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List of Publications by Year in descending order

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38
papers

6,477
citations

172457

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315739

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docs citations

39
times ranked

11638
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular subtypes of diffuse large B cell lymphoma are associated with distinct pathogenic mechanisms and outcomes. <i>Nature Medicine</i> , 2018, 24, 679-690.	30.7	1,224
2	Genomic sequencing of meningiomas identifies oncogenic SMO and AKT1 mutations. <i>Nature Genetics</i> , 2013, 45, 285-289.	21.4	532
3	High-Throughput Detection of Actionable Genomic Alterations in Clinical Tumor Samples by Targeted, Massively Parallel Sequencing. <i>Cancer Discovery</i> , 2012, 2, 82-93.	9.4	484
4	Integration of gene mutations in risk prognostication for patients receiving first-line immunochemotherapy for follicular lymphoma: a retrospective analysis of a prospective clinical trial and validation in a population-based registry. <i>Lancet Oncology</i> , The, 2015, 16, 1111-1122.	10.7	483
5	Targetable genetic features of primary testicular and primary central nervous system lymphomas. <i>Blood</i> , 2016, 127, 869-881.	1.4	429
6	Validation of OncoPanel: A Targeted Next-Generation Sequencing Assay for the Detection of Somatic Variants in Cancer. <i>Archives of Pathology and Laboratory Medicine</i> , 2017, 141, 751-758.	2.5	350
7	Institutional implementation of clinical tumor profiling on an unselected cancer population. <i>JCI Insight</i> , 2016, 1, e87062.	5.0	340
8	Genomic Heterogeneity as a Barrier to Precision Medicine in Gastroesophageal Adenocarcinoma. <i>Cancer Discovery</i> , 2018, 8, 37-48.	9.4	248
9	MYB-QKI rearrangements in angiocentric glioma drive tumorigenicity through a tripartite mechanism. <i>Nature Genetics</i> , 2016, 48, 273-282.	21.4	214
10	Genomic analysis of diffuse pediatric low-grade gliomas identifies recurrent oncogenic truncating rearrangements in the transcription factor <i>MYBL1</i> . <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 8188-8193.	7.1	188
11	Novel Peptide Inhibitors of Angiotensin-converting Enzyme 2. <i>Journal of Biological Chemistry</i> , 2003, 278, 15532-15540.	3.4	185
12	<i>MAP2K1</i> and <i>MAP3K1</i> mutations in langerhans cell histiocytosis. <i>Genes Chromosomes and Cancer</i> , 2015, 54, 361-368.	2.8	167
13	Somatic activating ARAF mutations in Langerhans cell histiocytosis. <i>Blood</i> , 2014, 123, 3152-3155.	1.4	161
14	BreakMer: detection of structural variation in targeted massively parallel sequencing data using kmers. <i>Nucleic Acids Research</i> , 2015, 43, e19-e19.	14.5	161
15	Association of Alterations in Main Driver Genes With Outcomes of Patients With Resected Pancreatic Ductal Adenocarcinoma. <i>JAMA Oncology</i> , 2018, 4, e173420.	7.1	155
16	The impact of tumor profiling approaches and genomic data strategies for cancer precision medicine. <i>Genome Medicine</i> , 2016, 8, 79.	8.2	151
17	Germline cancer susceptibility gene variants, somatic second hits, and survival outcomes in patients with resected pancreatic cancer. <i>Genetics in Medicine</i> , 2019, 21, 213-223.	2.4	151
18	Genomic analyses of flow-sorted Hodgkin Reed-Sternberg cells reveal complementary mechanisms of immune evasion. <i>Blood Advances</i> , 2019, 3, 4065-4080.	5.2	99

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19	Landscape of Genomic Alterations in Pituitary Adenomas. <i>Clinical Cancer Research</i> , 2017, 23, 1841-1851.	7.0	94
20	Comparison of Prevalence and Types of Mutations in Lung Cancers Among Black and White Populations. <i>JAMA Oncology</i> , 2017, 3, 801.	7.1	78
21	Prospective Enterprise-Level Molecular Genotyping of a Cohort of Cancer Patients. <i>Journal of Molecular Diagnostics</i> , 2014, 16, 660-672.	2.8	70
22	Diffuse large B-cell lymphoma patient-derived xenograft models capture the molecular and biological heterogeneity of the disease. <i>Blood</i> , 2016, 127, 2203-2213.	1.4	68
23	Clinical targeted exome-based sequencing in combination with genome-wide copy number profiling: precision medicine analysis of 203 pediatric brain tumors. <i>Neuro-Oncology</i> , 2017, 19, now294.	1.2	54
24	Recurrent genetic HLA loss in AML relapsed after matched unrelated allogeneic hematopoietic cell transplantation. <i>Blood Advances</i> , 2019, 3, 2199-2204.	5.2	52
25	Next-generation sequencing of cytologic preparations: An analysis of quality metrics. <i>Cancer Cytopathology</i> , 2017, 125, 786-794.	2.4	51
26	OncoTree: A Cancer Classification System for Precision Oncology. <i>JCO Clinical Cancer Informatics</i> , 2021, 5, 221-230.	2.1	51
27	Hedgehog pathway mutations drive oncogenic transformation in high-risk T-cell acute lymphoblastic leukemia. <i>Leukemia</i> , 2018, 32, 2126-2137.	7.2	48
28	Genomic aberrations in cervical adenocarcinomas in Hong Kong Chinese women. <i>International Journal of Cancer</i> , 2015, 137, 776-783.	5.1	39
29	Correlation Between Surrogate End Points and Overall Survival in a Multi-institutional Clinicogenomic Cohort of Patients With Non-Small Cell Lung or Colorectal Cancer. <i>JAMA Network Open</i> , 2021, 4, e2117547.	5.9	20
30	Molecular Characterization and Therapeutic Targeting of Colorectal Cancers Harboring Receptor Tyrosine Kinase Fusions. <i>Clinical Cancer Research</i> , 2021, 27, 1695-1705.	7.0	19
31	Somatic mutations in <i>CDH1</i> and <i>CTNNB1</i> in primary carcinomas at 13 anatomic sites. <i>Oncotarget</i> , 2017, 8, 85680-85691.	1.8	16
32	Clinical and mutational spectrum of highly differentiated, paired box 3:forkhead box protein o1 fusion-negative rhabdomyosarcoma: A report from the Children's Oncology Group. <i>Cancer</i> , 2018, 124, 1973-1981.	4.1	14
33	Insulin-Like Growth Factor-1 Receptor Expression and Disease Recurrence and Survival in Patients with Resected Pancreatic Ductal Adenocarcinoma. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2020, 29, 1586-1595.	2.5	8
34	ViroPanel. <i>Journal of Molecular Diagnostics</i> , 2020, 22, 476-487.	2.8	6
35	Diffuse Large B-Cell Lymphoma Patient-Derived Xenograft Models Capture Molecular and Biologic Heterogeneity and Inform Therapy. <i>Blood</i> , 2015, 126, 817-817.	1.4	5
36	Dissection of PIK3CA Aberration for Cervical Adenocarcinoma Outcomes. <i>Cancers</i> , 2021, 13, 3218.	3.7	2

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37	Comprehensive Genomic Analysis of Flow-Sorted Hodgkin Reed Sternberg Cells Reveals Additional Genetic Bases of Immune Evasion. <i>Blood</i> , 2018, 132, 1559-1559.	1.4	2
38	Recurrent Genetic HLA Loss in Acute Myeloid Leukemia Relapsed after Matched Unrelated Allogeneic Hematopoietic Cell Transplant. <i>Blood</i> , 2018, 132, 817-817.	1.4	0