Traver Hart

List of Publications by Year in descending order

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53 6,510 29
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72

docs citations

h-index g-index

72 11021
times ranked citing authors

182427

51

#	Article	IF	CITATIONS
1	HumanNet v3: an improved database of human gene networks for disease research. Nucleic Acids Research, 2022, 50, D632-D639.	14.5	53
2	Genome-wide CRISPR screens using isogenic cells reveal vulnerabilities conferred by loss of tumor suppressors. Science Advances, 2022, 8, eabm6638.	10.3	17
3	Integrated screens uncover a cell surface tumor suppressor gene <i>KIRREL</i> involved in Hippo pathway. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, .	7.1	10
4	Dynamic rewiring of biological activity across genotype and lineage revealed by context-dependent functional interactions. Genome Biology, 2022, 23, .	8.8	9
5	Genetic vulnerabilities upon inhibition of DNA damage response. Nucleic Acids Research, 2021, 49, 8214-8231.	14.5	17
6	AR-negative prostate cancer is vulnerable to loss of JMJD1C demethylase. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118 , .	7.1	6
7	Improved analysis of CRISPR fitness screens and reduced off-target effects with the BAGEL2 gene essentiality classifier. Genome Medicine, 2021, 13, 2.	8.2	60
8	Discovery of putative tumor suppressors from CRISPR screens reveals rewired lipid metabolism in acute myeloid leukemia cells. Nature Communications, 2021, 12, 6506.	12.8	13
9	Common computational tools for analyzing CRISPR screens. Emerging Topics in Life Sciences, 2021, 5, 779-788.	2.6	10
10	Multiplex enCas12a screens detect functional buffering among paralogs otherwise masked in monogenic Cas9 knockout screens. Genome Biology, 2020, 21, 262.	8.8	62
11	Copper bioavailability is a KRAS-specific vulnerability in colorectal cancer. Nature Communications, 2020, 11, 3701.	12.8	128
12	C17orf53 is identified as a novel gene involved in inter-strand crosslink repair. DNA Repair, 2020, 95, 102946.	2.8	14
13	Functional genomic landscape of cancer-intrinsic evasion of killing by T cells. Nature, 2020, 586, 120-126.	27.8	249
14	DNA polymerase \hat{l}^1 compensates for Fanconi anemia pathway deficiency by countering DNA replication stress. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 33436-33445.	7.1	13
15	CRISPR/CAS9-based DNA damage response screens reveal gene-drug interactions. DNA Repair, 2020, 87, 102803.	2.8	23
16	Genome-wide InÂVivo CNS Screening Identifies Genes that Modify CNS Neuronal Survival and mHTT Toxicity. Neuron, 2020, 106, 76-89.e8.	8.1	62
17	Genome-wide CRISPR screen uncovers a synergistic effect of combining Haspin and Aurora kinase B inhibition. Oncogene, 2020, 39, 4312-4322.	5.9	16
18	IPO11 mediates \hat{I}^2 catenin nuclear import in a subset of colorectal cancers. Journal of Cell Biology, 2020, 219, .	5.2	27

