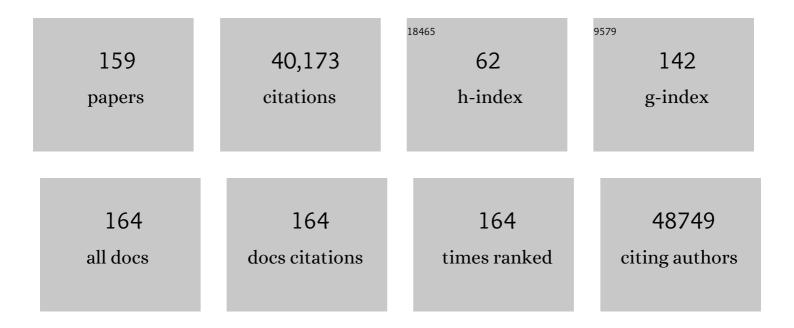
List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Signatures of mutational processes in human cancer. Nature, 2013, 500, 415-421.	13.7	8,060
2	Genomic Aberrations and Survival in Chronic Lymphocytic Leukemia. New England Journal of Medicine, 2000, 343, 1910-1916.	13.9	2,967
3	Driver mutations in histone H3.3 and chromatin remodelling genes in paediatric glioblastoma. Nature, 2012, 482, 226-231.	13.7	2,129
4	International network of cancer genome projects. Nature, 2010, 464, 993-998.	13.7	2,114
5	DNA methylation-based classification of central nervous system tumours. Nature, 2018, 555, 469-474.	13.7	1,872
6	Hotspot Mutations in H3F3A and IDH1 Define Distinct Epigenetic and Biological Subgroups of Glioblastoma. Cancer Cell, 2012, 22, 425-437.	7.7	1,551
7	The landscape of genomic alterations across childhood cancers. Nature, 2018, 555, 321-327.	13.7	1,068
8	Resistance Mechanisms for the Bruton's Tyrosine Kinase Inhibitor Ibrutinib. New England Journal of Medicine, 2014, 370, 2286-2294.	13.9	1,042
9	Molecular Classification of Ependymal Tumors across All CNS Compartments, Histopathological Grades, and Age Groups. Cancer Cell, 2015, 27, 728-743.	7.7	933
10	A Biologic Definition of Burkitt's Lymphoma from Transcriptional and Genomic Profiling. New England Journal of Medicine, 2006, 354, 2419-2430.	13.9	915
11	Matrix-based comparative genomic hybridization: Biochips to screen for genomic imbalances. Genes Chromosomes and Cancer, 1997, 20, 399-407.	1.5	879
12	The whole-genome landscape of medulloblastoma subtypes. Nature, 2017, 547, 311-317.	13.7	787
13	Dissecting the genomic complexity underlying medulloblastoma. Nature, 2012, 488, 100-105.	13.7	765
14	Genome Sequencing of Pediatric Medulloblastoma Links Catastrophic DNA Rearrangements with TP53 Mutations. Cell, 2012, 148, 59-71.	13.5	743
15	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. Cell, 2016, 164, 1060-1072.	13.5	702
16	Recurrent somatic alterations of FGFR1 and NTRK2 in pilocytic astrocytoma. Nature Genetics, 2013, 45, 927-932.	9.4	674
17	Genome Sequencing of SHH Medulloblastoma Predicts Genotype-Related Response to Smoothened Inhibition. Cancer Cell, 2014, 25, 393-405.	7.7	627
18	Detection of complete and partial chromosome gains and losses by comparative genomic in situ hybridization. Human Genetics, 1993, 90, 590-610.	1.8	544

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19	Enhancer hijacking activates GFI1 family oncogenes in medulloblastoma. Nature, 2014, 511, 428-434.	13.7	520
20	Paediatric and adult glioblastoma: multiform (epi)genomic culprits emerge. Nature Reviews Cancer, 2014, 14, 92-107.	12.8	469
21	Atypical Teratoid/Rhabdoid Tumors Are Comprised of Three Epigenetic Subgroups with Distinct Enhancer Landscapes. Cancer Cell, 2016, 29, 379-393.	7.7	438
22	Gene mutations and treatment outcome in chronic lymphocytic leukemia: results from the CLL8 trial. Blood, 2014, 123, 3247-3254.	0.6	428
23	Exosomes released by chronic lymphocytic leukemia cells induce the transition of stromal cells into cancer-associated fibroblasts. Blood, 2015, 126, 1106-1117.	0.6	399
24	Recurrent mutation of the ID3 gene in Burkitt lymphoma identified by integrated genome, exome and transcriptome sequencing. Nature Genetics, 2012, 44, 1316-1320.	9.4	389
25	BCAT1 promotes cell proliferation through amino acid catabolism in gliomas carrying wild-type IDH1. Nature Medicine, 2013, 19, 901-908.	15.2	388
26	Decoding the regulatory landscape of medulloblastoma using DNA methylation sequencing. Nature, 2014, 510, 537-541.	13.7	378
