tumor cells. Cancer and Metastasis Reviews, 1985, 4, 7-26.	5.9	22

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127	ETNK1 mutations induce a mutator phenotype that can be reverted with phosphoethanolamine. Nature Communications, 2020, 11, 5938.	12.8	22
128	IL10RA Modulates Crizotinib Sensitivity in NPM1-ALK-positive Anaplastic Large Cell Lymphoma. Blood, 2020, 136, 1657-1669.	1.4	22
129	Tyrosine phosphatases regulate resistance to ALK inhibitors in ALK+ anaplastic large cell lymphoma. Blood, 2022, 139, 717-731.	1.4	22
130	Acute Promyelocytic Leukaemia Cells Resistant to Retinoic Acid Show Further Perturbation of the RARα Signal Transduction System. Leukemia and Lymphoma, 1995, 16, 289-295.	1.3	21
131	Patient-reported outcomes in the phase 3 BFORE trial of bosutinib versus imatinib for newly diagnosed chronic phase chronic myeloid leukemia. Journal of Cancer Research and Clinical Oncology, 2019, 145, 1589-1599.	2.5	21
132	<i>De novo UBE2A</i> mutations are recurrently acquired during chronic myeloid leukemia progression and interfere with myeloid differentiation pathways. Haematologica, 2019, 104, 1789-1797.	3.5	21
133	Phase two study of crizotinib in patients with anaplastic lymphoma kinase (<scp>ALK</scp>)â€positive anaplastic large cell lymphoma relapsed/refractory to chemotherapy. American Journal of Hematology, 2020, 95, E319-E321.	4.1	21
134	Effects of 1,25-Dihydroxy Vitamin D3 on All-Trans Retinoic Acid Sensitive and Resistant Acute Promyelocytic Leukemia Cells. Biochemical and Biophysical Research Communications, 1996, 224, 50-56.	2.1	20
135	CEQer: A Graphical Tool for Copy Number and Allelic Imbalance Detection from Whole-Exome Sequencing Data. PLoS ONE, 2013, 8, e74825.	2.5	20
136	Synthesis and biological evaluation of benzo[4,5]imidazo[1,2-c]pyrimidine and benzo[4,5]imidazo[1,2-a]pyrazine derivatives as anaplastic lymphoma kinase inhibitors. Bioorganic and Medicinal Chemistry, 2014, 22, 1303-1312.	3.0	20
137	In vitro and in vivo identification of ABCB1 as an efflux transporter of bosutinib. Journal of Hematology and Oncology, 2015, 8, 81.	17.0	20
138	Phase 1 Trial of Vodobatinib, a Novel Oral BCR-ABL1 Tyrosine Kinase Inhibitor (TKI): Activity in CML Chronic Phase Patients Failing TKI Therapies Including Ponatinib. Blood, 2020, 136, 51-52.	1.4	20
139	Lysis by interleukin 2-stimulated tumor-infiltrating lymphocytes of autologous and allogeneic tumor target cells. Cancer Immunology, Immunotherapy, 1989, 28, 67-73.	4.2	19
140	Sensitivity to the abl inhibitor STI571 in fresh leukaemic cells obtained from chronic myelogenous leukaemia patients in different stages of disease. British Journal of Haematology, 2001, 112, 972-974.	2.5	19
141	Morgana acts as an oncosuppressor in chronic myeloid leukemia. Blood, 2015, 125, 2245-2253.	1.4	19
142	RET kinase inhibitors: a review of recent patents (2012–2015). Expert Opinion on Therapeutic Patents, 2017, 27, 91-99.	5.0	19
143	Longâ€ŧerm patientâ€reported outcomes from an openâ€ŀabel safety and efficacy study of bosutinib in Philadelphia chromosome–positive chronic myeloid leukemia patients resistant or intolerant to prior therapy. Cancer, 2018, 124, 587-595.	4.1	19
144	Design, Synthesis, and Biological Activity of Urea Derivatives as Anaplastic Lymphoma Kinase Inhibitors. ChemMedChem, 2011, 6, 1680-1692.	3.2	18

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145	How <scp>I</scp> treat newly diagnosed chronic myeloid leukemia in 2015. American Journal of Hematology, 2015, 90, 156-161.	4.1	18
146	Telomere length shortening is associated with treatment-free remission in chronic myeloid leukemia patients. Journal of Hematology and Oncology, 2016, 9, 63.	17.0	18
147	β-catenin knockdown promotes NHERF1-mediated survival of colorectal cancer cells: implications for a double-targeted therapy. Oncogene, 2018, 37, 3301-3316.	5.9	18
148	Acute myeloid leukaemia niche regulates response to Lâ€asparaginase. British Journal of Haematology, 2019, 186, 420-430.	2.5	18
149	Identification of novel point mutations in splicing sites integrating wholeâ€exome and <scp>RNA</scp> â€seq data in myeloproliferative diseases. Molecular Genetics & Genomic Medicine, 2013, 1, 246-259.	1.2	17
150	Pregnancy outcomes in patients treated with bosutinib. International Journal of Hematologic Oncology, 2020, 9, IJH26.	1.6	17
151	Bcr-Abl mutations, resistance to imatinib, and imatinib plasma levels. Blood, 2003, 102, 1933-1935.	1.4	16
152	A Mechanistic Design Principle for Protein Tyrosine Kinase Sensors:  Application to a Validated Cancer Target. Organic Letters, 2008, 10, 301-304.	4.6	16
153	Bosutinib: a review of preclinical and clinical studies in chronic myelogenous leukemia. Expert Opinion on Pharmacotherapy, 2014, 15, 701-710.	1.8	16
154	Imatinib—A New Tyrosine Kinase Inhibitor for First-Line Treatment of Chronic Myeloid Leukemia in 2015. JAMA Oncology, 2015, 1, 143.	7.1	16
155	Long-term safety review of tyrosine kinase inhibitors in chronic myeloid leukemia - What to look for when treatment-free remission is not an option. Blood Reviews, 2022, 56, 100968.	5.7	16
156	Evidence for D276G and L364I Bcr-Abl mutations in Ph+ leukaemic cells obtained from patients resistant to Imatinib. Leukemia, 2005, 19, 132-134.	7.2	15
157	Synergistic activity of the Src/Abl inhibitor bosutinib in combination with imatinib. Leukemia, 2010, 24, 1223-1227.	7.2	15
158	Molecular cytogenetics of the acute promyelocytic leukemia-derived cell line NB4 and of four all-trans retinoic acid-resistant subclones. Genes Chromosomes and Cancer, 2002, 35, 261-270.	2.8	14
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