

Brian D Lehmann

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

61
papers

9,294
citations

34
h-index

67
g-index

67
ext. papers

10,925
ext. citations

7.2
avg, IF

5.91
L-index

#	Paper	IF	Citations
61	Abstract PD3-04: Multi-omics characterization of triple-negative breast cancer identifies therapeutic vulnerabilities and epigenetic immune suppression in the mesenchymal subtype. <i>Cancer Research</i> , 2022 , 82, PD3-04-PD3-04	10.1	
60	Abstract P5-09-01: Using isogenic model systems to determine mechanisms regulating mutant p53 protein stability in breast cancer cells. <i>Cancer Research</i> , 2022 , 82, P5-09-01-P5-09-01	10.1	
59	Multi-omics analysis identifies therapeutic vulnerabilities in triple-negative breast cancer subtypes. <i>Nature Communications</i> , 2021 , 12, 6276	17.4	10
58	Tissue-specific expression of p73 and p63 isoforms in human tissues. <i>Cell Death and Disease</i> , 2021 , 12, 745	9.8	0
57	LR Hunting: A Random Forest Based Cell-Cell Interaction Discovery Method for Single-Cell Gene Expression Data. <i>Frontiers in Genetics</i> , 2021 , 12, 708835	4.5	1
56	Acquisition of aneuploidy drives mutant p53-associated gain-of-function phenotypes. <i>Nature Communications</i> , 2021 , 12, 5184	17.4	5
55	Targeting MYCN-expressing triple-negative breast cancer with BET and MEK inhibitors. <i>Science Translational Medicine</i> , 2020 , 12,	17.5	22
54	TBCRC 032 IB/II Multicenter Study: Molecular Insights to AR Antagonist and PI3K Inhibitor Efficacy in Patients with AR Metastatic Triple-Negative Breast Cancer. <i>Clinical Cancer Research</i> , 2020 , 26, 2111-2123	12.9	41
53	Reciprocal expression of Annexin A6 and RasGRF2 discriminates rapidly growing from invasive triple negative breast cancer subsets. <i>PLoS ONE</i> , 2020 , 15, e0231711	3.7	5
52	Identification of Targetable Recurrent MAP3K8 Rearrangements in Melanomas Lacking Known Driver Mutations. <i>Molecular Cancer Research</i> , 2019 , 17, 1842-1853	6.6	7
51	Implication of calcium activated RasGRF2 in Annexin A6-mediated breast tumor cell growth and motility. <i>Oncotarget</i> , 2019 , 10, 133-151	3.3	5
50	The Landscape of Small Non-Coding RNAs in Triple-Negative Breast Cancer. <i>Genes</i> , 2018 , 9,	4.2	14
49	A Randomized Phase II Neoadjuvant Study of Cisplatin, Paclitaxel With or Without Everolimus in Patients with Stage II/III Triple-Negative Breast Cancer (TNBC): Responses and Long-term Outcome Correlated with Increased Frequency of DNA Damage Response Gene Mutations, TNBC Subtype, AR Status, and Ki67. <i>Clinical Cancer Research</i> , 2017 , 23, 4035-4045	12.9	79
48	Comparison of triple-negative breast cancer molecular subtyping using RNA from matched fresh-frozen versus formalin-fixed paraffin-embedded tissue. <i>BMC Cancer</i> , 2017 , 17, 241	4.8	20
47	The Utilization of Formalin Fixed-Paraffin-Embedded Specimens in High Throughput Genomic Studies. <i>International Journal of Genomics</i> , 2017 , 2017, 1926304	2.5	44
46	Estimating relative mitochondrial DNA copy number using high throughput sequencing data. <i>Genomics</i> , 2017 , 109, 457-462	4.3	13
45	Mitochondria sequence mapping strategies and practicability of mitochondria variant detection from exome and RNA sequencing data. <i>Briefings in Bioinformatics</i> , 2016 , 17, 224-32	13.4	23