27	Longitudinal molecular trajectories of diffuse glioma in adults. Nature, 2019, 576, 112-120.	13.7	320
28	Active medulloblastoma enhancers reveal subgroup-specific cellular origins. Nature, 2016, 530, 57-62.	13.7	318
29	Quantitative analysis of comparative genomic hybridization. Cytometry, 1995, 19, 27-41.	1.8	286
30	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. Lancet Oncology, The, 2018, 19, 785-798.	5.1	268
31	A comprehensive assessment of somatic mutation detection in cancer using whole-genome sequencing. Nature Communications, 2015, 6, 10001.	5.8	266
32	Next-generation personalised medicine for high-risk paediatric cancer patients – The INFORM pilot study. European Journal of Cancer, 2016, 65, 91-101.	1.3	262
33	BCAT1 restricts αKG levels in AML stem cells leading to IDHmut-like DNA hypermethylation. Nature, 2017, 551, 384-388.	13.7	261
34	The landscape of viral associations in human cancers. Nature Genetics, 2020, 52, 320-330.	9.4	261
35	DNA methylation dynamics during B cell maturation underlie a continuum of disease phenotypes in chronic lymphocytic leukemia. Nature Genetics, 2016, 48, 253-264.	9.4	254
36	Somatic CRISPR/Cas9-mediated tumour suppressor disruption enables versatile brain tumour modelling. Nature Communications, 2015, 6, 7391.	5.8	244

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37	Tumor-derived exosomes modulate PD-L1 expression in monocytes. Science Immunology, 2017, 2, .	5.6	236
38	Oncogenic FAM131B–BRAF fusion resulting from 7q34 deletion comprises an alternative mechanism of MAPK pathway activation in pilocytic astrocytoma. Acta Neuropathologica, 2011, 121, 763-774.	3.9	211
39	Clonal evolution in chronic lymphocytic leukemia: acquisition of high-risk genomic aberrations associated with unmutated VH, resistance to therapy, and short survival. Haematologica, 2007, 92, 1242-1245.	1.7	195
40	Minimal sizes of deletions detected by comparative genomic hybridization. Genes Chromosomes and Cancer, 1998, 21, 172-175.	1.5	191
41	High-resolution genomic profiling of chronic lymphocytic leukemia reveals new recurrent genomic alterations. Blood, 2012, 120, 4783-4794.	0.6	179
42	MAPK pathway activation in pilocytic astrocytoma. Cellular and Molecular Life Sciences, 2012, 69, 1799-1811.	2.4	177
43	Evolutionary Trajectories of IDHWT Glioblastomas Reveal a Common Path of Early Tumorigenesis Instigated Years ahead of Initial Diagnosis. Cancer Cell, 2019, 35, 692-704.e12.	7.7	172
44	Therapeutic targeting of ependymoma as informed by oncogenic enhancer profiling. Nature, 2018, 553, 101-105.	13.7	170
45	PD-L1 checkpoint blockade prevents immune dysfunction and leukemia development in a mouse model of chronic lymphocytic leukemia. Blood, 2015, 126, 203-211.	0.6	158
46	Risk-adapted therapy for young children with medulloblastoma (SJYC07): therapeutic and molecular outcomes from a multicentre, phase 2 trial. Lancet Oncology, The, 2018, 19, 768-784.	5.1	151
47	Evolution of DNA Methylation Is Linked to Genetic Aberrations in Chronic Lymphocytic Leukemia. Cancer Discovery, 2014, 4, 348-361.	7.7	135
48	A biobank of patient-derived pediatric brain tumor models. Nature Medicine, 2018, 24, 1752-1761.	15.2	124
49	Targeting Self-Renewal in High-Grade Brain Tumors Leads to Loss of Brain Tumor Stem Cells and Prolonged Survival. Cell Stem Cell, 2014, 15, 185-198.	5.2	123
50	A cellâ€based model system links chromothripsis with hyperploidy. Molecular Systems Biology, 2015, 11, 828.	3.2	118
51	Chd7 is indispensable for mammalian brain development through activation of a neuronal differentiation programme. Nature Communications, 2017, 8, 14758.	5.8	118
