

Greta Pintacuda

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/60333/publications.pdf>

Version: 2024-02-01

16
papers

2,143
citations

623699

14
h-index

940516

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g-index

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all docs

20
docs citations

20
times ranked

2594
citing authors

#	ARTICLE	IF	CITATIONS
1	piRNAs Can Trigger a Multigenerational Epigenetic Memory in the Germline of <i>C.Âelegans</i> . <i>Cell</i> , 2012, 150, 88-99.	28.9	673
2	hnRNPk Recruits PCGF3/5-PRC1 to the Xist RNA B-Repeat to Establish Polycomb-Mediated Chromosomal Silencing. <i>Molecular Cell</i> , 2017, 68, 955-969.e10.	9.7	255
3	A Pooled shRNA Screen Identifies Rbm15, Spen, and Wtap as Factors Required for Xist RNA-Mediated Silencing. <i>Cell Reports</i> , 2015, 12, 562-572.	6.4	226
4	PCGF3/5-PRC1 initiates Polycomb recruitment in X chromosome inactivation. <i>Science</i> , 2017, 356, 1081-1084.	12.6	220
5	Xist localization and function: new insights from multiple levels. <i>Genome Biology</i> , 2015, 16, 166.	8.8	151
6	The nuclear matrix protein CIZ1 facilitates localization of Xist RNA to the inactive X-chromosome territory. <i>Genes and Development</i> , 2017, 31, 876-888.	5.9	104
7	Systematic allelic analysis defines the interplay of key pathways in X chromosome inactivation. <i>Nature Communications</i> , 2019, 10, 3129.	12.8	93
8	The non-canonical SMC protein SmcHD1 antagonises TAD formation and compartmentalisation on the inactive X chromosome. <i>Nature Communications</i> , 2019, 10, 30.	12.8	87
9	Function by Structure: Spotlights on Xist Long Non-coding RNA. <i>Frontiers in Molecular Biosciences</i> , 2017, 4, 90.	3.5	84
10	SmcHD1 Targeting to the Inactive X Is Dependent on the Xist-HnrnpK-PRC1 Pathway. <i>Cell Reports</i> , 2018, 25, 1912-1923.e9.	6.4	56
11	Coexpression network architecture reveals the brain-wide and multiregional basis of disease susceptibility. <i>Nature Neuroscience</i> , 2021, 24, 1313-1323.	14.8	44
12	Connecting TDP-43 Pathology with Neuropathy. <i>Trends in Neurosciences</i> , 2021, 44, 424-440.	8.6	42
13	Acute depletion of METTL3 implicates <i>m⁶A</i> -methyladenosine in alternative intron/exon inclusion in the nascent transcriptome. <i>Genome Research</i> , 2021, 31, 1395-1408.	5.5	37
14	Loss of mouse Stmn2 function causes motor neuropathy. <i>Neuron</i> , 2022, 110, 1671-1688.e6.	8.1	37
15	Mind the translational gap: using iPS cell models to bridge from genetic discoveries to perturbed pathways and therapeutic targets. <i>Molecular Autism</i> , 2021, 12, 10.	4.9	15
16	X Inactivation Lessons from Differentiating Mouse Embryonic Stem Cells. <i>Stem Cell Reviews and Reports</i> , 2015, 11, 699-705.	5.6	12