Ruo-Min Di

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/9821567/publications.pdf

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8 papers	155 citations	1307594 7 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	245 citing authors

#	Article	IF	Citations
1	GATA6 loss-of-function mutation contributes to congenital bicuspid aortic valve. Gene, 2018, 663, 115-120.	2.2	36
2	A SHOX2 loss-of-function mutation underlying familial atrial fibrillation. International Journal of Medical Sciences, 2018, 15, 1564-1572.	2.5	33
3	ISL1 loss-of-function mutation contributes to congenital heart defects. Heart and Vessels, 2019, 34, 658-668.	1.2	21
4	Identification and functional characterization of KLF5 as a novel disease gene responsible for familial dilated cardiomyopathy. European Journal of Medical Genetics, 2020, 63, 103827.	1.3	17
5	HAND2 loss-of-function mutation causes familial dilated cardiomyopathy. European Journal of Medical Genetics, 2019, 62, 103540.	1.3	16
6	Identification and Functional Characterization of an ISL1 Mutation Predisposing to Dilated Cardiomyopathy. Journal of Cardiovascular Translational Research, 2019, 12, 257-267.	2.4	14
7	NR2F2 lossâ€ofâ€function mutation is responsible for congenital bicuspid aortic valve. International Journal of Molecular Medicine, 2019, 43, 1839-1846.	4.0	11
8	Atrial Arrhythmias in Patients with Severe COVID-19. Cardiology Research and Practice, 2021, 2021, 1-7.	1.1	7