Andrew K Dingwall

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/1976877/publications.pdf

Version: 2024-02-01

23 papers 916 citations

430874 18 h-index 642732 23 g-index

26 all docs

 $\begin{array}{c} 26 \\ \\ \text{docs citations} \end{array}$

26 times ranked

1144 citing authors

#	Article	IF	CITATIONS
1	The cancer COMPASS: navigating the functions of MLL complexes in cancer. Cancer Genetics, 2015, 208, 178-191.	0.4	122
2	COMPASS Ascending: Emerging clues regarding the roles of MLL3/KMT2C and MLL2/KMT2D proteins in cancer. Cancer Letters, 2019, 458, 56-65.	7.2	121
3	Cascade regulation of Caulobacter flagellar and chemotaxis genes. Journal of Molecular Biology, 1987, 194, 71-80.	4.2	81
4	Plasmid and chromosomal DNA replication and partitioning during the Caulobacter crescentus cell cycle. Journal of Molecular Biology, 1990, 212, 709-722.	4.2	65
5	The Drosophila Brahma (SWI/SNF) chromatin remodeling complex exhibits cell-type specific activation and repression functions. Developmental Biology, 2004, 267, 279-293.	2.0	54
6	SNR1 is an essential subunit in a subset of drosophila brm complexes, targeting specific functions during development. Developmental Biology, 2003, 253, 291-308.	2.0	48
7	Hormone-response Genes Are Direct in Vivo Regulatory Targets of Brahma (SWI/SNF) Complex Function. Journal of Biological Chemistry, 2006, 281, 35305-35315.	3.4	44
8	Drosophila cyclin E interacts with components of the Brahma complex. EMBO Journal, 2002, 21, 3377-3389.	7.8	42
9	Organization and ordered expression of Caulobacter genes encoding flagellar basal body rod and ring proteins. Journal of Molecular Biology, 1992, 228, 1147-1162.	4.2	41
10	The chromatin remodeling and mRNA splicing functions of the Brahma (SWI/SNF) complex are mediated by the SNR1/SNF5 regulatory subunit. Nucleic Acids Research, 2012, 40, 5975-5987.	14.5	40
11	The Drosophila SNR1 (SNF5/INI1) Subunit Directs Essential Developmental Functions of the Brahma Chromatin Remodeling Complex. Molecular and Cellular Biology, 2003, 23, 289-305.	2.3	35
12	Histone recognition and nuclear receptor co-activator functions of <i>Drosophila </i> Cara Mitad, a homolog of the N-terminal portion of mammalian MLL2 and MLL3. Development (Cambridge), 2012, 139, 1997-2008.	2.5	35
13	Identification of Cis and Trans-elements involved in the timed control of a Caulobacter flagellar gene. Journal of Molecular Biology, 1991, 217, 247-257.	4.2	30
14	DAXX Suppresses Tumor-Initiating Cells in Estrogen Receptor–Positive Breast Cancer Following Endocrine Therapy. Cancer Research, 2019, 79, 4965-4977.	0.9	27
15	Temporal regulation and overlap organization of two Caulobacter flagellar genes. Journal of Molecular Biology, 1989, 205, 71-83.	4.2	26
16	Histone lysine demethylases function as co-repressors of SWI/SNF remodeling activities during Drosophila wing development. Developmental Biology, 2011, 350, 534-547.	2.0	26
17	SNR1 (INI1/SNF5) Mediates Important Cell Growth Functions of the Drosophila Brahma (SWI/SNF) Chromatin Remodeling Complex. Genetics, 2004, 168, 199-214.	2.9	22
18	Congenital anomalies and rhabdoid tumor associated with 22q11 germline deletion and somatic inactivation of the <i>SMARCB1</i> tumor suppressor. Genes Chromosomes and Cancer, 2011, 50, 379-388.	2.8	20

#	Article	IF	CITATIONS
19	The Drosophila COMPASS-like Cmi-Trr coactivator complex regulates dpp/BMP signaling in pattern formation. Developmental Biology, 2013, 380, 185-198.	2.0	13
20	The Drosophila MLR COMPASS complex is essential for programming cis-regulatory information and maintaining epigenetic memory during development. Nucleic Acids Research, 2020, 48, 3476-3495.	14.5	8
21	Genetic and Molecular Analysis of Region 88E9;88F2 in Drosophila melanogaster, Including the ear Gene Related to Human Factors Involved in Lineage-Specific Leukemias. Genetics, 2002, 160, 1051-1065.	2.9	8
22	Drosophila LSD1â€CoREST demethylase complex regulates DPP/TGFβ signaling during wing development. Genesis, 2013, 51, 16-31.	1.6	6
23	The Drosophila MLR COMPASS-like complex regulates bantam miRNA expression differentially in the context of cell fate. Developmental Biology, 2020, 468, 41-53.	2.0	2