52	Spatial heterogeneity in medulloblastoma. Nature Genetics, 2017, 49, 780-788.	9.4	112
53	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. Cancer Discovery, 2021, 11, 2764-2779.	7.7	110
54	LifeTime and improving European healthcare through cell-based interceptive medicine. Nature, 2020, 587, 377-386.	13.7	108

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55	Recurrent copy number gain of transcription factor <i>SOX2</i> and corresponding high protein expression in oral squamous cell carcinoma. Genes Chromosomes and Cancer, 2010, 49, 9-16.	1.5	106
56	Experimental evidence for the influence of molecular crowding on nuclear architecture. Journal of Cell Science, 2007, 120, 1673-1680.	1.2	104
57	Chromothripsis in cancer cells: An update. International Journal of Cancer, 2016, 138, 2322-2333.	2.3	101
58	Genomic and transcriptomic changes complement each other in the pathogenesis of sporadic Burkitt lymphoma. Nature Communications, 2019, 10, 1459.	5.8	99
59	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. Nature, 2020, 580, 396-401.	13.7	94
60	Epithelioid glioblastomas stratify into established diagnostic subsets upon integrated molecular analysis. Brain Pathology, 2018, 28, 656-662.	2.1	89
61	Inflammatory cytokines and signaling pathways are associated with survival of primary chronic lymphocytic leukemia cells in vitro: a dominant role of CCL2. Haematologica, 2011, 96, 408-416.	1.7	80
62	Molecular signatures classify astrocytic gliomas by <i>IDH1</i> mutation status. International Journal of Cancer, 2011, 128, 1095-1103.	2.3	75
63	The landscape of chromothripsis across adult cancer types. Nature Communications, 2020, 11, 2320.	5.8	75
64	Branchedâ€chain ketoacids secreted by glioblastoma cells via <scp>MCT</scp> 1 modulate macrophage phenotype. EMBO Reports, 2017, 18, 2172-2185.	2.0	74
65	Capture and Amplification by Tailing and Switching (CATS). RNA Biology, 2014, 11, 817-828.	1.5	68
66	Hypermutation of the Inactive X Chromosome Is a Frequent Event in Cancer. Cell, 2013, 155, 567-581.	13.5	67
67	Defective DNA damage repair leads to frequent catastrophic genomic events in murine and human tumors. Nature Communications, 2018, 9, 4760.	5.8	66
68	YAP1 subgroup supratentorial ependymoma requires TEAD and nuclear factor I-mediated transcriptional programmes for tumorigenesis. Nature Communications, 2019, 10, 3914.	5.8	65
69	Comprehensive Analysis of Chromatin States in Atypical Teratoid/Rhabdoid Tumor Identifies Diverging Roles for SWI/SNF and Polycomb in Gene Regulation. Cancer Cell, 2019, 35, 95-110.e8.	7.7	65
70	Medulloblastoma-associated DDX3 variant selectively alters the translational response to stress. Oncotarget, 2016, 7, 28169-28182.	0.8	62
71	Soluble CD14 is a novel monocyte-derived survival factor for chronic lymphocytic leukemia cells, which is induced by CLL cells in vitro and present at abnormally high levels in vivo. Blood, 2010, 116, 4223-4230.	0.6	60
72	Loss of SOX2 expression induces cell motility via vimentin upâ€regulation and is an unfavorable risk factor for survival ofÂhead and neck squamous cell carcinoma. Molecular Oncology, 2015, 9, 1704-1719.	2.1	60

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73	Interleukin-10 receptor signaling promotes the maintenance of a PD-1int TCF-1+ CD8+ TÂcell population that sustains anti-tumor immunity. Immunity, 2021, 54, 2825-2841.e10.	6.6	57
74	TelomereHunter – in silico estimation of telomere content and composition from cancer genomes. BMC Bioinformatics, 2019, 20, 272.	1.2	56
