

Matthew D Ducar

List of Publications by Year in descending order

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38
papers

6,477
citations

172457

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#	ARTICLE	IF	CITATIONS
1	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
2	OncoTree: A Cancer Classification System for Precision Oncology. <i>JCO Clinical Cancer Informatics</i> , 2021, 5, 221-230.	2.1	51
3	Dissection of PIK3CA Aberration for Cervical Adenocarcinoma Outcomes. <i>Cancers</i> , 2021, 13, 3218.	3.7	2
4	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
5	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
6	ViroPanel. <i>Journal of Molecular Diagnostics</i> , 2020, 22, 476-487.	2.8	6
7	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
8	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
9	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
10	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
11	Hedgehog pathway mutations drive oncogenic transformation in high-risk T-cell acute lymphoblastic leukemia. <i>Leukemia</i> , 2018, 32, 2126-2137.	7.2	48
12	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
13	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
14	Genomic Heterogeneity as a Barrier to Precision Medicine in Gastroesophageal Adenocarcinoma. <i>Cancer Discovery</i> , 2018, 8, 37-48.	9.4	248
15	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
16	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
17	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
18	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

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19	Validation of OncoPanel: A Targeted Next-Generation Sequencing Assay for the Detection of Somatic Variants in Cancer. Archives of Pathology and Laboratory Medicine, 2017, 141, 751-758.	2.5	350
20	Next-generation sequencing of cytologic preparations: An analysis of quality metrics. Cancer Cytopathology, 2017, 125, 786-794.	2.4	51
21	Landscape of Genomic Alterations in Pituitary Adenomas. Clinical Cancer Research, 2017, 23, 1841-1851.	7.0	94
22	Somatic mutations in <i>CDH1</i> and <i>CTNNB1</i> in primary carcinomas at 13 anatomic sites. Oncotarget, 2017, 8, 85680-85691.	1.8	16
23	Institutional implementation of clinical tumor profiling on an unselected cancer population. JCI Insight, 2016, 1, e87062.	5.0	340
24	Targetable genetic features of primary testicular and primary central nervous system lymphomas. Blood, 2016, 127, 869-881.	1.4	429
25	The impact of tumor profiling approaches and genomic data strategies for cancer precision medicine. Genome Medicine, 2016, 8, 79.	8.2	151
26	Diffuse large B-cell lymphoma patient-derived xenograft models capture the molecular and biological heterogeneity of the disease. Blood, 2016, 127, 2203-2213.	1.4	68
27	MYB-QKI rearrangements in angiocentric glioma drive tumorigenicity through a tripartite mechanism. Nature Genetics, 2016, 48, 273-282.	21.4	214
28	<i>MAP2K1</i> and <i>MAP3K1</i> mutations in langerhans cell histiocytosis. Genes Chromosomes and Cancer, 2015, 54, 361-368.	2.8	167
29	Genomic aberrations in cervical adenocarcinomas in Hong Kong Chinese women. International Journal of Cancer, 2015, 137, 776-783.	5.1	39
30	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. Lancet Oncology, The, 2015, 16, 1111-1122.	10.7	483
31	BreaKmer: detection of structural variation in targeted massively parallel sequencing data using kmers. Nucleic Acids Research, 2015, 43, e19-e19.	14.5	161
32	Diffuse Large B-Cell Lymphoma Patient-Derived Xenograft Models Capture Molecular and Biologic Heterogeneity and Inform Therapy. Blood, 2015, 126, 817-817.	1.4	5
33	Prospective Enterprise-Level Molecular Genotyping of a Cohort of Cancer Patients. Journal of Molecular Diagnostics, 2014, 16, 660-672.	2.8	70
34	Somatic activating ARAF mutations in Langerhans cell histiocytosis. Blood, 2014, 123, 3152-3155.	1.4	161
35	Genomic sequencing of meningiomas identifies oncogenic SMO and AKT1 mutations. Nature Genetics, 2013, 45, 285-289.	21.4	532
36	Genomic analysis of diffuse pediatric low-grade gliomas identifies recurrent oncogenic truncating rearrangements in the transcription factor <i>MYBL1</i> . Proceedings of the National Academy of Sciences of the United States of America, 2013, 110, 8188-8193.	7.1	188

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37	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
38	Novel Peptide Inhibitors of Angiotensin-converting Enzyme 2. <i>Journal of Biological Chemistry</i> , 2003, 278, 15532-15540.	3.4	185