

Matthew Jensen

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4863003/publications.pdf>

Version: 2024-02-01

23
papers

485
citations

933447

10
h-index

996975

15
g-index

29
all docs

29
docs citations

29
times ranked

1162
citing authors

#	ARTICLE	IF	CITATIONS
1	Rare variants in the genetic background modulate cognitive and developmental phenotypes in individuals carrying disease-associated variants. <i>Genetics in Medicine</i> , 2019, 21, 816-825.	2.4	127
2	Pervasive genetic interactions modulate neurodevelopmental defects of the autism-associated 16p11.2 deletion in <i>Drosophila melanogaster</i> . <i>Nature Communications</i> , 2018, 9, 2548.	12.8	56
3	Novel metrics to measure coverage in whole exome sequencing datasets reveal local and global non-uniformity. <i>Scientific Reports</i> , 2017, 7, 885.	3.3	43
4	A machine-learning approach for accurate detection of copy number variants from exome sequencing. <i>Genome Research</i> , 2019, 29, 1134-1143.	5.5	41
5	An interaction-based model for neuropsychiatric features of copy-number variants. <i>PLoS Genetics</i> , 2019, 15, e1007879.	3.5	39
6	Dissecting the genetic basis of comorbid epilepsy phenotypes in neurodevelopmental disorders. <i>Genome Medicine</i> , 2019, 11, 65.	8.2	35
7	Mapping a shared genetic basis for neurodevelopmental disorders. <i>Genome Medicine</i> , 2017, 9, 109.	8.2	31
8	NCBP2 modulates neurodevelopmental defects of the 3q29 deletion in <i>Drosophila</i> and <i>Xenopus laevis</i> models. <i>PLoS Genetics</i> , 2020, 16, e1008590.	3.5	30
9	A higher rare CNV burden in the genetic background potentially contributes to intellectual disability phenotypes in 22q11.2 deletion syndrome. <i>European Journal of Medical Genetics</i> , 2018, 61, 209-212.	1.3	17
10	Transcriptome Analyses of Heart and Liver Reveal Novel Pathways for Regulating Songbird Migration. <i>Scientific Reports</i> , 2019, 9, 6058.	3.3	16
11	Gene discoveries in autism are biased towards comorbidity with intellectual disability. <i>Journal of Medical Genetics</i> , 2020, 57, 647-652.	3.2	12
12	Functional assessment of the "two-hit" model for neurodevelopmental defects in <i>Drosophila</i> and <i>X. laevis</i> . <i>PLoS Genetics</i> , 2021, 17, e1009112.	3.5	12
13	<i>Drosophila</i> models of pathogenic copy-number variant genes show global and non-neuronal defects during development. <i>PLoS Genetics</i> , 2020, 16, e1008792.	3.5	9
14	The chromatin remodeler ISWI acts during <i>Drosophila</i> development to regulate adult sleep. <i>Science Advances</i> , 2021, 7, .	10.3	9
15	Combinatorial patterns of gene expression changes contribute to variable expressivity of the developmental delay-associated 16p12.1 deletion. <i>Genome Medicine</i> , 2021, 13, 163.	8.2	5
16	Title is missing!. , 2020, 16, e1008792.		0
17	Title is missing!. , 2020, 16, e1008792.		0
18	Title is missing!. , 2020, 16, e1008792.		0

#	ARTICLE	IF	CITATIONS
19	Title is missing!. , 2020, 16, e1008792.		0
20	Title is missing!. , 2020, 16, e1008590.		0
21	Title is missing!.. , 2020, 16, e1008590.		0
22	Title is missing!. , 2020, 16, e1008590.		0
23	Title is missing!.. , 2020, 16, e1008590.		0