44	Generation of an algorithm based on minimal gene sets to clinically subtype triple negative breast cancer patients. <i>BMC Cancer</i> , 2016 , 16, 143	4.8	42
43	RNA Sequencing of Formalin-Fixed, Paraffin-Embedded Specimens for Gene Expression Quantification and Data Mining. <i>International Journal of Genomics</i> , 2016 , 2016, 9837310	2.5	14
42	Refinement of Triple-Negative Breast Cancer Molecular Subtypes: Implications for Neoadjuvant Chemotherapy Selection. <i>PLoS ONE</i> , 2016 , 11, e0157368	3.7	590
41	Diverse, Biologically Relevant, and Targetable Gene Rearrangements in Triple-Negative Breast Cancer and Other Malignancies. <i>Cancer Research</i> , 2016 , 76, 4850-60	10.1	28
40	Clinical implications of molecular heterogeneity in triple negative breast cancer. <i>Breast</i> , 2015 , 24 Suppl 2, S36-40	3.6	76
39	Practicality of identifying mitochondria variants from exome and RNAseq data. <i>BMC Bioinformatics</i> , 2015 , 16, P6	3.6	3
38	RNAseq by Total RNA Library Identifies Additional RNAs Compared to Poly(A) RNA Library. <i>BioMed Research International</i> , 2015 , 2015, 862130	3	26
37	Triple-negative breast cancer: molecular subtypes and new targets for therapy. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2015 , e31-9	7.1	83
36	Subtyping of triple-negative breast cancer: implications for therapy. <i>Cancer</i> , 2015 , 121, 8-16	6.4	212
35	A Synthetic Lethal Screen Identifies DNA Repair Pathways that Sensitize Cancer Cells to Combined ATR Inhibition and Cisplatin Treatments. <i>PLoS ONE</i> , 2015 , 10, e0125482	3.7	76
34	Aberrant over-expression of COX-1 intersects multiple pro-tumorigenic pathways in high-grade serous ovarian cancer. <i>Oncotarget</i> , 2015 , 6, 21353-68	3.3	31
33	Identification and use of biomarkers in treatment strategies for triple-negative breast cancer subtypes. <i>Journal of Pathology</i> , 2014 , 232, 142-50	9.4	267
32	New strategies for triple-negative breast cancer--deciphering the heterogeneity. <i>Clinical Cancer Research</i> , 2014 , 20, 782-90	12.9	210
31	PIK3CA mutations in androgen receptor-positive triple negative breast cancer confer sensitivity to the combination of PI3K and androgen receptor inhibitors. <i>Breast Cancer Research</i> , 2014 , 16, 406	8.3	199
30	Mislocalization of the cell polarity protein scribble promotes mammary tumorigenesis and is associated with basal breast cancer. <i>Cancer Research</i> , 2014 , 74, 3180-94	10.1	76
29	Molecular profiling of the residual disease of triple-negative breast cancers after neoadjuvant chemotherapy identifies actionable therapeutic targets. <i>Cancer Discovery</i> , 2014 , 4, 232-45	24.4	310
28	Transforming growth factor beta receptor type III is a tumor promoter in mesenchymal-stem like triple negative breast cancer. <i>Breast Cancer Research</i> , 2014 , 16, R69	8.3	37
27	Detection of internal exon deletion with exon Del. <i>BMC Bioinformatics</i> , 2014 , 15, 332	3.6	10