75	Control of chronic lymphocytic leukemia development by clonally-expanded CD8+ T-cells that undergo functional exhaustion in secondary lymphoid tissues. Leukemia, 2019, 33, 625-637.	3.3	55
76	The genomic and transcriptional landscape of primary central nervous system lymphoma. Nature Communications, 2022, 13, 2558.	5.8	52
77	Brainstem biopsy in pediatric diffuse intrinsic pontine glioma in the era of precision medicine: the INFORM study experience. European Journal of Cancer, 2019, 114, 27-35.	1.3	51
78	PI3Kδ inhibition modulates regulatory and effector T-cell differentiation and function in chronic lymphocytic leukemia. Leukemia, 2019, 33, 1427-1438.	3.3	51
79	Molecular subgrouping of primary pineal parenchymal tumors reveals distinct subtypes correlated with clinical parameters and genetic alterations. Acta Neuropathologica, 2020, 139, 243-257.	3.9	50
80	Glioblastoma epigenome profiling identifies SOX10 as a master regulator of molecular tumour subtype. Nature Communications, 2020, 11, 6434.	5.8	48
81	Kinome-wide shRNA Screen Identifies the Receptor Tyrosine Kinase AXL as a Key Regulator for Mesenchymal Glioblastoma Stem-like Cells. Stem Cell Reports, 2015, 4, 899-913.	2.3	47
82	Telomere dysfunction and chromothripsis. International Journal of Cancer, 2016, 138, 2905-2914.	2.3	42
83	Evidence for multi-copy Mega-NUMT <i>s</i> in the human genome. Nucleic Acids Research, 2021, 49, 1517-1531.	6.5	42
84	The need for a worldwide consensus for cell line authentication: Experience implementing a mandatory requirement at the International Journal of Cancer. PLoS Biology, 2017, 15, e2001438.	2.6	41
85	17p Deletion Predicts for Inferior Overall Survival after Fludarabine - Based First Line Therapy in Chronic Lymphocytic Leukemia: First Analysis of Genetics in the CLL4 Trial of the GCLLSG Blood, 2005, 106, 715-715.	0.6	41
86	Linking aberrant chromatin features in chronic lymphocytic leukemia to transcription factor networks. Molecular Systems Biology, 2019, 15, e8339.	3.2	39
87	Reprogramming of the ERRα and ERα Target Gene Landscape Triggers Tamoxifen Resistance in Breast Cancer. Cancer Research, 2015, 75, 720-731.	0.4	36
88	Rapid analysis of mouse-hamster hybrid cell lines by in situ hybridization. Genomics, 1990, 7, 127-130.	1.3	34
89	Mutational mechanisms shaping the coding and noncoding genome of germinal center derived B-cell lymphomas. Leukemia, 2021, 35, 2002-2016.	3.3	34
90	<i>NFATC1</i> activation by <scp>DNA</scp> hypomethylation in chronic lymphocytic leukemia correlates with clinical staging and can be inhibited by ibrutinib. International Journal of Cancer, 2018, 142, 322-333.	2.3	33

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91	No correlation between NF1 mutation position and risk of optic pathway glioma in 77 unrelated NF1 patients. Human Genetics, 2016, 135, 469-475.	1.8	29
92	EOMES and IL-10 regulate antitumor activity of T regulatory type 1 CD4+ T cells in chronic lymphocytic leukemia. Leukemia, 2021, 35, 2311-2324.	3.3	27
93	Low-dose Actinomycin-D treatment re-establishes the tumoursuppressive function of P53 in RELA-positive ependymoma. Oncotarget, 2016, 7, 61860-61873.	0.8	27
94	Kruppel-like factor 4 (KLF4) inactivation in chronic lymphocytic leukemia correlates with promoter DNA-methylation and can be reversed by inhibition of NOTCH signaling. Haematologica, 2016, 101, e249-e253.	1.7	26
95	Tumor necrosis factor receptor signaling is a driver of chronic lymphocytic leukemia that can be therapeutically targeted by the flavonoid wogonin. Haematologica, 2018, 103, 688-697.	1.7	26
