## **Benedikt Brors**

List of Publications by Year in descending order

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Version: 2024-02-01

235 papers

25,185 citations

71
h-index

147

g-index

250 all docs

250 docs citations

times ranked

250

41293 citing authors

#	Article	IF	Citations
1	Whole-exome sequencing in eccrine porocarcinoma indicates promising therapeutic strategies. Cancer Gene Therapy, 2022, 29, 697-708.	4.6	10
2	A scoping review of distributed ledger technology in genomics: thematic analysis and directions for future research. Journal of the American Medical Informatics Association: JAMIA, 2022, 29, 1433-1444.	4.4	6
3	The genomic and transcriptional landscape of primary central nervous system lymphoma. Nature Communications, 2022, 13, 2558.	12.8	52
4	Rare Germline Variants Are Associated with Rapid Biochemical Recurrence After Radical Prostate Cancer Treatment: A Pan Prostate Cancer Group Study. European Urology, 2022, 82, 201-211.	1.9	2
5	Gene expression-based prediction of pazopanib efficacy in sarcoma. European Journal of Cancer, 2022, 172, 107-118.	2.8	O
6	Aggressive PDACs Show Hypomethylation of Repetitive Elements and the Execution of an Intrinsic IFN Program Linked to a Ductal Cell of Origin. Cancer Discovery, 2021, 11, 638-659.	9.4	65
7	Integrating proteomics into precision oncology. International Journal of Cancer, 2021, 148, 1438-1451.	5.1	15
8	Accurate and efficient detection of gene fusions from RNA sequencing data. Genome Research, 2021, 31, 448-460.	5 <b>.</b> 5	215
9	Characteristics and outcome of patients with acute myeloid leukaemia and t(8;16)(p11;p13): results from an International Collaborative Study*. British Journal of Haematology, 2021, 192, 832-842.	2.5	15
10	Alternative lengthening of telomeres in childhood neuroblastoma from genome to proteome. Nature Communications, 2021, 12, 1269.	12.8	46
11	Selective elimination of immunosuppressive T cells in patients with multiple myeloma. Leukemia, 2021, 35, 2602-2615.	7.2	27
12	Single-cell chromatin accessibility landscape identifies tissue repair program in human regulatory TÂcells. Immunity, 2021, 54, 702-720.e17.	14.3	78
13	CATCH: A Prospective Precision Oncology Trial in Metastatic Breast Cancer. JCO Precision Oncology, 2021, 5, 676-686.	3.0	20
14	Deconvolution of sarcoma methylomes reveals varying degrees of immune cell infiltrates with association to genomic aberrations. Journal of Translational Medicine, 2021, 19, 204.	4.4	5
15	Mutational mechanisms shaping the coding and noncoding genome of germinal center derived B-cell lymphomas. Leukemia, 2021, 35, 2002-2016.	7.2	34
16	Comprehensive Genomic and Transcriptomic Analysis for Guiding Therapeutic Decisions in Patients with Rare Cancers. Cancer Discovery, 2021, 11, 2780-2795.	9.4	125
17	RosettaSX: Reliable gene expression signature scoring of cancer models and patients. Neoplasia, 2021, 23, 1069-1077.	5.3	3
18	Sarcoma classification by DNA methylation profiling. Nature Communications, 2021, 12, 498.	12.8	237

