

Thomas R Webb

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

1,205
citations

1051969

10
h-index

1051228

16
g-index

17
all docs

17
docs citations

17
times ranked

2436
citing authors

#	ARTICLE	IF	CITATIONS
1	The spliceosome is a therapeutic vulnerability in MYC-driven cancer. <i>Nature</i> , 2015, 525, 384-388.	13.7	392
2	Anaplastic lymphoma kinase: role in cancer pathogenesis and small-molecule inhibitor development for therapy. <i>Expert Review of Anticancer Therapy</i> , 2009, 9, 331-356.	1.1	206
3	Sudemycins, Novel Small Molecule Analogues of FR901464, Induce Alternative Gene Splicing. <i>ACS Chemical Biology</i> , 2011, 6, 582-589.	1.6	155
4	Mutant U2AF1-expressing cells are sensitive to pharmacological modulation of the spliceosome. <i>Nature Communications</i> , 2017, 8, 14060.	5.8	99
5	The development and application of small molecule modulators of SF3b as therapeutic agents for cancer. <i>Drug Discovery Today</i> , 2013, 18, 43-49.	3.2	89
6	The splicing modulator sudemycin induces a specific antitumor response and cooperates with ibrutinib in chronic lymphocytic leukemia. <i>Oncotarget</i> , 2015, 6, 22734-22749.	0.8	60
7	Optimization of Antitumor Modulators of Pre-mRNA Splicing. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 10033-10044.	2.9	57
8	Inhibition of SF3B1 by molecules targeting the spliceosome results in massive aberrant exon skipping. <i>Rna</i> , 2018, 24, 1056-1066.	1.6	42
9	USP39 Deubiquitinase Is Essential for KRAS Oncogene-driven Cancer. <i>Journal of Biological Chemistry</i> , 2017, 292, 4164-4175.	1.6	37
10	A triple exon-skipping luciferase reporter assay identifies a new CLK inhibitor pharmacophore. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 406-412.	1.0	16
11	Hereditary retinoblastoma iPSC model reveals aberrant spliceosome function driving bone malignancies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2117857119.	3.3	13
12	Pharmacodynamic assays to facilitate preclinical and clinical development of pre-mRNA splicing modulatory drug candidates. <i>Pharmacology Research and Perspectives</i> , 2015, 3, e00158.	1.1	12
13	An exon skipping screen identifies antitumor drugs that are potent modulators of pre-mRNA splicing, suggesting new therapeutic applications. <i>PLoS ONE</i> , 2020, 15, e0233672.	1.1	11
14	Changes in Alternative Splicing as Pharmacodynamic Markers for Sudemycin D6. <i>Biomarker Insights</i> , 2017, 12, 117727191773055.	1.0	9
15	Improving the Efficiency of the Drug Development by Expanding the Scope of the Role of Medicinal Chemists in Drug Discovery. <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 1153-1155.	1.3	4
16	Sudemycin Selectively Inhibits Growth of Primary Murine Hematopoietic Cells Expressing Mutant U2AF1. <i>Blood</i> , 2012, 120, 554-554.	0.6	3