Mostafa Hotait

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/4466430/publications.pdf

Version: 2024-02-01

14	171	1684188 5	1281871 11
papers	citations	h-index	g-index

16	16	16	247
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Swine Atrioventricular Node Ablation Using Stereotactic Radiosurgery: Methods and In Vivo Feasibility Investigation for Catheterâ€Free Ablation of Cardiac Arrhythmias. Journal of the American Heart Association, 2017, 6, .	3.7	39
2	Utility of the Exercise Electrocardiogram Testing in Sudden Cardiac Death Risk Stratification. , 2014, 19, 311-318.		32
3	Genetics of Sudden Cardiac Death. Current Cardiology Reports, 2015, 17, 606.	2.9	30
4	Efficacy and tolerability of treatment with lacosamide in children: Postmarketing experience from the Middle East. Seizure: the Journal of the British Epilepsy Association, 2020, 79, 75-79.	2.0	14
5	FARS2 Mutations: More Than Two Phenotypes? A Case Report. Frontiers in Genetics, 2020, 11, 787.	2.3	13
6	Brugada Syndrome. Cardiac Electrophysiology Clinics, 2016, 8, 239-245.	1.7	8
7	The Muscle-Bound Heart. Cardiac Electrophysiology Clinics, 2016, 8, 223-231.	1.7	6
8	The Mutation P.T613a in the Pore Helix of the Kv11.1 Potassium Channel is Associated with Long QT Syndrome. PACE - Pacing and Clinical Electrophysiology, 2015, 38, 1304-1309.	1.2	5
9	A novel possible familial cause of epilepsy of infancy with migrating focal seizures related to SZT2 gene variant. Epilepsia Open, 2021, 6, 73-78.	2.4	5
10	Patients' and family members' views on pacemaker reuse: An international survey. Journal of Cardiovascular Electrophysiology, 2022, , .	1.7	5
11	Non-familial cardiomyopathies in Lebanon: exome sequencing results for five idiopathic cases. BMC Medical Genomics, 2019, 12, 33.	1.5	4
12	Homozygous factor V Leiden mutation in type IV Ehlers-Danlos patient. World Journal of Clinical Cases, 2014, 2, 75.	0.8	4
13	Iron Overload Leading to Torsades de Pointes in Î²-Thalassemia and Long QT Syndrome. Cardiac Electrophysiology Clinics, 2016, 8, 247-256.	1.7	3
14	Case Report: Distinctive EEG Patterns in SCARB-2 Related Progressive Myoclonus Epilepsy. Frontiers in Genetics, 2020, 11, 581253.	2.3	1