

Maen D Abou Ziki

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/151279/publications.pdf>

Version: 2024-02-01

19
papers

284
citations

1163117

8
h-index

1058476

14
g-index

23
all docs

23
docs citations

23
times ranked

630
citing authors

#	ARTICLE	IF	CITATIONS
1	Metabolic syndrome. <i>Current Opinion in Lipidology</i> , 2016, 27, 162-171.	2.7	58
2	Application of Whole Exome Sequencing in the Clinical Diagnosis and Management of Inherited Cardiovascular Diseases in Adults. <i>Circulation: Cardiovascular Genetics</i> , 2017, 10, .	5.1	55
3	Pulmonary Embolism and Atrial Fibrillation: Two Sides of the Same Coin? A Systematic Review. <i>Seminars in Thrombosis and Hemostasis</i> , 2017, 43, 849-863.	2.7	45
4	The interplay of canonical and noncanonical Wnt signaling in metabolic syndrome. <i>Nutrition Research</i> , 2019, 70, 18-25.	2.9	26
5	Wnt signaling, a novel pathway regulating blood pressure? State of the art review. <i>Atherosclerosis</i> , 2017, 262, 171-178.	0.8	24
6	Point of Care Exome Sequencing Reveals Allelic and Phenotypic Heterogeneity Underlying Mendelian disease in Qatar. <i>Human Molecular Genetics</i> , 2019, 28, 3970-3981.	2.9	16
7	Prevalence of the Apolipoprotein E Arg145Cys Dyslipidemia At-Risk Polymorphism in African-Derived Populations. <i>American Journal of Cardiology</i> , 2014, 113, 302-308.	1.6	13
8	Nurse-led theory-based educational intervention improves glycemic and metabolic parameters in South Asian patients with type II diabetes: a randomized controlled trial. <i>Diabetology International</i> , 2017, 8, 95-103.	1.4	11
9	Epistatic interaction of PDE4DIP and DES mutations in familial atrial fibrillation with slow conduction. <i>Human Mutation</i> , 2021, 42, 1279-1293.	2.5	10
10	Deleterious protein-altering mutations in the <i>SCN10A</i> voltage-gated sodium channel gene are associated with prolonged QT. <i>Clinical Genetics</i> , 2018, 93, 741-751.	2.0	9
11	The Value of the History and Physical Examination "Sailing Through Medicine With Modern Tools. <i>JAMA Internal Medicine</i> , 2015, 175, 1901.	5.1	8
12	Dyrk1b promotes autophagy during skeletal muscle differentiation by upregulating 4e-bp1. <i>Cellular Signalling</i> , 2022, 90, 110186.	3.6	4
13	Rare mutation in the SLC26A3 transporter causes life-long diarrhoea with metabolic alkalosis. <i>BMJ Case Reports</i> , 2015, 2015, bcr2014206849-bcr2014206849.	0.5	3
14	Tachyarrhythmia Onset Captured on Telemetry Deciphers the Diagnosis. <i>JAMA Internal Medicine</i> , 2017, 177, 1673.	5.1	2
15	The Yin-Yang of Atrial Fibrillation and Pulmonary Embolism. <i>Chest</i> , 2016, 150, 1189A.	0.8	0
16	Beware of Limb Lead Reversal "Reply. <i>JAMA Internal Medicine</i> , 2018, 178, 435.	5.1	0
17	Genetics of Dyslipidemias and the Role of the ApoE Arg145Cys Mutation in African-derived Populations. <i>Qscience Proceedings</i> , 2012, , .	0.0	0
18	Collaborative leadership: organisational structure and institutional investment to multiply innovative educational efforts among trainees. <i>BMJ Leader</i> , 0, , leader-2020-000233.	1.5	0

#	ARTICLE	IF	CITATIONS
19	Abstract 17195: Hydroxychloroquine Associated Electrocardiographic Changes in Patients With COVID-19 Extend Beyond QTc Prolongation and Are Accentuated in Those With Myocardial Injury. Circulation, 2020, 142, .	1.6	0