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19	Clinical Impact of Molecular Subtyping of Pancreatic Cancer. Frontiers in Cell and Developmental Biology, 2021, 9, 743908.	3.7	29
20	Distributed Ledger Technology in genomics: a call for Europe. European Journal of Human Genetics, 2020, 28, 139-140.	2.8	19
21	Identification and characterization of a BRAF fusion oncoprotein with retained autoinhibitory domains. Oncogene, 2020, 39, 814-832.	5.9	19
22	Precursors for Nonlymphoid-Tissue Treg Cells Reside in Secondary Lymphoid Organs and Are Programmed by the Transcription Factor BATF. Immunity, 2020, 52, 295-312.e11.	14.3	140
23	Identification of BCL-XL as highly active survival factor and promising therapeutic target in colorectal cancer. Cell Death and Disease, 2020, 11, 875.	6.3	17
24	Successful BRAF/MEK inhibition in a patient with <i>BRAF</i> <sup>V600E</sup> -mutated extrapancreatic acinar cell carcinoma. Journal of Physical Education and Sports Management, 2020, 6, a005553.	1.2	13
25	The landscape of chromothripsis across adult cancer types. Nature Communications, 2020, 11, 2320.	12.8	75
26	Dissecting intratumour heterogeneity of nodal B-cell lymphomas at the transcriptional, genetic and drug-response levels. Nature Cell Biology, 2020, 22, 896-906.	10.3	93
27	Germline <i>SDHB</i> â€inactivating mutation in gastric spindle cell sarcoma. Genes Chromosomes and Cancer, 2020, 59, 601-608.	2.8	4
28	Comprehensive genomic characterization of gene therapy-induced T-cell acute lymphoblastic leukemia. Leukemia, 2020, 34, 2785-2789.	7.2	4
29	Macrophages/Microglia Represent the Major Source of Indolamine 2,3-Dioxygenase Expression in Melanoma Metastases of the Brain. Frontiers in Immunology, 2020, 11, 120.	4.8	28
30	Assessment of modelling strategies for drug response prediction in cell lines and xenografts. Scientific Reports, 2020, 10, 2849.	3.3	31
31	Genomic footprints of activated telomere maintenance mechanisms in cancer. Nature Communications, 2020, 11, 733.	12.8	87
32	Comprehensive analysis of chromothripsis in 2,658 human cancers using whole-genome sequencing. Nature Genetics, 2020, 52, 331-341.	21.4	431
33	Whole-genome fingerprint of the DNA methylome during chemically induced differentiation of the human AML cell line HL-60/S4. Biology Open, 2020, 9, .	1.2	3
34	Genome-Wide DNA Methylation Profiling in Early Stage I Lung Adenocarcinoma Reveals Predictive Aberrant Methylation in the Promoter Region of the Long Noncoding RNA PLUT: An Exploratory Study. Journal of Thoracic Oncology, 2020, 15, 1338-1350.	1.1	8
35	Targetable ERBB2 mutations identified in neurofibroma/schwannoma hybrid nerve sheath tumors. Journal of Clinical Investigation, 2020, 130, 2488-2495.	8.2	23
36	Abstract 821: Comprehensive genomic analysis of rare cancers: Results of the MASTER precision oncology trial of the German Cancer Consortium. , 2020, , .		0

