Michael A Iacocca

List of Publications by Year
in descending order

14	869	759233 12	1058476 14
papers	citations	h-index	g-index

15	15	15	1199
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The Clinical Genome Resource (ClinGen) Familial Hypercholesterolemia Variant Curation Expert Panel consensus guidelines for LDLR variant classification. Genetics in Medicine, 2022, 24, 293-306.	2.4	53
2	ClinGen Variant Curation Interface: a variant classification platform for the application of evidence criteria from ACMG/AMP guidelines. Genome Medicine, 2022, 14, 6.	8.2	34
3	Improving reporting standards for polygenic scores in risk prediction studies. Nature, 2021, 591, 211-219.	27.8	265
4	Six yearsâ€™ experience with LipidSeq: clinical and research learnings from a hybrid, targeted sequencing panel for dyslipidemias. BMC Medical Genomics, 2020, 13, 23.	1.5	52
5	Genetic Determinants of Myocardial Infarction Risk in Familial Hypercholesterolemia. CJC Open, 2019, 1, 225-230.	1.5	10
6	Efficacy of Evolocumab in Monogenic vs Polygenic Hypercholesterolemia. CJC Open, 2019, 1, 115-118.	1.5	8
7	Progress in finding pathogenic DNA copy number variations in dyslipidemia. Current Opinion in Lipidology, 2019, 30, 63-70.	2.7	18
8	Severe hypertriglyceridemia is primarily polygenic. Journal of Clinical Lipidology, 2019, 13, 80-88.	1.5	136
9	Role of DNA copy number variation in dyslipidemias. Current Opinion in Lipidology, 2018, 29, 125-132.	2.7	33
10	ClinVar database of global familial hypercholesterolemiaâ€associated DNA variants. Human Mutation, 2018, 39, 1631-1640.	2.5	84
11	Whole-Gene Duplication of PCSK9 as a Novel Genetic Mechanism for Severe Familial Hypercholesterolemia. Canadian Journal of Cardiology, 2018, 34, 1316-1324.	1.7	34
12	Large-scale deletions of the ABCA1 gene in patients with hypoalphalipoproteinemia. Journal of Lipid Research, 2018, 59, 1529-1535.	4.2	22
13	Recent advances in genetic testing for familial hypercholesterolemia. Expert Review of Molecular Diagnostics, 2017, 17, 641-651.	3.1	53
14	Use of next-generation sequencing to detect LDLR gene copy number variation in familial hypercholesterolemia. Journal of Lipid Research, 2017, 58, 2202-2209.	4.2	65