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19	Identifying chemogenetic interactions from CRISPR screens with drugZ. Genome Medicine, 2019, 11, 52.	8.2	127
20	Pooled library screening with multiplexed Cpf1 library. Nature Communications, 2019, 10, 3144.	12.8	37
21	A consensus set of genetic vulnerabilities to ATR inhibition. Open Biology, 2019, 9, 190156.	3.6	81
22	Genome-Wide CRISPR-Cas9 Screens Expose Genetic Vulnerabilities and Mechanisms of Temozolomide Sensitivity in Glioblastoma Stem Cells. Cell Reports, 2019, 27, 971-986.e9.	6.4	139
23	Chemogenetic interactions in human cancer cells. Computational and Structural Biotechnology Journal, 2019, 17, 1318-1325.	4.1	8
24	HumanNet v2: human gene networks for disease research. Nucleic Acids Research, 2019, 47, D573-D580.	14.5	161
25	A network of human functional gene interactions from knockout fitness screens in cancer cells. Life Science Alliance, 2019, 2, e201800278.	2.8	81
26	PICKLES: the database of pooled in-vitro CRISPR knockout library essentiality screens. Nucleic Acids Research, 2018, 46, D776-D780.	14.5	74
27	Microdroplet PCR for Highly Multiplexed Targeted Bisulfite Sequencing. Methods in Molecular Biology, 2018, 1708, 333-348.	0.9	2
28	Expression of the miR-150 tumor suppressor is restored by and synergizes with rapamycin in a human leukemia T-cell line. Leukemia Research, 2018, 74, 1-9.	0.8	9
29	CRISPR screens identify genomic ribonucleotides as a source of PARP-trapping lesions. Nature, 2018, 559, 285-289.	27.8	297
30	The shieldin complex mediates 53BP1-dependent DNA repair. Nature, 2018, 560, 117-121.	27.8	445
31	Identification of FZD5 as Genetic Vulnerability in RNF43 Mutant Cancer. FASEB Journal, 2018, 32, 804.31.	0.5	0
32	All for One, and One for All. Cell Systems, 2017, 5, 314-316.	6.2	2
33	Evaluation and Design of Genome-Wide CRISPR/SpCas9 Knockout Screens. G3: Genes, Genomes, Genetics, 2017, 7, 2719-2727.	1.8	417
34	Genome-wide CRISPR screens reveal a Wnt–FZD5 signaling circuit as a druggable vulnerability of RNF43-mutant pancreatic tumors. Nature Medicine, 2017, 23, 60-68.	30.7	261
35	BAGEL: a computational framework for identifying essential genes from pooled library screens. BMC Bioinformatics, 2016, 17, 164.	2.6	216
36	Extensive mapping of an innate immune network with <scp>CRISPR</scp> . Molecular Systems Biology, 2015, 11, 821.	7. 2	2

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37	Genome-wide CRISPR-Cas9 Screens Reveal Loss of Redundancy between PKMYT1 and WEE1 in Glioblastoma Stem-like Cells. Cell Reports, 2015, 13, 2425-2439.	6.4	146
38	Defining CD4 T Cell Memory by the Epigenetic Landscape of CpG DNA Methylation. Journal of Immunology, 2015, 194, 1565-1579.	0.8	59
39	Natural Variation in Gene Expression Modulates the Severity of Mutant Phenotypes. Cell, 2015, 162, 391-402.	28.9	129
40	High-Resolution CRISPR Screens Reveal Fitness Genes and Genotype-Specific Cancer Liabilities. Cell, 2015, 163, 1515-1526.	28.9	1,339
41	Genome-wide RNAi analysis reveals that simultaneous inhibition of specific mevalonate pathway genes potentiates tumor cell death. Oncotarget, 2015, 6, 26909-26921.	1.8	52
42	Measuring error rates in genomic perturbation screens: gold standards for human functional genomics. Molecular Systems Biology, 2014, 10, 733.	7.2	322
43	Finding the active genes in deep RNA-seq gene expression studies. BMC Genomics, 2013, 14, 778.	2.8	193
44	Scaling up the systematic hunt for mammalian genetic interactions. Nature Methods, 2013, 10, 397-399.	19.0	4
45	A Census of Human Soluble Protein Complexes. Cell, 2012, 150, 1068-1081.	28.9	781
46	Method for improved Illumina sequencing library preparation using NuGEN Ovation RNA-Seq System. BioTechniques, 2011, 50, 177-181.	1.8	27
47	Application of microdroplet PCR for large-scale targeted bisulfite sequencing. Genome Research, 2011, 21, 1738-1745.	5.5	36
48	MicroRNA Regulation of Molecular Networks Mapped by Global MicroRNA, mRNA, and Protein Expression in Activated T Lymphocytes. Journal of Immunology, 2011, 187, 2233-2243.	0.8	86
49	Systematic Definition of Protein Constituents along the Major Polarization Axis Reveals an Adaptive Reuse of the Polarization Machinery in Pheromone-Treated Budding Yeast. Journal of Proteome Research, 2009, 8, 6-19.	3.7	28
50	Human Cell Chips: Adapting DNA Microarray Spotting Technology to Cell-Based Imaging Assays. PLoS ONE, 2009, 4, e7088.	2.5	22
51	A map of human protein interactions derived from coâ€expression of human mRNAs and their orthologs. Molecular Systems Biology, 2008, 4, 180.	7.2	77
52	Systematic profiling of cellular phenotypes and gene function using spotted cellular microarrays. FASEB Journal, 2006, 20, LB61.	0.5	0
53	The Functional Genomic Circuitry of Human Glioblastoma Stem Cells. SSRN Electronic Journal, 0, , .	0.4	0