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37	Abstract 820: Genomics based personalized oncology of cancer of unknown primary. , 2020, , .		O
38	Homology Modelling and Molecular Docking Studies of Selected Substituted  Lyase Receptor. Bioinformatics and Biology Insights, 2019, 13, 117793221986553.	2.0	35
39	KIT-Dependent and KIT-Independent Genomic Heterogeneity of Resistance in Gastrointestinal Stromal Tumors — TORC1/2 Inhibition as Salvage Strategy. Molecular Cancer Therapeutics, 2019, 18, 1985-1996.	4.1	22
40	TelomereHunter – in silico estimation of telomere content and composition from cancer genomes. BMC Bioinformatics, 2019, 20, 272.	2.6	56
41	Linking aberrant chromatin features in chronic lymphocytic leukemia to transcription factor networks. Molecular Systems Biology, 2019, 15, e8339.	7.2	39
42	Impact of post-surgical freezing delay on brain tumor metabolomics. Metabolomics, 2019, 15, 78.	3.0	9
43	Variant classification in precision oncology. International Journal of Cancer, 2019, 145, 2996-3010.	5.1	76
44	MDM4 Is Targeted by 1q Gain and Drives Disease in Burkitt Lymphoma. Cancer Research, 2019, 79, 3125-3138.	0.9	19
45	Response to olaparib in a <i>PALB2</i> germline mutated prostate cancer and genetic events associated with resistance. Journal of Physical Education and Sports Management, 2019, 5, a003657.	1.2	36
46	Defective homologous recombination DNA repair as therapeutic target in advanced chordoma. Nature Communications, 2019, 10, 1635.	12.8	64
47	Genomic and transcriptomic changes complement each other in the pathogenesis of sporadic Burkitt lymphoma. Nature Communications, 2019, 10, 1459.	12.8	99
48	Generation of Whole Genome Bisulfite Sequencing Libraries from Very Low DNA Input. Methods in Molecular Biology, 2019, 1956, 229-248.	0.9	1
49	Community-driven development of a modified progression-free survival ratio for precision oncology. ESMO Open, 2019, 4, e000583.	4.5	22
50	The mutational landscape underlying carfilzomib and pomalidomide resistance in relapsed /refractory multiple myeloma. Clinical Lymphoma, Myeloma and Leukemia, 2019, 19, e83-e84.	0.4	0
51	N2M2 (NOA-20) phase I/II trial of molecularly matched targeted therapies plus radiotherapy in patients with newly diagnosed non-MGMT hypermethylated glioblastoma. Neuro-Oncology, 2019, 21, 95-105.	1.2	100
52	<i>RSPO2</i> gene rearrangement: a powerful driver of $\hat{l}^2$ -catenin activation in liver tumours. Gut, 2019, 68, 1287-1296.	12.1	29
53	Abstract 919: Clinical relevance of comprehensive genomic analysis in patients with advanced-stage neuroendocrine neoplasms: Results from the MASTER trial of the German Cancer Consortium. Cancer Research, 2019, 79, 919-919.	0.9	2
54	Glioblastoma evolution pattern under surgery and radio(chemo)therapy (RCHT) to identify novel methylome based glioma subtypes Journal of Clinical Oncology, 2019, 37, 2012-2012.	1.6	3

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55	Abstract 1686: Comprehensive genomic and transcriptomic profiling of gastrointestinal stromal tumors. , 2019, , .		O
56	Abstract 468: Clinical relevance of comprehensive genomic analysis in advanced-stage cancers and rare malignancies: Results from the MASTER trial of the German Cancer Consortium., 2019,,.		0
57	Abstract 2723: Defective homologous recombination DNA repair as therapeutic target in advanced chordoma., 2019,,.		0
58	A Comprehensive Analysis of Single-Cell Chromatin Accessibility and Gene Expression Identifies Intra-Tumor Heterogeneity and Molecular Treatment Responses in Relapsed/Refractory Multiple Myeloma. Blood, 2019, 134, 575-575.	1.4	0
59	No Evidence for Hematopoietic Stem Cell Self-Renewal in-Vivo Following Inflammatory Challenge. Blood, 2019, 134, 456-456.	1.4	1
60	Dissecting Heterogeneity of Tumor Cells and Their Microenvironment in Refractory Multiple Myeloma. Blood, 2019, 134, 571-571.	1.4	0
61	Biallelic Inactivation of Multiple Tumor Suppressors Is Associated with Early Relapse after Stem Cell Transplant in Newly Diagnosed Myeloma. Blood, 2019, 134, 1783-1783.	1.4	3
62	Deconvolution of Hematopoietic Commitment Decisions By Genome-Wide Analysis of Progressive DNA Methylation Changes. Blood, 2019, 134, 1179-1179.	1.4	0
63	Abstract LB-B08: Identification and characterization of an unusual BRAF fusion oncoprotein with retained autoinhibitory domains. , $2019$ , , .		0
64	The landscape of genomic alterations across childhood cancers. Nature, 2018, 555, 321-327.	27.8	1,068
65	Methylation profiling identifies two subclasses of squamous cell carcinoma related to distinct cells of origin. Nature Communications, 2018, 9, 577.	12.8	64
66	Feasibility of real-time molecular profiling for patients with newly diagnosed glioblastoma without MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology, 2018, 20, 826-837.	1.2	32
66	MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology,		32
	MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology, 2018, 20, 826-837.  Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9,	1.2	
67	MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology, 2018, 20, 826-837.  Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9, 144.	1.2	197
67 68	MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology, 2018, 20, 826-837.  Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9, 144.  MetaboDiff: an R package for differential metabolomic analysis. Bioinformatics, 2018, 34, 3417-3418.  From somatic variants towards precision oncology: Evidence-driven reporting of treatment options	1.2 12.8 4.1	197
67 68 69	MGMT promoter hypermethylationâ€"the NCT Neuro Master Match (N2M2) pilot study. Neuro-Oncology, 2018, 20, 826-837.  Integrative genomic and transcriptomic analysis of leiomyosarcoma. Nature Communications, 2018, 9, 144.  MetaboDiff: an R package for differential metabolomic analysis. Bioinformatics, 2018, 34, 3417-3418.  From somatic variants towards precision oncology: Evidence-driven reporting of treatment options in molecular tumor boards. Genome Medicine, 2018, 10, 18.  Validating Comprehensive Next-Generation Sequencing Results for Precision Oncology: The NCT/DKTK Molecularly Aided Stratification for Tumor Eradication Research Experience. JCO Precision Oncology,	1.2 12.8 4.1 8.2	197 41 36

