Sandy Elbitar

List of Publications by Year
in descending order

13	201	1478280 6	1372474 10
papers	citations	h-index	g-index

14	14	14	478
all docs	docs citations	times ranked	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. Current Atherosclerosis Reports, 2014, 16, 439.	2.0	87
2	PCSK9 Mutations in Familial Hypercholesterolemia: from a Groundbreaking Discovery to Anti-PCSK9 Therapies. Current Atherosclerosis Reports, 2017, 19, 49.	2.0	31
3	New Sequencing technologies help revealing unexpected mutations in Autosomal Dominant Hypercholesterolemia. Scientific Reports, 2018, 8, 1943.	1.6	25
4	Proprotein convertase subtilisin / kexin 9 (PCSK9) inhibitors and the future of dyslipidemia therapy: an updated patent review (2011-2015). Expert Opinion on Therapeutic Patents, 2016, 26, 1377-1392.	2.4	23
5	Plasma proproteinâ€convertaseâ€subtilisin/kexin type 9 (PCSK9) and cardiovascular events in type 2 diabetes. Diabetes, Obesity and Metabolism, 2018, 20, 943-953.	2.2	17
6	Identification of the first Tangier disease patient in Lebanon carrying a new pathogenic variant in ABCA1. Journal of Clinical Lipidology, 2018, 12, 1374-1382.	0.6	6
7	Identification of a Variant in APOB Gene as a Major Cause of Hypobetalipoproteinemia in Lebanese Families. Metabolites, 2021, 11, 564.	1.3	5
8	High prevalence of ventricular repolarization abnormalities in people carrying TGFÎ²R2 mutations. Scientific Reports, 2018, 8, 13019.	1.6	4
9	Plasma PCSK9 and cardiovascular events in type 2 diabetes. Atherosclerosis, 2017, 263, e81.	0.4	1
10	Whole Exome/Genome Sequencing Joint Analysis of a Family with Oligogenic Familial Hypercholesterolemia. Metabolites, 2022, 12, 262.	1.3	1
11	Circulating PCSK9 Linked to Dyslipidemia in Lebanese Schoolchildren. Metabolites, 2022, 12, 504.	1.3	1
12	Identification of a new mutation in the N-terminal region of the apolipoprotein B gene in familial hypercholesterolemia. Atherosclerosis, 2016, 252, e34.	0.4	0
13	Usefulness of the genetic risk score to identify phenocopies in families with autosomal dominant hypercholesterolemia?. Atherosclerosis, 2017, 263, e83.	0.4	0