96	Combining ibrutinib and checkpoint blockade improves CD8+ T-cell function and control of chronic lymphocytic leukemia in Em-TCL1 mice. Haematologica, 2021, 106, 968-977.	1.7	26
97	A novel cloning strategy for one-step assembly of multiplex CRISPR vectors. Scientific Reports, 2018, 8, 17499.	1.6	25
98	Altered nuclear envelope structure and proteasome function of micronuclei. Experimental Cell Research, 2018, 371, 353-363.	1.2	25
99	DECIPHER pooled shRNA library screen identifies PP2A and FGFR signaling as potential therapeutic targets for diffuse intrinsic pontine gliomas. Neuro-Oncology, 2019, 21, 867-877.	0.6	24
100	Cytogenetic characterization of head and neck squamous cell carcinoma cell lines as model systems for the functional analyses of tumorâ€associated genes. Journal of Oral Pathology and Medicine, 2010, 39, 382-389.	1.4	23
101	Oxidative stress as candidate therapeutic target to overcome microenvironmental protection of CLL. Leukemia, 2020, 34, 115-127.	3.3	23
102	Recollections of a scientific journey published in human genetics: from chromosome territories to interphase cytogenetics and comparative genome hybridization. Human Genetics, 2014, 133, 403-416.	1.8	22
103	Antagonizing inactivated tumor suppressor genes and activated oncogenes by a versatile transgenesis system: application in mantle cell lymphoma. FASEB Journal, 2006, 20, 1188-1190.	0.2	21
104	CATCH: A Prospective Precision Oncology Trial in Metastatic Breast Cancer. JCO Precision Oncology, 2021, 5, 676-686.	1.5	20
105	Biologic and Clinical Markers for Outcome after Fludarabine (F) or F Plus Cyclophosphamide (FC) - Comprehensive Analysis of the CLL4 Trial of the GCLLSG Blood, 2008, 112, 2089-2089.	0.6	18
106	CD8 <sup>+</sup> T-cells of CLL-bearing mice acquire a transcriptional program of T-cell activation and exhaustion. Leukemia and Lymphoma, 2020, 61, 351-356.	0.6	17
107	Extracellular vesicles in chronic lymphocytic leukemia. Leukemia and Lymphoma, 2013, 54, 1826-1830.	0.6	15
108	CITED4 gene silencing in colorectal cancer cells modulates adherens/tight junction gene expression and reduces cell proliferation. Journal of Cancer Research and Clinical Oncology, 2016, 142, 225-237.	1.2	15

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109	A PRDX1â€p38α heterodimer amplifies METâ€driven invasion of <i>IDH</i> â€wildtype and <i>IDH</i> â€mutant gliomas. International Journal of Cancer, 2018, 143, 1176-1187.	2.3	14
110	Mismatch Repair Deficiency Drives Durable Complete Remission by Targeting Programmed Death Receptor 1 in a Metastatic Luminal Breast Cancer Patient. Breast Care, 2019, 14, 53-59.	0.8	13
111	Rejection of adoptively transferred Eµ-TCL1 chronic lymphocytic leukemia cells in C57BL/6 substrains or knockout mouse lines. Leukemia, 2019, 33, 1514-1539.	3.3	12
112	Association of mutation signature effectuating processes with mutation hotspots in driver genes and non-coding regions. Nature Communications, 2022, 13, 178.	5.8	12
113	TBETâ€expressing Th1 CD4 <sup>+</sup> T cells accumulate in chronic lymphocytic leukaemia without affecting disease progression in Eµâ€TCL1 mice. British Journal of Haematology, 2020, 189, 133-145.	1.2	11
114	Chromothripsis in Human Breast Cancer. Cancer Research, 2020, 80, 4918-4931.	0.4	11
115	Clonal evolution in chronic lymphocytic leukemia is scant in relapsed but accelerated in refractory cases after chemo(immune) therapy. Haematologica, 2022, 107, 604-614.	1.7	11
116	Genetics of Patients with F-Refractory CLL or Early Relapse After FC or FCR: Results From the CLL8 Trial of the GCLLSG. Blood, 2010, 116, 2427-2427.	0.6	11