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73	Molecular Evolution of Early-Onset Prostate Cancer Identifies Molecular Risk Markers and Clinical Trajectories. Cancer Cell, 2018, 34, 996-1011.e8.	16.8	190
74	IGF1R upregulation confers resistance to isoform-specific inhibitors of PI3K in PIK3CA-driven ovarian cancer. Cell Death and Disease, 2018, 9, 944.	6.3	33
75	<i>NRG1</i> Fusions in <i>KRAS</i> Wild-Type Pancreatic Cancer. Cancer Discovery, 2018, 8, 1087-1095.	9.4	189
76	HiGHmed – An Open Platform Approach to Enhance Care and Research across Institutional Boundaries. Methods of Information in Medicine, 2018, 57, e66-e81.	1.2	64
77	Drug-based perturbation screen uncovers synergistic drug combinations in Burkitt lymphoma. Scientific Reports, 2018, 8, 12046.	3.3	22
78	Distinct human circulating NKp30 $<$ sup>+ $<$ /sup> FclµRll³ $<$ sup>+ $<$ /sup> CD8 $<$ sup>+ $<$ /sup> T cell population exhibiting high natural killer-like antitumor potential. Proceedings of the National Academy of Sciences of the United States of America, 2018, 115, E5980-E5989.	7.1	43
79	Towards a molecular algorithm predicting glioma treatment response and resistance: A biomarker analysis and path to real time profiling in N2M2 Journal of Clinical Oncology, 2018, 36, 12090-12090.	1.6	0
80	Recurrent Mutations in EGR2 Direct Specific Epigenetic Reconfiguration in Chronic Lymphocytic Leukemia. Blood, 2018, 132, 650-650.	1.4	0
81	Cooperative Effects of a Dnmt Inhibitor and All-Trans Retinoic Acid in an AML Cell Line Model Lacking PML-Rara. Blood, 2018, 132, 1359-1359.	1.4	0
82	PD-L1 (CD274) copy number gain, expression, and immune cell infiltration as candidate predictors for response to immune checkpoint inhibitors in soft-tissue sarcoma. Oncolmmunology, 2017, 6, e1279777.	4.6	50
83	Genetic and epigenetic profiling of a solitary Peutz–Jeghers colon polyp. Journal of Physical Education and Sports Management, 2017, 3, a001610.	1.2	10
84	Precision oncology based on omics data: The NCT Heidelberg experience. International Journal of Cancer, 2017, 141, 877-886.	5.1	133
85	Succession of transiently active tumorâ€initiating cell clones in human pancreatic cancer xenografts. EMBO Molecular Medicine, 2017, 9, 918-932.	6.9	36
86	Genetic subclone architecture of tumor clone-initiating cells in colorectal cancer. Journal of Experimental Medicine, 2017, 214, 2073-2088.	8.5	30
87	DNMT and HDAC inhibitors induce cryptic transcription start sites encoded in long terminal repeats. Nature Genetics, 2017, 49, 1052-1060.	21.4	235
88	Patient-derived xenografts of gastrointestinal cancers are susceptible to rapid and delayed B-lymphoproliferation. International Journal of Cancer, 2017, 140, 1356-1363.	5.1	26
89	Revisiting the Road Map of Medullary Thymic Epithelial Cell Differentiation. Journal of Immunology, 2017, 199, 3488-3503.	0.8	32
90	The whole-genome landscape of medulloblastoma subtypes. Nature, 2017, 547, 311-317.	27.8	787

