

Leon Raskin

List of Publications by Year in descending order

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Version: 2024-02-01

51
papers

2,372
citations

257450

24
h-index

233421

45
g-index

54
all docs

54
docs citations

54
times ranked

5625
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic architectures of proximal and distal colorectal cancer are partly distinct. <i>Gut</i> , 2021, 70, 1325-1334.	12.1	44
2	Treatment patterns of malignant melanoma in the United States from 2011 to 2016: a retrospective cohort study. <i>Current Medical Research and Opinion</i> , 2020, 36, 63-72.	1.9	4
3	High mobility group A protein-2 as a tumor cancer diagnostic and prognostic marker: a systematic review and meta-analysis. <i>European Journal of Cancer Prevention</i> , 2020, 29, 565-581.	1.3	2
4	Variations in hospitalization and emergency department/observation stays using the oncology care model methodology in Medicare data. <i>Current Medical Research and Opinion</i> , 2020, 36, 1519-1527.	1.9	0
5	Observational study of talimogene laherparepvec use in the anti-PD-1 era for melanoma in the US (COSMUS-2). <i>Melanoma Management</i> , 2020, 7, MMT41.	0.5	3
6	Novel Common Genetic Susceptibility Loci for Colorectal Cancer. <i>Journal of the National Cancer Institute</i> , 2019, 111, 146-157.	6.3	129
7	Observational study of talimogene laherparepvec use for melanoma in clinical practice in the United States (COSMUS-1). <i>Melanoma Management</i> , 2019, 6, MMT19.	0.5	21
8	Shared heritability and functional enrichment across six solid cancers. <i>Nature Communications</i> , 2019, 10, 431.	12.8	88
9	Protein kinase C- δ is upregulated by IMP1 in melanoma and is linked to poor survival. <i>Melanoma Research</i> , 2019, 29, 539-543.	1.2	9
10	Treatment patterns of melanoma by <i>BRAF</i> mutation status in the USA from 2011 to 2017: a retrospective cohort study. <i>Melanoma Management</i> , 2019, 6, MMT31.	0.5	3
11	Discovery of common and rare genetic risk variants for colorectal cancer. <i>Nature Genetics</i> , 2019, 51, 76-87.	21.4	377
12	<i>CDKN2A</i> Germline Rare Coding Variants and Risk of Pancreatic Cancer in Minority Populations. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2018, 27, 1364-1370.	2.5	23
13	Activation of cancer-associated fibroblasts is required for tumor neovascularization in a murine model of melanoma. <i>Matrix Biology</i> , 2018, 74, 52-61.	3.6	52
14	A human <i>MUTYH</i> variant linking colonic polyposis to redox degradation of the [4Fe4S] ₂ ⁺ cluster. <i>Nature Chemistry</i> , 2018, 10, 873-880.	13.6	20
15	Prevalence and Predictors of Renal Impairment Among Patients with Multiple Myeloma (MM): An Analysis of Oncology Clinic Electronic Health Records Linked to Commercial Claims in the United States. <i>Blood</i> , 2018, 132, 5657-5657.	1.4	0
16	Targeted sequencing of established and candidate colorectal cancer genes in the Colon Cancer Family Registry Cohort. <i>Oncotarget</i> , 2017, 8, 93450-93463.	1.8	23
17	Transcriptional dissection of melanoma identifies a high-risk subtype underlying TP53 family genes and epigenome deregulation. <i>JCI Insight</i> , 2017, 2, .	5.0	48
18	Microsatellite Instability Is Common in Colorectal Cancer in Native Nigerians. <i>Anticancer Research</i> , 2017, 37, 2649-2654.	1.1	22

