

Sandy Elbitar

List of Publications by Year in descending order

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Version: 2024-02-01

13
papers

201
citations

1478505

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1372567

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14
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14
docs citations

14
times ranked

478
citing authors

#	ARTICLE	IF	CITATIONS
1	Whole Exome/Genome Sequencing Joint Analysis of a Family with Oligogenic Familial Hypercholesterolemia. <i>Metabolites</i> , 2022, 12, 262.	2.9	1
2	Circulating PCSK9 Linked to Dyslipidemia in Lebanese Schoolchildren. <i>Metabolites</i> , 2022, 12, 504.	2.9	1
3	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
4	New Sequencing technologies help revealing unexpected mutations in Autosomal Dominant Hypercholesterolemia. <i>Scientific Reports</i> , 2018, 8, 1943.	3.3	25
5	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
6	High prevalence of ventricular repolarization abnormalities in people carrying TGFÎ²R2 mutations. <i>Scientific Reports</i> , 2018, 8, 13019.	3.3	4
7	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
8	PCSK9 Mutations in Familial Hypercholesterolemia: from a Groundbreaking Discovery to Anti-PCSK9 Therapies. <i>Current Atherosclerosis Reports</i> , 2017, 19, 49.	4.8	31
9	Usefulness of the genetic risk score to identify phenocopies in families with autosomal dominant hypercholesterolemia?. <i>Atherosclerosis</i> , 2017, 263, e83.	0.8	0
10	Plasma PCSK9 and cardiovascular events in type 2 diabetes. <i>Atherosclerosis</i> , 2017, 263, e81.	0.8	1
11	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
12	Identification of a new mutation in the N-terminal region of the apolipoprotein B gene in familial hypercholesterolemia. <i>Atherosclerosis</i> , 2016, 252, e34.	0.8	0
13	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