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91	Genome-wide DNA-methylation landscape defines specialization of regulatory T cells in tissues. Nature Immunology, 2017, 18, 1160-1172.	14.5	193
92	Molecular Classification Substitutes for the Prognostic Variables Stage, Age, and MYCN Status in Neuroblastoma Risk Assessment. Neoplasia, 2017, 19, 982-990.	5.3	26
93	Targeting Fibroblast Growth Factor Receptor 1 for Treatment of Soft-Tissue Sarcoma. Clinical Cancer Research, 2017, 23, 962-973.	7.0	29
94	Mutant KIT as imatinib-sensitive target in metastatic sinonasal carcinoma. Annals of Oncology, 2017, 28, 142-148.	1.2	30
95	Combining transcription factor binding affinities with open-chromatin data for accurate gene expression prediction. Nucleic Acids Research, 2017, 45, 54-66.	14.5	112
96	Genetic Contribution to Alcohol Dependence: Investigation of a Heterogeneous German Sample of Individuals with Alcohol Dependence, Chronic Alcoholic Pancreatitis, and Alcohol-Related Cirrhosis. Genes, 2017, 8, 183.	2.4	11
97	Abstract 4807:Brafmutations initiate the development of rat gliomas induced by postnatal exposure toN-ethyl-N-nitrosourea (ENU)., 2017,,.		0
98	Abstract LB-287: Identification of patients at risk for tumor predisposition syndromes based on the evaluation of sporadic cancer exome sequencing data: experiences from the NCT/DKTK MASTER program., 2017,,.		0
99	Abstract 1023: Activation of proto-oncogenes by enhancer-hijacking in highâ€risk neuroblastoma. , 2017, ,		0
100	Alterations of microRNA and microRNA-regulated messenger RNA expression in germinal center B-cell lymphomas determined by integrative sequencing analysis. Haematologica, 2016, 101, 1380-1389.	3.5	43
101	Between-species differences in gene copy number are enriched among functions critical for adaptive evolution in Arabidopsis halleri. BMC Genomics, 2016, 17, 1034.	2.8	28
102	Braf Mutations Initiate the Development of Rat Gliomas Induced by Postnatal Exposure to N-Ethyl-N-Nitrosourea. American Journal of Pathology, 2016, 186, 2569-2576.	3.8	7
103	Next-generation personalised medicine for high-risk paediatric cancer patients – The INFORM pilot study. European Journal of Cancer, 2016, 65, 91-101.	2.8	262
104	Epigenetic dynamics of monocyte-to-macrophage differentiation. Epigenetics and Chromatin, 2016, 9, 33.	3.9	73
105	Epigenomic Profiling of Human CD4+ T Cells Supports a Linear Differentiation Model and Highlights Molecular Regulators of Memory Development. Immunity, 2016, 45, 1148-1161.	14.3	174
106	Recurrent MET fusion genes represent a drug target in pediatric glioblastoma. Nature Medicine, 2016, 22, 1314-1320.	30.7	183
107	The International Human Epigenome Consortium: A Blueprint for Scientific Collaboration and Discovery. Cell, 2016, 167, 1145-1149.	28.9	404
108	MYC/MIZ1-dependent gene repression inversely coordinates the circadian clock with cell cycle and proliferation. Nature Communications, 2016, 7, 11807.	12.8	103

