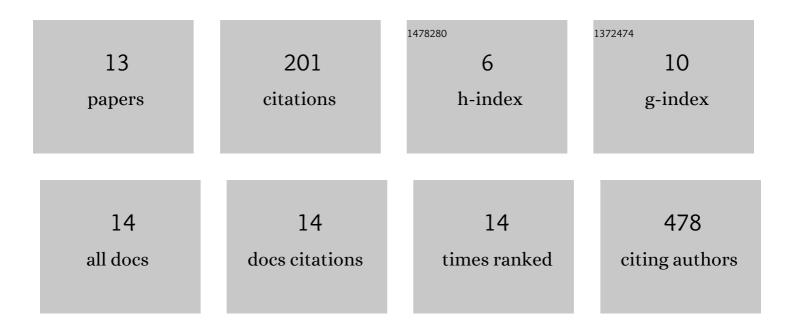
Sandy Elbitar

List of Publications by Year in descending order

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| # | 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 TGFβR2 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 |