117	All You Need Is a Mir-acle: The Role of Nontranslated RNAs in the Suppression of B Cell Chronic Lymphocytic Leukemia. Cancer Cell, 2010, 17, 3-4.	7.7	10
118	Pilocytic astrocytoma demethylation and transcriptional landscapes link bZIP transcription factors to immune response. Neuro-Oncology, 2020, 22, 1327-1338.	0.6	10
119	In Acute Myeloid Leukemia with Complex Karyotype TP53 Alterations Are Associated with Specific Genomic Aberrations and Predict Inferior Survival Blood, 2009, 114, 2632-2632.	0.6	10
120	IGF2 knockdown in two colorectal cancer cell lines decreases survival, adhesion and modulates survival-associated genes. Tumor Biology, 2016, 37, 12485-12495.	0.8	9
121	confFuse: High-Confidence Fusion Gene Detection across Tumor Entities. Frontiers in Genetics, 2017, 8, 137.	1.1	9
122	TALEN/CRISPR-mediated engineering of a promoterless anti-viral RNAi hairpin into an endogenous miRNA locus. Nucleic Acids Research, 2017, 45, e3-e3.	6.5	8
123	Selective BTK inhibition improves bendamustine therapy response and normalizes immune effector functions in chronic lymphocytic leukemia. International Journal of Cancer, 2019, 144, 2762-2773.	2.3	8
124	Cited4 is a sexâ€biased mediator of the antidiabetic glitazone response in adipocyte progenitors. EMBO Molecular Medicine, 2018, 10, .	3.3	7
125	Gene Mutations and Treatment Outcome in Chronic Lymphocytic Leukemia: Results From the CLL8 Trial. Blood, 2012, 120, 433-433.	0.6	7
126	Cortactin expression: Association with disease progression and survival in oral squamous cell carcinoma. Head and Neck, 2018, 40, 2685-2694.	0.9	6

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127	DNA methylation of chronic lymphocytic leukemia with differential response to chemotherapy. Scientific Data, 2020, 7, 133.	2.4	6
128	Dissecting the Prognostic Significance and Functional Role of Progranulin in Chronic Lymphocytic Leukemia. Cancers, 2019, 11, 822.	1.7	5
129	A synergistic interaction between HDAC―and PARP inhibitors in childhood tumors with chromothripsis. International Journal of Cancer, 2022, 151, 590-606.	2.3	5
130	TP53 Alterations in Acute Myeloid Leukemia with Complex Karyotype Correlate with Specific Copy Number Alterations, Monosomal Karyotype, and Dismal Outcome,. Blood, 2011, 118, 3558-3558.	0.6	4
131	Correlation of Quantitative Gene Expression by RQ-PCR with Clinical and Biologic Factors in Mantle Cell Lymphoma Blood, 2004, 104, 696-696.	0.6	4
132	A versatile system to introduce clusters of genomic doubleâ€strand breaks in large cell populations. Genes Chromosomes and Cancer, 2021, 60, 303-313.	1.5	3
133	Longitudinal analyses of CLL in mice identify leukemia-related clonal changes including a Myc gain predicting poor outcome in patients. Leukemia, 2021, , .	3.3	3
134	Quantitative Gene Expression Analysis of Surrogate Markers for Genetic Risk Groups and Survival in CLL. Blood, 2008, 112, 4170-4170.	0.6	2
135	Focal structural variants revealed by whole genome sequencing disrupt the histone demethylase KDM4C in B-cell lymphomas. Haematologica, 2023, 108, 543-554.	1.7	2
136	CRISPR-mediated Loss of Function Analysis in Cerebellar Granule Cells Using <em>In Utero</em> Electroporation-based Gene Transfer. Journal of Visualized Experiments, 2018, , .	0.2	1
137	Clonal Evolution in Chronic Lymphocytic Leukemia: Acquisition of High-Risk Genomic Aberrations Associated with Unmutated VH, Resistance to Therapy, and Short Survival Blood, 2006, 108, 296-296.	0.6	1
138	MicroRNA-130a Targets ATG2B, AGO4 and DICER1, Inhibits Autophagy and Induces Cell Death in Chronic Lymphocytic Leukemia. Blood, 2011, 118, 1768-1768.	0.6	1
