Kevin Whitehead

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11076946/publications.pdf

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9	950	1306789 7	1588620 8
papers	citations	h-index	g-index

9	9	9	1887
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Intracranial aneurysm and coarctation of the aorta: prevalence in the current era. Cardiology in the Young, 2021, 31, 229-232.	0.4	8
2	Cavernous Malformation Hemorrhagic Presentation at Diagnosis Associated with Low 25-Hydroxy-Vitamin D Level. Cerebrovascular Diseases, 2020, 49, 216-222.	0.8	9
3	CuraÃ§ao diagnostic criteria for hereditary hemorrhagic telangiectasia is highly predictive of a pathogenic variant in ENG or ACVRL1 (HHT1 and HHT2). Genetics in Medicine, 2020, 22, 1201-1205.	1.1	37
4	Phenotype of CM-AVM2 caused by variants in EPHB4: how much overlap with hereditary hemorrhagic telangiectasia (HHT)?. Genetics in Medicine, 2019, 21, 2007-2014.	1.1	38
5	Lateâ€inâ€Life Treadmillâ€Training Ameliorates the Decline in Cardiac Autophagy Associated with Aging in Mice. FASEB Journal, 2019, 33, 693.4.	0.2	0
6	Vitamin D levels are associated with epistaxis severity and bleeding duration in hereditary hemorrhagic telangiectasia. Biomarkers in Medicine, 2018, 12, 365-371.	0.6	3
7	Synopsis of Guidelines for the Clinical Management of Cerebral Cavernous Malformations: Consensus Recommendations Based on Systematic Literature Review by the Angioma Alliance Scientific Advisory Board Clinical Experts Panel. Neurosurgery, 2017, 80, 665-680.	0.6	334
8	Hereditary hemorrhagic telangiectasia: genetics and molecular diagnostics in a new era. Frontiers in Genetics, 2015, 6, 1.	1.1	489
9	Lack of CCM1 induces hypersprouting and impairs response to flow. Human Molecular Genetics, 2014, 23, 6223-6234.	1.4	32