

# Ruo-Min Di

## List of Publications by Year in descending order

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8  
papers

155  
citations

1307594

7  
h-index

1588992

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g-index

8  
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8  
docs citations

8  
times ranked

245  
citing authors

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