26	Multi-perspective quality control of Illumina exome sequencing data using QC3. <i>Genomics</i> , 2014 , 103, 323-8	4.3	61
25	Identification of prognosis-relevant subgroups in patients with chemoresistant triple-negative breast cancer. <i>Clinical Cancer Research</i> , 2013 , 19, 2723-33	12.9	119
24	Patient-derived breast tumor xenografts facilitating personalized cancer therapy. <i>Breast Cancer Research</i> , 2013 , 15, 201	8.3	69
23	SPARCL1 suppresses metastasis in prostate cancer. <i>Molecular Oncology</i> , 2013 , 7, 1019-30	7.9	23
22	Comparative study of exome copy number variation estimation tools using array comparative genomic hybridization as control. <i>BioMed Research International</i> , 2013 , 2013, 915636	3	36
21	BRAF fusions define a distinct molecular subset of melanomas with potential sensitivity to MEK inhibition. <i>Clinical Cancer Research</i> , 2013 , 19, 6696-702	12.9	122
20	Differential response to neoadjuvant chemotherapy among 7 triple-negative breast cancer molecular subtypes. <i>Clinical Cancer Research</i> , 2013 , 19, 5533-40	12.9	476
19	A data similarity-based strategy for meta-analysis of transcriptional profiles in cancer. <i>PLoS ONE</i> , 2013 , 8, e54979	3.7	4
18	Differential pathologic complete response rates after neoadjuvant chemotherapy among molecular subtypes of triple-negative breast cancer.. <i>Journal of Clinical Oncology</i> , 2013 , 31, 1005-1005	2.2	5
17	TNBCtype: A Subtyping Tool for Triple-Negative Breast Cancer. <i>Cancer Informatics</i> , 2012 , 11, 147-56	2.4	161
16	p53 expression controls prostate cancer sensitivity to chemotherapy and the MDM2 inhibitor Nutlin-3. <i>Cell Cycle</i> , 2012 , 11, 4579-88	4.7	73
15	Targeting mutant p53 in human tumors. <i>Journal of Clinical Oncology</i> , 2012 , 30, 3648-50	2.2	59
14	Identification of human triple-negative breast cancer subtypes and preclinical models for selection of targeted therapies. <i>Journal of Clinical Investigation</i> , 2011 , 121, 2750-67	15.9	3125
13	Cooperative effects of Akt-1 and Raf-1 on the induction of cellular senescence in doxorubicin or tamoxifen treated breast cancer cells. <i>Oncotarget</i> , 2011 , 2, 610-26	3.3	38
12	Attenuation of myocardial injury in mice with functional deletion of the circadian rhythm gene mPer2. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2010 , 298, H1088-95	5.2	34
11	RNA interference (RNAi) screening approach identifies agents that enhance paclitaxel activity in breast cancer cells. <i>Breast Cancer Research</i> , 2010 , 12, R41	8.3	58
10	Suppression of PTEN function increases breast cancer chemotherapeutic drug resistance while conferring sensitivity to mTOR inhibitors. <i>Oncogene</i> , 2008 , 27, 4086-95	9.2	128
9	Senescence-associated exosome release from human prostate cancer cells. <i>Cancer Research</i> , 2008 , 68, 7864-71	10.1	310

8	Targeting prostate cancer based on signal transduction and cell cycle pathways. <i>Cell Cycle</i> , 2008 , 7, 1745-47	4.7	80
7	Distinct roles for p107 and p130 in Rb-independent cellular senescence. <i>Cell Cycle</i> , 2008 , 7, 1262-8	4.7	15
6	Alteration of Akt activity increases chemotherapeutic drug and hormonal resistance in breast cancer yet confers an achilles heel by sensitization to targeted therapy. <i>Advances in Enzyme Regulation</i> , 2008 , 48, 113-35		20
5	Targeting Survival Cascades Induced by Activation of Ras/Raf/MEK/ERK and PI3K/Akt Pathways to Sensitize Cancer Cells to Therapy 2008 , 81-114		
4	Roles of the Raf/MEK/ERK pathway in cell growth, malignant transformation and drug resistance. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2007 , 1773, 1263-84	4.9	1532
3	Targeting the RAF/MEK/ERK, PI3K/AKT and p53 pathways in hematopoietic drug resistance. <i>Advances in Enzyme Regulation</i> , 2007 , 47, 64-103		63
2	A dominant role for p53-dependent cellular senescence in radiosensitization of human prostate cancer cells. <i>Cell Cycle</i> , 2007 , 6, 595-605	4.7	80
1	Radiosensitization of prostate cancer by priming the wild-type p53-dependent cellular senescence pathway. <i>Cancer Biology and Therapy</i> , 2007 , 6, 1165-70	4.6	12