139	CLL Exosome-Derived Y RNA hY4 Induces TLR7/8-Mediated Inflammation and PD-L1 Expression in Monocytes. Blood, 2016, 128, 3217-3217.	0.6	1
140	Disclosure of Copy Number Alterations in AML with Normal Karyotype Using Matrix-Based Comparative Genomic Hybridization Blood, 2005, 106, 759-759.	0.6	1
141	NF1 Alterations Are Common In AML with Complex Karyotype and Correlate with Specific Genomic Imbalances. Blood, 2010, 116, 4179-4179.	0.6	1
142	GENE-06. DISTINCT MOLECULAR SUBGROUPS OF TUMORS OF THE PINEAL REGION CORRELATE WITH CLINICAL PARAMETERS AND GENETIC ALTERATIONS. Neuro-Oncology, 2019, 21, ii81-ii82.	0.6	0
143	High Resolution Screening of Genomic Aberrations in Follicular Lymphoma Using Microarray Based Comparative Genomic Hybridization (MATRIX-CGH) Blood, 2004, 104, 2271-2271.	0.6	Ο
144	Identification of Genomic Imbalances in AML with Complex Karyotype Using Matrix-Based Comparative Genomic Hybridization Blood, 2004, 104, 3382-3382.	0.6	0

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145	High ZAP-70 and Differential Expression of B-Cell Receptor Related Genes in Chronic Lymphocytic Leukemia with V3-21 Gene Usage Blood, 2004, 104, 773-773.	0.6	0
146	Protein Expression Analysis of Chromosome 12 Candidate Genes in Chronic Lymphocytic Leukemia (B-CLL) Blood, 2004, 104, 4797-4797.	0.6	0
147	A Lymphoid Progenitor Propagates AML in a Mouse Model of CALM/AF10 Positive Leukemia Blood, 2005, 106, 101-101.	0.6	0
148	Parallel Analyses Disclose Novel Genomic Imbalances in Acute Myeloid Leukemia with Complex Karyotypes Blood, 2005, 106, 2357-2357.	0.6	0
149	Identification of High-Level DNA Amplifications in AML with Complex Karyotype Using Array-CGH Blood, 2006, 108, 1914-1914.	0.6	0
150	Uncovering the Epigenetic Pathomechanism in 13q14 Blood, 2007, 110, 487-487.	0.6	0
151	Epimutation of the Tumor Suppressor Mechanism in 13q14.3 Involves Monoallelic Expression, Non-Coding RNA Genes and Deregulation of NFkB Signalling. Blood, 2008, 112, 783-783.	0.6	0
152	High-Resolution Genomic Profiling (array-CGH) of Childhood T-ALL Identifies Deletions at 6q15-16.1 as a Predictive Marker for Early Treatment Response Blood, 2008, 112, 1484-1484.	0.6	0
153	The Toll-Like Receptor-Like Molecule CD180 and Soluble CD14 Transmit Survival Signals in B-Cell Chronic Lymphocytic Leukemia Cells Presumably by Acting As Co-Receptors,. Blood, 2011, 118, 3883-3883.	0.6	0
154	Neoadjuvant epirubicin, gemcitabine, and docetaxel for primary breast cancer: Survival and prognostic factors in two consecutive neoadjuvant phase I/II trials Journal of Clinical Oncology, 2012, 30, 1096-1096.	0.8	0
155	Lenalidomide Reduces Survival of Chronic Lymphocytic Leukemia Cells in Primary Co-Cultures by Altering the Myeloid Microenvironment. Blood, 2012, 120, 3894-3894.	0.6	0
156	Microenvironmental Factors and The Role Of Tumor Necrosis Factor Receptor Type 1 (TNFR-1) In Chronic Lymphocytic Leukemia. Blood, 2013, 122, 4149-4149.	0.6	0
157	Heterogeneity and Evolution Of DNA Methylation In Chronic Lymphocytic Leukemia. Blood, 2013, 122, 1626-1626.	0.6	0
158	Modelling Single Cell B-Cell Receptor Signaling Reveals Enhanced Activity in Primary CLL Cells Compared to Non-Malignant Cells While Fundamental Network Circuit Topology Remains Stable Even with Novel Therapeutic Inhibitors. Blood, 2019, 134, 4275-4275.	0.6	0
159	Modeling the Bâ€cell receptor signaling on single cell level reveals a stable network circuit topology between nonâ€malignant B cells and chronic lymphocytic leukemia cells and between untreated cells and cells treated with kinase inhibitors. International Journal of Cancer, 2022, , .	2.3	0