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109	Integration of genomics and histology revises diagnosis and enables effective therapy of refractory cancer of unknown primary with $\langle i \rangle PDL1 \langle i \rangle$ amplification. Journal of Physical Education and Sports Management, 2016, 2, a001180.	1.2	57
110	Genome-wide association study of pathological gambling. European Psychiatry, 2016, 36, 38-46.	0.2	82
111	DNA methylation dynamics during B cell maturation underlie a continuum of disease phenotypes in chronic lymphocytic leukemia. Nature Genetics, 2016, 48, 253-264.	21.4	254
112	Evolutionary conserved gene co-expression drives generation ofÂself-antigen diversity in medullary thymic epithelial cells. Journal of Autoimmunity, 2016, 67, 65-75.	6.5	19
113	Cooperation of BRAFF595L and mutant HRAS in histiocytic sarcoma provides new insights into oncogenic BRAF signaling. Leukemia, 2016, 30, 937-946.	7.2	52
114	Umbrella protocol for phase I/IIa trials of molecularly matched targeted therapies plus radiotherapy in patients with newly diagnosed glioblastoma without MGMT promoter methylation Neuro Master Match ( $N\hat{A}^2M\hat{A}^2$ ) Journal of Clinical Oncology, 2016, 34, TPS2084-TPS2084.	1.6	4
115	Abstract LB-287: Combining immunomics and genomics for immunotherapy of refractory and rare cancers. , 2016, , .		0
116	Abstract 910: Genetic subclone heterogeneity of the human colon cancer initiating cell compartment. , 2016, , .		0
117	Hematopoietic Stem Cells Fail to Regenerate In Vivo Following Inflammatory Stress. Blood, 2016, 128, 1472-1472.	1.4	0
118	Epigenetic Drug Treatment Globally Induces Cryptic Transcription Start Sites Encoded in Long Terminal Repeats. Blood, 2016, 128, 3931-3931.	1.4	0
119	Iroquois homeobox 2 suppresses cellular motility and chemokine expression in breast cancer cells. BMC Cancer, 2015, 15, 896.	2.6	18
120	In Silico Gene Regulatory Network of the Maurer's Cleft Pathway in Plasmodium falciparum. Evolutionary Bioinformatics, 2015, 11, EBO.S25585.	1.2	2
121	Thymic B Cells Are Licensed to Present Self Antigens for Central T Cell Tolerance Induction. Immunity, 2015, 42, 1048-1061.	14.3	201
122	Molecular profiling of long-term survivors identifies a subgroup of glioblastoma characterized by chromosome 19/20 co-gain. Acta Neuropathologica, 2015, 130, 419-434.	7.7	74
123	A comprehensive assessment of somatic mutation detection in cancer using whole-genome sequencing. Nature Communications, 2015, 6, 10001.	12.8	266
124	Comparison of RNA-seq and microarray-based models for clinical endpoint prediction. Genome Biology, 2015, 16, 133.	8.8	325
125	Recurrent Mutations within the Amino-Terminal Region of $\hat{l}^2$ -Catenin Are Probable Key Molecular Driver Events in Sinonasal Hemangiopericytoma. American Journal of Pathology, 2015, 185, 563-571.	3.8	49
126	Suppression of Early Hematogenous Dissemination of Human Breast Cancer Cells to Bone Marrow by Retinoic Acid–Induced 2. Cancer Discovery, 2015, 5, 506-519.	9.4	45

