

Youmna Ghaleb

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7725286/publications.pdf>

Version: 2024-02-01

10
papers

218
citations

1478505

6
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

486
citing authors

#	ARTICLE	IF	CITATIONS
1	Living the PCSK9 Adventure: from the Identification of a New Gene in Familial Hypercholesterolemia Towards a Potential New Class of Anticholesterol Drugs. <i>Current Atherosclerosis Reports</i> , 2014, 16, 439.	4.8	87
2	PCSK9 Mutations in Familial Hypercholesterolemia: from a Groundbreaking Discovery to Anti-PCSK9 Therapies. <i>Current Atherosclerosis Reports</i> , 2017, 19, 49.	4.8	31
3	New Sequencing technologies help revealing unexpected mutations in Autosomal Dominant Hypercholesterolemia. <i>Scientific Reports</i> , 2018, 8, 1943.	3.3	25
4	Proprotein convertase subtilisin / kexin 9 (PCSK9) inhibitors and the future of dyslipidemia therapy: an updated patent review (2011-2015). <i>Expert Opinion on Therapeutic Patents</i> , 2016, 26, 1377-1392.	5.0	23
5	Usefulness of the genetic risk score to identify phenocopies in families with familial hypercholesterolemia?. <i>European Journal of Human Genetics</i> , 2018, 26, 570-578.	2.8	22
6	Plasma proproteinâ€convertaseâ€subtilisin/kexin type 9 (PCSK9) and cardiovascular events in type 2 diabetes. <i>Diabetes, Obesity and Metabolism</i> , 2018, 20, 943-953.	4.4	17
7	Identification of the first Tangier disease patient in Lebanon carrying a new pathogenic variant in ABCA1. <i>Journal of Clinical Lipidology</i> , 2018, 12, 1374-1382.	1.5	6
8	Identification of a Variant in APOB Gene as a Major Cause of Hypobetalipoproteinemia in Lebanese Families. <i>Metabolites</i> , 2021, 11, 564.	2.9	5
9	Whole Exome/Genome Sequencing Joint Analysis of a Family with Oligogenic Familial Hypercholesterolemia. <i>Metabolites</i> , 2022, 12, 262.	2.9	1
10	Circulating PCSK9 Linked to Dyslipidemia in Lebanese Schoolchildren. <i>Metabolites</i> , 2022, 12, 504.	2.9	1