Xiangyu

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/8193722/publications.pdf

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12	152	1163117 8	1199594 12
papers	citations	h-index	g-index

12	12	12	219
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Interaction of PDK1 with Phosphoinositides Is Essential for Neuronal Differentiation but Dispensable for Neuronal Survival. Molecular and Cellular Biology, 2013, 33, 1027-1040.	2.3	38
2	Fine-tuning the intensity of the PKB/Akt signal enables diverse physiological responses. Cell Cycle, 2014, 13, 3164-3168.	2.6	20
3	Bi-Allelic Mutations in NUP205 and NUP210 Are Associated With Abnormal Cardiac Left-Right Patterning. Circulation Genomic and Precision Medicine, 2019, 12, e002492.	3.6	14
4	Acox2 is a regulator of lysine crotonylation that mediates hepatic metabolic homeostasis in mice. Cell Death and Disease, 2022, 13, 279.	6.3	12
5	Rare mutations in apoptosis related genes APAF1, CASP9, and CASP3 contribute to human neural tube defects. Cell Death and Disease, 2018, 9, 43.	6.3	11
6	Two sides of NNMT in alcoholic and non-alcoholic fatty liver development. Journal of Hepatology, 2021, 74, 1250-1253.	3.7	11
7	Natural killer cells: functional differences in recurrent spontaneous abortionâ€. Biology of Reproduction, 2020, 102, 524-531.	2.7	10
8	Biallelic loss of function NEK3 mutations deacetylate Î±-tubulin and downregulate NUP205 that predispose individuals to cilia-related abnormal cardiac leftâ€“right patterning. Cell Death and Disease, 2020, 11, 1005.	6.3	10
9	Combined effects of FH (E404D) and ACOX2 (R409H) cause metabolic defects in primary cardiac malignant tumor. Cell Death Discovery, 2018, 4, 18.	4.7	9
10	ACOX2 deficiency in primary malignant cardiac tumors. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E3590-E3591.	7.1	8
11	Biallelic DNAH9 mutations are identified in Chinese patients with defective leftâ€“right patterning and cilia-related complex congenital heart disease. Human Genetics, 2022, 141, 1339-1353.	3.8	7
12	A hypoxia-inducible factor 1Î± null splice variant lacking exon 10. Cell Death and Disease, 2017, 8, e2873-e2873.	6.3	2