

Eric Karlins

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

1,757
citations

471509

17
h-index

610901

24
g-index

30
all docs

30
docs citations

30
times ranked

3961
citing authors

#	ARTICLE	IF	CITATIONS
1	Age-related DNA methylation in paired normal and tumour breast tissue in Chinese breast cancer patients. <i>Epigenetics</i> , 2021, 16, 677-691.	2.7	9
2	Low-frequency variation near common germline susceptibility loci are associated with risk of Ewing sarcoma. <i>PLoS ONE</i> , 2020, 15, e0237792.	2.5	6
3	Subsequent Neoplasm Risk Associated With Rare Variants in DNA Damage Response and Clinical Radiation Sensitivity Syndrome Genes in the Childhood Cancer Survivor Study. <i>JCO Precision Oncology</i> , 2020, 4, 926-936.	3.0	9
4	Frequency of Pathogenic Germline Variants in Cancer-Susceptibility Genes in Patients With Osteosarcoma. <i>JAMA Oncology</i> , 2020, 6, 724.	7.1	139
5	Title is missing!. , 2020, 15, e0237792.		0
6	Title is missing!. , 2020, 15, e0237792.		0
7	Title is missing!. , 2020, 15, e0237792.		0
8	Title is missing!. , 2020, 15, e0237792.		0
9	Identifying, understanding, and correcting technical artifacts on the sex chromosomes in next-generation sequencing data. <i>GigaScience</i> , 2019, 8, .	6.4	65
10	Genome-Wide Association Study in Irradiated Childhood Cancer Survivors Identifies HTR2A for Subsequent Basal Cell Carcinoma. <i>Journal of Investigative Dermatology</i> , 2019, 139, 2042-2045.e8.	0.7	18
11	Blood DNA methylation and breast cancer risk: a meta-analysis of four prospective cohort studies. <i>Breast Cancer Research</i> , 2019, 21, 62.	5.0	34
12	Successful use of whole genome amplified DNA from multiple source types for high-density Illumina SNP microarrays. <i>BMC Genomics</i> , 2018, 19, 182.	2.8	16
13	Genome-wide association study identifies the <i>GLDC</i> / <i>IL33</i> locus associated with survival of osteosarcoma patients. <i>International Journal of Cancer</i> , 2018, 142, 1594-1601.	5.1	31
14	Genome-wide association study identifies multiple new loci associated with Ewing sarcoma susceptibility. <i>Nature Communications</i> , 2018, 9, 3184.	12.8	50
15	Genome-Wide Association Study to Identify Susceptibility Loci That Modify Radiation-Related Risk for Breast Cancer After Childhood Cancer. <i>Journal of the National Cancer Institute</i> , 2017, 109, .	6.3	66
16	Evaluating the Causal Link Between Malaria Infection and Endemic Burkitt Lymphoma in Northern Uganda: A Mendelian Randomization Study. <i>EBioMedicine</i> , 2017, 25, 58-65.	6.1	37
17	Higher-than-expected population prevalence of potentially pathogenic germline <i>TP53</i> variants in individuals unselected for cancer history. <i>Human Mutation</i> , 2017, 38, 1723-1730.	2.5	40
18	Whole exome sequencing in 75 high-risk families with validation and replication in independent case-control studies identifies <i>TANGO2</i> , <i>OR5H14</i> , and <i>CHAD</i> as new prostate cancer susceptibility genes. <i>Oncotarget</i> , 2017, 8, 1495-1507.	1.8	11

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19	Biallelic BRCA2 Mutations Shape the Somatic Mutational Landscape of Aggressive Prostate Tumors. <i>American Journal of Human Genetics</i> , 2016, 98, 818-829.	6.2	34
20	Female chromosome X mosaicism is age-related and preferentially affects the inactivated X chromosome. <i>Nature Communications</i> , 2016, 7, 11843.	12.8	86
21	Genome-wide association study of meningioma as a subsequent neoplasm: A report from the Childhood Cancer Survivor Study (CCSS) and St. Jude Lifetime Cohort (SJLIFE).. <i>Journal of Clinical Oncology</i> , 2016, 34, 10510-10510.	1.6	0
22	Homologous Mutation to Human BRAF V600E Is Common in Naturally Occurring Canine Bladder Cancer—Evidence for a Relevant Model System and Urine-Based Diagnostic Test. <i>Molecular Cancer Research</i> , 2015, 13, 993-1002.	3.4	117
23	Comparison against 186 canid whole-genome sequences reveals survival strategies of an ancient clonally transmissible canine tumor. <i>Genome Research</i> , 2015, 25, 1646-1655.	5.5	63
24	A Copy Number Variant at the KITLG Locus Likely Confers Risk for Canine Squamous Cell Carcinoma of the Digit. <i>PLoS Genetics</i> , 2013, 9, e1003409.	3.5	60
25	Fine scale mapping of the breast cancer 16q12 locus. <i>Human Molecular Genetics</i> , 2010, 19, 2507-2515.	2.9	68
26	FGFR2 variants and breast cancer risk: fine-scale mapping using African American studies and analysis of chromatin conformation. <i>Human Molecular Genetics</i> , 2009, 18, 1692-1703.	2.9	110
27	No evidence of BRCA2 mutations in chromosome 13q-linked Utah high-risk prostate cancer pedigrees. <i>BMC Research Notes</i> , 2009, 2, 94.	1.4	2
28	A Single <i>IGF1</i> Allele Is a Major Determinant of Small Size in Dogs. <i>Science</i> , 2007, 316, 112-115.	12.6	587
29	Case-Control Study of the Parkin Gene in Early-Onset Parkinson Disease. <i>Archives of Neurology</i> , 2006, 63, 548.	4.5	95