

Jason W Hoskins

List of Publications by Year in descending order

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Version: 2024-02-01

29
papers

1,342
citations

361413

20
h-index

610901

24
g-index

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all docs

30
docs citations

30
times ranked

2912
citing authors

#	ARTICLE	IF	CITATIONS
1	A 584Åbp deletion in CTRB2 inhibits chymotrypsin B2 activity and secretion and confers risk of pancreatic cancer. <i>American Journal of Human Genetics</i> , 2021, 108, 1852-1865.	6.2	15
2	Inferred expression regulator activities suggest genes mediating cardiometabolic genetic signals. <i>PLoS Computational Biology</i> , 2021, 17, e1009563.	3.2	3
3	A Transcriptome-Wide Association Study Identifies Novel Candidate Susceptibility Genes for Pancreatic Cancer. <i>Journal of the National Cancer Institute</i> , 2020, 112, 1003-1012.	6.3	59
4	Genome-wide meta-analysis identifies five new susceptibility loci for pancreatic cancer. <i>Nature Communications</i> , 2018, 9, 556.	12.8	188
5	Characterising <i>cis</i> -regulatory variation in the transcriptome of histologically normal and tumour-derived pancreatic tissues. <i>Gut</i> , 2018, 67, 521-533.	12.1	26
6	Functional characterization of a multi-cancer risk locus on chr5p15.33 reveals regulation of TERT by ZNF148. <i>Nature Communications</i> , 2017, 8, 15034.	12.8	40
7	Abstract 1442: Analysis of <i>cis</i> -eQTLs in normal and tumor-derived pancreatic tissues reveals functional insights, including for the 9q34.1ABOpancreatic cancer risk locus. , 2017, , .		0
8	Three new pancreatic cancer susceptibility signals identified on chromosomes 1q32.1, 5p15.33 and 8q24.21. <i>Oncotarget</i> , 2016, 7, 66328-66343.	1.8	88
9	Functional characterization of a chr13q22.1 pancreatic cancer risk locus reveals long-range interaction and allele-specific effects on <i>DIS3</i> expression. <i>Human Molecular Genetics</i> , 2016, 25, ddw300.	2.9	24
10	Abstract B111: Functional analysis of the chr13q22.1 pancreatic cancer risk locus suggests allele-specific effects on DIS3 expression with prognostic implications. , 2015, , .		0
11	Abstract B112: Transcriptome analysis in pancreatic cancer reveals a tumor suppressor function for HNF1A. , 2015, , .		0
12	Abstract 4610: Functional characterization of a multicancer risk locus on chr5p15.33 reveals regulation of TERT by ZNF148. , 2015, , .		0
13	Abstract A1-09: Functional analysis of the chr13q22.1 pancreatic cancer risk locus suggests allele-specific effects on DIS3 expression with prognostic implications. , 2015, , .		0
14	Transcriptome analysis of pancreatic cancer reveals a tumor suppressor function for HNF1A. <i>Carcinogenesis</i> , 2014, 35, 2670-2678.	2.8	46
15	Imputation and subset-based association analysis across different cancer types identifies multiple independent risk loci in the TERT-CLPTM1L region on chromosome 5p15.33. <i>Human Molecular Genetics</i> , 2014, 23, 6616-6633.	2.9	90
16	Genetic Susceptibility and Risk of Pancreatic Cancer. , 2014, , 169-194.		0
17	CLPTM1L Promotes Growth and Enhances Aneuploidy in Pancreatic Cancer Cells. <i>Cancer Research</i> , 2014, 74, 2785-2795.	0.9	48
18	Lomofungin and dilomofungin: inhibitors of MBNL1-CUG RNA binding with distinct cellular effects. <i>Nucleic Acids Research</i> , 2014, 42, 6591-6602.	14.5	46

#	ARTICLE	IF	CITATIONS
19	Induction and reversal of myotonic dystrophy type 1 pre-mRNA splicing defects by small molecules. <i>Nature Communications</i> , 2013, 4, 2044.	12.8	76
20	An integrated transcriptome and epigenome analysis identifies a novel candidate gene for pancreatic cancer. <i>BMC Medical Genomics</i> , 2013, 6, 33.	1.5	31
21	A Resequencing Analysis of Genomic Loci on Chromosomes 1q32.1, 5p15.33, and 13q22.1 Associated With Pancreatic Cancer Risk. <i>Pancreas</i> , 2013, 42, 209-215.	1.1	5
22	From dynamic combinatorial <i>hit</i> ™ to lead: in vitro and in vivo activity of compounds targeting the pathogenic RNAs that cause myotonic dystrophy. <i>Nucleic Acids Research</i> , 2012, 40, 6380-6390.	14.5	69
23	Design of a Bioactive Small Molecule That Targets the Myotonic Dystrophy Type 1 RNA via an RNA Motif—Ligand Database and Chemical Similarity Searching. <i>Journal of the American Chemical Society</i> , 2012, 134, 4731-4742.	13.7	129
24	Rationally Designed Small Molecules Targeting the RNA That Causes Myotonic Dystrophy Type 1 Are Potently Bioactive. <i>ACS Chemical Biology</i> , 2012, 7, 856-862.	3.4	112
25	Two high-throughput screening assays for aberrant RNA–protein interactions in myotonic dystrophy type 1. <i>Analytical and Bioanalytical Chemistry</i> , 2012, 402, 1889-1898.	3.7	49
26	Properties of Nat4, an N ¹ -Acetyltransferase of <i>Saccharomyces cerevisiae</i> That Modifies N Termini of Histones H2A and H4. <i>Molecular and Cellular Biology</i> , 2009, 29, 2913-2924.	2.3	45
27	RNA-Based 5-Fluorouracil Toxicity Requires the Pseudouridylation Activity of Cbf5p. <i>Genetics</i> , 2008, 179, 323-330.	2.9	37
28	Evidence for distinct DNA- and RNA-based mechanisms of 5-fluorouracil cytotoxicity in <i>Saccharomyces cerevisiae</i> . <i>Yeast</i> , 2007, 24, 861-870.	1.7	39
29	5-Fluorouracil Enhances Exosome-Dependent Accumulation of Polyadenylated rRNAs. <i>Molecular and Cellular Biology</i> , 2004, 24, 10766-10776.	2.3	75