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127	Recurrent CDKN1B (p27) mutations in hairy cell leukemia. Blood, 2015, 126, 1005-1008.	1.4	88
128	XRCC5 as a Risk Gene for Alcohol Dependence: Evidence from a Genome-Wide Gene-Set-Based Analysis and Follow-up Studies in Drosophila and Humans. Neuropsychopharmacology, 2015, 40, 361-371.	5.4	12
129	BAZ2A (TIP5) is involved in epigenetic alterations in prostate cancer and its overexpression predicts disease recurrence. Nature Genetics, 2015, 47, 22-30.	21.4	141
130	Revised Risk Estimation and Treatment Stratification of Low- and Intermediate-Risk Neuroblastoma Patients by Integrating Clinical and Molecular Prognostic Markers. Clinical Cancer Research, 2015, 21, 1904-1915.	7.0	80
131	Progressive Epigenetic Programming during B Cell Maturation Is Reflected in a Continuum of Epigenetic Disease Phenotypes in Chronic Lymphocytic Leukemia. Blood, 2015, 126, 2436-2436.	1.4	1
132	<i>MYCN</i> amplification confers enhanced folate dependence and methotrexate sensitivity in neuroblastoma. Oncotarget, 2015, 6, 15510-15523.	1.8	13
133	FGFR2 is overexpressed in myxoid liposarcoma and inhibition of FGFR signaling impairs tumor growth <i>in vitro</i> . Oncotarget, 2015, 6, 20215-20230.	1.8	23
134	Abstract 1417: Clonal succession in pancreatic cancer progression is not driven by genetic instability. , $2015, \dots$		0
135	Cooperative Activity of BRAF F595L and Mutant HRAS in Histiocytic Sarcoma Provides New Insights into Oncogenic BRAF Signaling. Blood, 2015, 126, 1631-1631.	1.4	2
136	BRAF inhibitor–associated ERK activation drives development of chronic lymphocytic leukemia. Journal of Clinical Investigation, 2014, 124, 5074-5084.	8.2	56
137	Initiation of an Inflammatory Response in Resident Intestinal Lamina Propria Cells -Use of a Human Organ Culture Model. PLoS ONE, 2014, 9, e97780.	2.5	9
138	Integrated Pathway-Based Approach Identifies Association between Genomic Regions at CTCF and CACNB2 and Schizophrenia. PLoS Genetics, 2014, 10, e1004345.	3.5	44
139	Genetic variants in apoptosisâ€related genes associated with colorectal hyperplasia. Genes Chromosomes and Cancer, 2014, 53, 769-778.	2.8	2
140	Chromosome 17/17q gain and unaltered profiles in high resolution arrayâ€CGH are prognostically informative in neuroblastoma. Genes Chromosomes and Cancer, 2014, 53, 639-649.	2.8	34
141	Genome Sequencing of SHH Medulloblastoma Predicts Genotype-Related Response to Smoothened Inhibition. Cancer Cell, 2014, 25, 393-405.	16.8	627
142	Cell competition is a tumour suppressor mechanism in the thymus. Nature, 2014, 509, 465-470.	27.8	209
143	Decoding the regulatory landscape of medulloblastoma using DNA methylation sequencing. Nature, 2014, 510, 537-541.	27.8	378
144	Identification of DNA methylation changes at <i>cis</i> -regulatory elements during early steps of HSC differentiation using tagmentation-based whole genome bisulfite sequencing. Cell Cycle, 2014, 13, 3476-3487.	2.6	39

