## Klio Maratou

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/507731/publications.pdf

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567281 713466 1,187 21 15 21 h-index citations g-index papers 21 21 21 2222 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Two novel proteins recruited by synaptonemal complex protein 1 (SYCP1) are at the centre of meiosis. Journal of Cell Science, 2005, 118, 2755-2762.	2.0	190
2	Dazl binds in vivo to specific transcripts and can regulate the pre-meiotic translation of Mvh in germ cells. Human Molecular Genetics, 2005, 14, 3899-3909.	2.9	158
3	Genome Sequencing Reveals Loci under Artificial Selection that Underlie Disease Phenotypes in the Laboratory Rat. Cell, 2013, 154, 691-703.	28.9	154
4	Mouse MAELSTROM: the link between meiotic silencing of unsynapsed chromatin and microRNA pathway?. Human Molecular Genetics, 2006, 15, 2324-2334.	2.9	131
5	The zinc transporter ZIP12 regulates the pulmonary vascular response to chronic hypoxia. Nature, 2015, 524, 356-360.	27.8	113
6	<i>In silico</i> and <i>in vitro</i> evaluation of exonic and intronic off-target effects form a critical element of therapeutic ASO gapmer optimization. Nucleic Acids Research, 2015, 43, 8638-8650.	14.5	91
7	Comparison of dorsal root ganglion gene expression in rat models of traumatic and HIVâ€associated neuropathic pain. European Journal of Pain, 2009, 13, 387-398.	2.8	83
8	Expression profiling of the developing testis in wildâ€ŧype and <i>Dazl</i> knockout mice. Molecular Reproduction and Development, 2004, 67, 26-54.	2.0	45
9	Strategies for InÂVivo Screening and Mitigation of Hepatotoxicity Associated with Antisense Drugs. Molecular Therapy - Nucleic Acids, 2017, 8, 383-394.	5.1	37
10	Triple-target microarray experiments: a novel experimental strategy. BMC Genomics, 2004, 5, 13.	2.8	26
11	Heterotrisomy, a significant contributing factor to ventricular septal defect associated with Down syndrome?. Human Genetics, 2000, 107, 476-482.	3.8	25
12	Alternative Splicing and Transcriptome Profiling of Experimental Autoimmune Encephalomyelitis Using Genome-Wide Exon Arrays. PLoS ONE, 2009, 4, e7773.	2.5	20
13	Angiotensin-converting Enzyme Is a Modifier of Hypertensive End Organ Damage. Journal of Biological Chemistry, 2009, 284, 15564-15572.	3.4	20
14	Genetic Analysis of the Cardiac Methylome at Single Nucleotide Resolution in a Model of Human Cardiovascular Disease. PLoS Genetics, 2014, 10, e1004813.	3.5	19
15	MiMiR: an integrated platform for microarray data sharing, mining and analysis. BMC Bioinformatics, 2008, 9, 379.	2.6	16
16	Natural Polymorphisms in Tap2 Influence Negative Selection and CD4â^¶CD8 Lineage Commitment in the Rat. PLoS Genetics, 2014, 10, e1004151.	3.5	16
17	Isolation and high-throughput sequencing of two closely linked epistatic hypertension susceptibility loci with a panel of bicongenic strains. Physiological Genomics, 2013, 45, 729-736.	2.3	15
18	Variation in alphoid DNA size and trisomyÂ21: a possible cause of nondisjunction. Human Genetics, 2000, 106, 525-530.	3.8	11

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#	Article	IF	CITATION
19	Novel Methodology for the Detection of Chromosome 21-Specific $\hat{l}$ ±-Satellite DNA Sequences. Genomics, 1999, 57, 429-432.	2.9	8
20	Multiplexed DNA Methylation Analysis of Target Regions Using Microfluidics (Fluidigm). Methods in Molecular Biology, 2018, 1708, 349-363.	0.9	5
21	Variation in alphoid DNA size and trisomy 21: a possible cause of nondisjunction. Human Genetics, 2000, 106, 525-530.	3.8	4