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19	Fast detection of deletion breakpoints using quantitative PCR. <i>Genetics and Molecular Biology</i> , 2016, 39, 365-369.	1.3	1
20	Tu2061 Whole-Exome Analysis of Hereditary Microsatellite-Stable Colorectal Cancer in Israel. <i>Gastroenterology</i> , 2016, 150, S1013.	1.3	0
21	Early onset pancreatic malignancies: Clinical characteristics and survival associations. <i>International Journal of Cancer</i> , 2016, 139, 2169-2177.	5.1	22
22	Identification of Susceptibility Loci and Genes for Colorectal Cancer Risk. <i>Gastroenterology</i> , 2016, 150, 1633-1645.	1.3	97
23	A Germline Variant on Chromosome 4q31.1 Associates with Susceptibility to Developing Colon Cancer Metastasis. <i>PLoS ONE</i> , 2016, 11, e0146435.	2.5	2
24	Genome-wide association study of colorectal cancer identifies six new susceptibility loci. <i>Nature Communications</i> , 2015, 6, 7138.	12.8	138
25	Genome measures used for quality control are dependent on gene function and ancestry. <i>Bioinformatics</i> , 2015, 31, 318-323.	4.1	134
26	Abstract 2745: Exome sequencing analysis of 41 patients with Familial Colorectal Cancer Type X (FCCTX). , 2015, . .		1
27	Blood BDNF Level Is Gender Specific in Severe Depression. <i>PLoS ONE</i> , 2015, 10, e0127643.	2.5	73
28	A novel colorectal cancer risk locus at 4q32.2 identified from an international genome-wide association study. <i>Carcinogenesis</i> , 2014, 35, 2512-2519.	2.8	30
29	Correction: Interaction of Fatty Acid Genotype and Diet on Changes in Colonic Fatty Acids in a Mediterranean Diet Intervention Study. <i>Cancer Prevention Research</i> , 2014, 7, 372-372.	1.5	0
30	Distinct molecular features of colorectal cancer in Ghana. <i>Cancer Epidemiology</i> , 2013, 37, 556-561.	1.9	31
31	Transcriptome Profiling Identifies HMGA2 as a Biomarker of Melanoma Progression and Prognosis. <i>Journal of Investigative Dermatology</i> , 2013, 133, 2585-2592.	0.7	96
32	Interaction of Fatty Acid Genotype and Diet on Changes in Colonic Fatty Acids in a Mediterranean Diet Intervention Study. <i>Cancer Prevention Research</i> , 2013, 6, 1212-1221.	1.5	24
33	A Pilot Study of Microsatellite Instability and Endometrial Cancer Survival in White and African American Women. <i>International Journal of Gynecological Pathology</i> , 2012, 31, 66-72.	1.4	5
34	Risk of Non-Melanoma Cancers in First-Degree Relatives of CDKN2A Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2012, 104, 953-956.	6.3	42
35	Abstract B92: Molecular characterization of colorectal cancer in Ghana. <i>Cancer Prevention Research</i> , 2012, 5, B92-B92.	1.5	2
36	Identification and functional characterization of a novel MUTYH gene mutation.. <i>Journal of Clinical Oncology</i> , 2012, 30, e12026-e12026.	1.6	0

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37	Copy Number Variations and Clinical Outcome in Atypical Spitz Tumors. American Journal of Surgical Pathology, 2011, 35, 243-252.	3.7	105
38	Characterization of two Ashkenazi Jewish founder mutations in MSH6 gene causing Lynch syndrome. Clinical Genetics, 2011, 79, 512-522.	2.0	27
39	<i>MRE11</i> Deficiency Increases Sensitivity to Poly(ADP-ribose) Polymerase Inhibition in Microsatellite Unstable Colorectal Cancers. Cancer Research, 2011, 71, 2632-2642.	0.9	140
40	Disclosing Individual CDKN2A Research Results to Melanoma Survivors: Interest, Impact, and Demands on Researchers. Cancer Epidemiology Biomarkers and Prevention, 2011, 20, 522-529.	2.5	37
41	Human papillomavirus is not associated with colorectal cancer in a large international study. Cancer Causes and Control, 2010, 21, 737-743.	1.8	60
42	Returning Individual Research Results: Development of a Cancer Genetics Education and Risk Communication Protocol. Journal of Empirical Research on Human Research Ethics, 2010, 5, 17-30.	1.3	26
43	Abstract 2975: Two putative founder MSH6 mutations associated with inherited cancer susceptibility in Ashkenazi Jewish population. , 2010, , .		0
44	Abstract LB-349: Characterization of copy number variations in atypical Spitz tumors. , 2010, , .		0
45	Gene Expression Patterns in Mismatch Repair-Deficient Colorectal Cancers Highlight the Potential Therapeutic Role of Inhibitors of the Phosphatidylinositol 3-Kinase-AKT-Mammalian Target of Rapamycin Pathway. Clinical Cancer Research, 2009, 15, 2829-2839.	7.0	57
46	<i>BRCA1</i> Breast Cancer Risk Is Modified by <i>CYP19</i> Polymorphisms in Ashkenazi Jews. Cancer Epidemiology Biomarkers and Prevention, 2009, 18, 1617-1623.	2.5	11
47	FOXP3 germline polymorphisms are not associated with risk of breast cancer. Cancer Genetics and Cytogenetics, 2009, 190, 40-42.	1.0	26
48	Pediatric duodenal cancer and biallelic mismatch repair gene mutations. Pediatric Blood and Cancer, 2009, 53, 116-120.	1.5	13
49	<i>FGFR2</i> Is a Breast Cancer Susceptibility Gene in Jewish and Arab Israeli Populations. Cancer Epidemiology Biomarkers and Prevention, 2008, 17, 1060-1065.	2.5	52
50	Recreational Physical Activity Modifies the Association Between a Common GH1 Polymorphism and Colorectal Cancer Risk. Cancer Epidemiology Biomarkers and Prevention, 2008, 17, 3314-3318.	2.5	16
51	RAD51 135G→C Modifies Breast Cancer Risk among BRCA2 Mutation Carriers: Results from a Combined Analysis of 19 Studies. American Journal of Human Genetics, 2007, 81, 1186-1200.	6.2	217