

Meena Fatah

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1481934/publications.pdf>

Version: 2024-02-01

10
papers

409
citations

1163117

8
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

707
citing authors

#	ARTICLE	IF	CITATIONS
1	Evaluating the 12-Lead Electrocardiogram for Diagnosing ARVC in Young Populations: Implications for Preparticipation Screening of Athletes. <i>CJC Open</i> , 2021, 3, 498-503.	1.5	2
2	An autoantibody profile detects Brugada syndrome and identifies abnormally expressed myocardial proteins. <i>European Heart Journal</i> , 2020, 41, 2878-2890.	2.2	40
3	Initially unexplained cardiac arrest in children and adolescents: A national experience from the Canadian Pediatric Heart Rhythm Network. <i>Heart Rhythm</i> , 2020, 17, 975-981.	0.7	21
4	Calmodulin mutations and life-threatening cardiac arrhythmias: insights from the International Calmodulinopathy Registry. <i>European Heart Journal</i> , 2019, 40, 2964-2975.	2.2	116
5	Association of Echocardiographic Parameters of Right Ventricular Remodeling and Myocardial Performance With Modified Task Force Criteria in Adolescents With Arrhythmogenic Right Ventricular Cardiomyopathy. <i>Circulation: Cardiovascular Imaging</i> , 2019, 12, e007693.	2.6	30
6	Ankyrin-B dysfunction predisposes to arrhythmogenic cardiomyopathy and is amenable to therapy. <i>Journal of Clinical Investigation</i> , 2019, 129, 3171-3184.	8.2	42
7	An autoantibody identifies arrhythmogenic right ventricular cardiomyopathy and participates in its pathogenesis. <i>European Heart Journal</i> , 2018, 39, 3932-3944.	2.2	114
8	Clinical utility of endomyocardial biopsies in the diagnosis of arrhythmogenic right ventricular cardiomyopathy in children. <i>Pediatric Research</i> , 2018, 84, 552-557.	2.3	4
9	Left Ventricular Function in Children and Adolescents With Arrhythmogenic Right Ventricular Cardiomyopathy. <i>American Journal of Cardiology</i> , 2017, 119, 778-784.	1.6	21
10	Novel CALM3 mutations in pediatric long QT syndrome patients support a CALM3 -specific calmodulinopathy. <i>HeartRhythm Case Reports</i> , 2016, 2, 250-254.	0.4	19