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145	TMPRSS2-ERG Fusions Are Strongly Linked to Young Patient Age in Low-grade Prostate Cancer. European Urology, 2014, 66, 978-981.	1.9	54
146	Whole-genome bisulfite sequencing of HSCs and their immediate progeny identifies novel regulatory elements involved in self-renewal and early hematopoietic commitment. Experimental Hematology, 2014, 42, S20.	0.4	0
147	Intratumor DNA Methylation Heterogeneity Reflects Clonal Evolution in Aggressive Prostate Cancer. Cell Reports, 2014, 8, 798-806.	6.4	219
148	Identification of Regulatory Networks in HSCs and Their Immediate Progeny via Integrated Proteome, Transcriptome, and DNA Methylome Analysis. Cell Stem Cell, 2014, 15, 507-522.	11.1	439
149	<i>circlize</i> iiplements and enhances circular visualization in R. Bioinformatics, 2014, 30, 2811-2812.	4.1	2,736
150	Investigation of manic and euthymic episodes identifies state- and trait-specific gene expression and STAB1 as a new candidate gene for bipolar disorder. Translational Psychiatry, 2014, 4, e426-e426.	4.8	30
151	Enhancer hijacking activates GFI1 family oncogenes in medulloblastoma. Nature, 2014, 511, 428-434.	27.8	520
152	Solitary Fibrous Tumors/Hemangiopericytomas with Different Variants of the NAB2-STAT6 Gene Fusion Are Characterized by Specific Histomorphology and Distinct Clinicopathological Features. American Journal of Pathology, 2014, 184, 1209-1218.	3.8	198
153	Recurrent somatic alterations of FGFR1 and NTRK2 in pilocytic astrocytoma. Nature Genetics, 2013, 45, 927-932.	21.4	674
154	Hypermutation of the Inactive X Chromosome Is a Frequent Event in Cancer. Cell, 2013, 155, 567-581.	28.9	67
155	Autotaxin is expressed in FLT3-ITD positive acute myeloid leukemia and hematopoietic stem cells and promotes cell migration and proliferation. Experimental Hematology, 2013, 41, 444-461.e4.	0.4	25
156	Integrative Genomic Analyses Reveal an Androgen-Driven Somatic Alteration Landscape in Early-Onset Prostate Cancer. Cancer Cell, 2013, 23, 159-170.	16.8	292
157	Hox-C9 activates the intrinsic pathway of apoptosis and is associated with spontaneous regression in neuroblastoma. Cell Death and Disease, 2013, 4, e586-e586.	6.3	184
158	Mining Quasi-Bicliques from HIV-1-Human Protein Interaction Network: A Multiobjective Biclustering Approach. IEEE/ACM Transactions on Computational Biology and Bioinformatics, 2013, 10, 423-435.	3.0	29
159	Overlapping gene coexpression patterns in human medullary thymic epithelial cells generate self-antigen diversity. Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, E3497-505.	7.1	70
160	Integrative DNA methylation and gene expression analysis in high-grade soft tissue sarcomas. Genome Biology, 2013, 14, r137.	9.6	78
161	Decision-Tree Based Model Analysis for Efficient Identification of Parameter Relations Leading to Different Signaling States. PLoS ONE, 2013, 8, e82593.	2.5	13
162	Coverage Bias and Sensitivity of Variant Calling for Four Whole-genome Sequencing Technologies. PLoS ONE, 2013, 8, e66621.	2.5	74

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163	Automated Universal BRAF State Detection within the Activation Segment in Skin Metastases by Pyrosequencing-Based Assay U-BRAFV600. PLoS ONE, 2013, 8, e59221.	2.5	13
164	Whole-genome analysis of gene expression associates the ubiquitin-proteasome system with the cardiomyopathy phenotype in disease-sensitized congenic mouse strains. Cardiovascular Research, 2012, 94, 87-95.	3.8	17
165	Multi-Parametric Analysis and Modeling of Relationships between Mitochondrial Morphology and Apoptosis. PLoS ONE, 2012, 7, e28694.	2.5	44
166	Genome Sequencing of Pediatric Medulloblastoma Links Catastrophic DNA Rearrangements with TP53 Mutations. Cell, 2012, 148, 59-71.	28.9	743
167	Recurrent mutation of the ID3 gene in Burkitt lymphoma identified by integrated genome, exome and transcriptome sequencing. Nature Genetics, 2012, 44, 1316-1320.	21.4	389
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