Scott T Younger

List of Publications by Year in descending order

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933447 1058476 14 793 10 14 citations h-index g-index papers 14 14 14 1767 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Multiplexable, locus-specific targeting of long RNAs with CRISPR-Display. Nature Methods, 2015, 12, 664-670.	19.0	268
2	Deubiquitinases Maintain Protein Homeostasis and Survival of Cancer Cells upon Glutathione Depletion. Cell Metabolism, 2019, 29, 1166-1181.e6.	16.2	121
3	Cells Lacking the <i>RB1</i> Tumor Suppressor Gene Are Hyperdependent on Aurora B Kinase for Survival. Cancer Discovery, 2019, 9, 230-247.	9.4	119
4	Integrative genomic analysis reveals widespread enhancer regulation by p53 in response to DNA damage. Nucleic Acids Research, 2015, 43, 4447-4462.	14.5	84
5	p53 regulates enhancer accessibility and activity in response to DNA damage. Nucleic Acids Research, 2017, 45, 9889-9900.	14.5	61
6	Genomic answers for children: Dynamic analyses of >1000 pediatric rare disease genomes. Genetics in Medicine, 2022, 24, 1336-1348.	2.4	37
7	'Lnc'-ing enhancers to MYC regulation. Cell Research, 2014, 24, 643-644.	12.0	23
8	SMAD4 represses FOSL1 expression and pancreatic cancer metastatic colonization. Cell Reports, 2021, 36, 109443.	6.4	23
9	Pooled Lentiviralâ€Delivery Genetic Screens. Current Protocols in Molecular Biology, 2018, 121, 32.1.1-32.1.21.	2.9	20
10	Identification of functional regulatory elements in the human genome using pooled CRISPR screens. BMC Genomics, 2020, 21, 107.	2.8	12
11	Expediting rare disease diagnosis: a call to bridge the gap between clinical and functional genomics. Molecular Medicine, 2020, 26, 117 .	4.4	10
12	Silent pericentromeric repeats speak out. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, 15008-15009.	7.1	8
13	FOXC2 Autoregulates Its Expression in the Pulmonary Endothelium After Endotoxin Stimulation in a Histone Acetylation-Dependent Manner. Frontiers in Cell and Developmental Biology, 2021, 9, 657662.	3.7	6
14	Massively parallel identification of functionally consequential noncoding genetic variants in undiagnosed rare disease patients. Scientific Reports, 2022, 12, 7576.	3.3	1