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List of Publications by Year in descending order

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Version: 2024-02-01

12
papers

958
citations

759233

12
h-index

1199594

12
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12
all docs

12
docs citations

12
times ranked

1588
citing authors

#	ARTICLE	IF	CITATIONS
1	ATR-X Syndrome Protein Targets Tandem Repeats and Influences Allele-Specific Expression in a Size-Dependent Manner. <i>Cell</i> , 2010, 143, 367-378.	28.9	365
2	ATRX Plays a Key Role in Maintaining Silencing at Interstitial Heterochromatic Loci and Imprinted Genes. <i>Cell Reports</i> , 2015, 11, 405-418.	6.4	152
3	The chromatin remodelling factor <scp>ATRX</scp> suppresses Râ€ loops in transcribed telomeric repeats. <i>EMBO Reports</i> , 2017, 18, 914-928.	4.5	99
4	New players in heterochromatin silencing: histone variant H3.3 and the ATRX/DAXX chaperone. <i>Nucleic Acids Research</i> , 2016, 44, 1496-1501.	14.5	80
5	Ribosomal DNA copy loss and repeat instability in ATRX-mutated cancers. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, 4737-4742.	7.1	72
6	Inhibition of a K9/K36 demethylase by an H3.3 point mutation found in paediatric glioblastoma. <i>Nature Communications</i> , 2018, 9, 3142.	12.8	49
7	Controlling $\hat{\alpha}$ -globin: a review of $\hat{\alpha}$ -globin expression and its impact on $\hat{\alpha}$ -thalassemia. <i>Haematologica</i> , 2008, 93, 1868-1876.	3.5	46
8	siRNA-mediated reduction of $\hat{\alpha}$ -globin results in phenotypic improvements in $\hat{\alpha}$ -thalassemic cells. <i>Haematologica</i> , 2008, 93, 1238-1242.	3.5	23
9	Mutations inhibiting KDM4B drive ALT activation in ATRX-mutated glioblastomas. <i>Nature Communications</i> , 2021, 12, 2584.	12.8	23
10	Maintaining memory of silencing at imprinted differentially methylated regions. <i>Cellular and Molecular Life Sciences</i> , 2016, 73, 1871-1879.	5.4	18
11	Compromised Telomeric Heterochromatin Promotes ALternative Lengthening of Telomeres. <i>Trends in Cancer</i> , 2016, 2, 114-116.	7.4	17
12	Co-inheritance of $\hat{\alpha}^+$ - and $\hat{\alpha}^2$ -thalassaemia in mice ameliorates thalassaemic phenotype. <i>Blood Cells, Molecules, and Diseases</i> , 2007, 39, 184-188.	1.4	14