Antoine B Abchee

List of Publications by Year
in descending order

13 papers	293 citations	1307594 7 h-index	1281871 11 g-index
13 all docs	13 docs citations	13 times ranked	521 citing authors

#	Article	IF	Citations
1	Founder Mutation in N Terminus of Cardiac Troponin I Causes Malignant Hypertrophic Cardiomyopathy. Circulation Genomic and Precision Medicine, 2020, 13, 444-452.	3.6	7
2	Artificial neural network-based model enhances risk stratification and reduces non-invasive cardiac stress imaging compared to Diamondâ€“Forrester and Morise risk assessment models: A prospective study. Journal of Nuclear Cardiology, 2018, 25, 1601-1609.	2.1	7
3	Type II diabetes mellitus and hyperhomocysteinemia: a complex interaction. Diabetology and Metabolic Syndrome, 2017, 9, 19.	2.7	20
4	Association of waterpipe smoking with myocardial infarction and determinants of metabolic syndrome among catheterized patients. Inhalation Toxicology, 2017, 29, 429-434.	1.6	12
5	Isolated Left Ventricular Diastolic Collapse Due to Extraâ€Thoracic Compression. Echocardiography, 2015, 32, 1314-1317.	0.9	0
6	Association of coronary artery disease and chronic kidney disease in Lebanese population. International Journal of Clinical and Experimental Medicine, 2015, 8, 15866-77.	1.3	1
7	Multivariate epidemiologic analysis of type 2 diabetes mellitus risks in the Lebanese population. Diabetology and Metabolic Syndrome, 2014, 6, 89.	2.7	17
8	Genetic and environmental influences on total plasma homocysteine and its role in coronary artery disease risk. Atherosclerosis, 2012, 222, 180-186.	0.8	27
9	Genome-Wide Association Study in a Lebanese Cohort Confirms PHACTR1 as a Major Determinant of Coronary Artery Stenosis. PLoS ONE, 2012, 7, e38663.	2.5	52
10	Molecular genetics of familial hypertrophic cardiomyopathy. Progress in Pediatric Cardiology, 1996, 6, 63-70.	0.4	0
11	Molecular and Clinical Aspects of Inherited Cardiomyopathies. Annals of Medicine, 1995, 27, 311-317.	3.8	22
12	901â€“88 Rapid Genetic Screen for Common Î²-Myosin Heavy Chain Mutations Causing Familial Hypertrophic Cardiomyopathy. Journal of the American College of Cardiology, 1995, 25, 26A.	2.8	5
13	Localization of a Gene Responsible for Familial Dilated Cardiomyopathy to Chromosome 1q32. Circulation, 1995, 92, 3387-3389.	1.6	123