

Zhiyuan Chen

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5628935/publications.pdf>

Version: 2024-02-01

16
papers

1,146
citations

643344

15
h-index

1051228

16
g-index

18
all docs

18
docs citations

18
times ranked

1437
citing authors

#	ARTICLE	IF	CITATIONS
1	Cell type-specific mechanism of Setd1a heterozygosity in schizophrenia pathogenesis. <i>Science Advances</i> , 2022, 8, eabm1077.	4.7	16
2	Distinct dynamics and functions of H2AK119ub1 and H3K27me3 in mouse preimplantation embryos. <i>Nature Genetics</i> , 2021, 53, 551-563.	9.4	83
3	DPPA2 and DPPA4 are dispensable for mouse zygotic genome activation and pre-implantation development. <i>Development (Cambridge)</i> , 2021, 148, .	1.2	17
4	The chromatin remodeler Snf2h is essential for oocyte meiotic cell cycle progression. <i>Genes and Development</i> , 2020, 34, 166-178.	2.7	21
5	Role of Mammalian DNA Methyltransferases in Development. <i>Annual Review of Biochemistry</i> , 2020, 89, 135-158.	5.0	182
6	Maternal H3K27me3-dependent autosomal and X chromosome imprinting. <i>Nature Reviews Genetics</i> , 2020, 21, 555-571.	7.7	53
7	Loss of DUX causes minor defects in zygotic genome activation and is compatible with mouse development. <i>Nature Genetics</i> , 2019, 51, 947-951.	9.4	138
8	Maternal-biased H3K27me3 correlates with paternal-specific gene expression in the human morula. <i>Genes and Development</i> , 2019, 33, 382-387.	2.7	47
9	Allelic H3K27me3 to allelic DNA methylation switch maintains noncanonical imprinting in extraembryonic cells. <i>Science Advances</i> , 2019, 5, eaay7246.	4.7	83
10	Maternal <i>Eed</i> knockout causes loss of H3K27me3 imprinting and random X inactivation in the extraembryonic cells. <i>Genes and Development</i> , 2018, 32, 1525-1536.	2.7	93
11	Colony-stimulating factor 2 acts from days 5 to 7 of development to modify programming of the bovine conceptus at day 86 of gestation. <i>Biology of Reproduction</i> , 2017, 96, 743-757.	1.2	30
12	Global misregulation of genes largely uncoupled to DNA methylome epimutations characterizes a congenital overgrowth syndrome. <i>Scientific Reports</i> , 2017, 7, 12667.	1.6	30
13	Global assessment of imprinted gene expression in the bovine conceptus by next generation sequencing. <i>Epigenetics</i> , 2016, 11, 501-516.	1.3	65
14	Characterization of global loss of imprinting in fetal overgrowth syndrome induced by assisted reproduction. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 4618-4623.	3.3	114
15	Large offspring syndrome. <i>Epigenetics</i> , 2013, 8, 591-601.	1.3	125
16	Expression of KCNQ1OT1, CDKN1C, H19, and PLAGL1 and the methylation patterns at the KvDMR1 and H19/IGF2 imprinting control regions is conserved between human and bovine. <i>Journal of Biomedical Science</i> , 2012, 19, 95.	2.6	48