

Mostafa Hotait

List of Publications by Year in descending order

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

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14
papers

171
citations

1684188

5
h-index

1281871

11
g-index

16
all docs

16
docs citations

16
times ranked

247
citing authors

#	ARTICLE	IF	CITATIONS
1	Patients' and family members' views on pacemaker reuse: An international survey. Journal of Cardiovascular Electrophysiology, 2022, , .	1.7	5
2	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
3	Case Report: Distinctive EEG Patterns in SCARB-2 Related Progressive Myoclonus Epilepsy. Frontiers in Genetics, 2020, 11, 581253.	2.3	1
4	FARS2 Mutations: More Than Two Phenotypes? A Case Report. Frontiers in Genetics, 2020, 11, 787.	2.3	13
5	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
6	Non-familial cardiomyopathies in Lebanon: exome sequencing results for five idiopathic cases. BMC Medical Genomics, 2019, 12, 33.	1.5	4
7	Swine Atrioventricular Node Ablation Using Stereotactic Radiosurgery: Methods and In Vivo Feasibility Investigation for Catheter-Free Ablation of Cardiac Arrhythmias. Journal of the American Heart Association, 2017, 6, .	3.7	39
8	Iron Overload Leading to Torsades de Pointes in β^2 -Thalassemia and Long QT Syndrome. Cardiac Electrophysiology Clinics, 2016, 8, 247-256.	1.7	3
9	Brugada Syndrome. Cardiac Electrophysiology Clinics, 2016, 8, 239-245.	1.7	8
10	The Muscle-Bound Heart. Cardiac Electrophysiology Clinics, 2016, 8, 223-231.	1.7	6
11	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
12	Genetics of Sudden Cardiac Death. Current Cardiology Reports, 2015, 17, 606.	2.9	30
13	Utility of the Exercise Electrocardiogram Testing in Sudden Cardiac Death Risk Stratification. , 2014, 19, 311-318.		32
14	Homozygous factor V Leiden mutation in type IV Ehlers-Danlos patient. World Journal of Clinical Cases, 2014, 2, 75.	0.8	4