

Xue Zhang

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6366392/publications.pdf>

Version: 2024-02-01

8
papers

67
citations

1684188
5
h-index

1720034
7
g-index

10
all docs

10
docs citations

10
times ranked

139
citing authors

#	ARTICLE	IF	CITATIONS
1	A comparison of bronchial blocker under artificial pneumothorax and double-lumen endobronchial tube for lung isolation in thoracoscopic enucleation of oesophageal leiomyoma. <i>Journal of Cardiothoracic Surgery</i> , 2021, 16, 322.	1.1	3
2	Homozygous missense variant in the TTN gene causing autosomal recessive limb-girdle muscular dystrophy type 10. <i>BMC Medical Genetics</i> , 2019, 20, 166.	2.1	12
3	Whole exome sequencing identified a novel truncation mutation in the NHS gene associated with Nance-Horan syndrome. <i>BMC Medical Genetics</i> , 2019, 20, 14.	2.1	12
4	Review of neuromyelitis optica spectrum disorder with pain-depression comorbidity. <i>Chinese Medical Sciences Journal</i> , 2019, 36, 1.	0.4	0
5	Mutation screening of NEK1 in Chinese ALS patients. <i>Neurobiology of Aging</i> , 2018, 71, 267.e1-267.e4.	3.1	13
6	ç¥žç»-çS'æ%«æœ-ç-...äºâ¿«éÿè¡€æ"â¼1âŠ»â¼¾æEææâ¿æœ-â¼â¼è¡€éâ¼...³ç³». <i>Chinese Medical Sciences Journal</i> , 2017, 32, 69-		
7	Identification of a novel loss-of-function C9orf72 splice site mutation in a patient with amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2016, 47, 219.e1-219.e5.	3.1	17
8	A Case of Novel Lamin A/C Mutation Manifesting as Atypical Progeroid Syndrome and Cardiomyopathy. <i>Canadian Journal of Cardiology</i> , 2016, 32, 1166.e29-1166.e31.	1.7	7