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List of Publications by Year in descending order

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Version: 2024-02-01

14
papers

869
citations

758635

12
h-index

1058022

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15
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15
docs citations

15
times ranked

1199
citing authors

#	ARTICLE	IF	CITATIONS
1	Improving reporting standards for polygenic scores in risk prediction studies. <i>Nature</i> , 2021, 591, 211-219.	13.7	265
2	Severe hypertriglyceridemia is primarily polygenic. <i>Journal of Clinical Lipidology</i> , 2019, 13, 80-88.	0.6	136
3	ClinVar database of global familial hypercholesterolemia-associated DNA variants. <i>Human Mutation</i> , 2018, 39, 1631-1640.	1.1	84
4	Use of next-generation sequencing to detect LDLR gene copy number variation in familial hypercholesterolemia. <i>Journal of Lipid Research</i> , 2017, 58, 2202-2209.	2.0	65
5	Recent advances in genetic testing for familial hypercholesterolemia. <i>Expert Review of Molecular Diagnostics</i> , 2017, 17, 641-651.	1.5	53
6	The Clinical Genome Resource (ClinGen) Familial Hypercholesterolemia Variant Curation Expert Panel consensus guidelines for LDLR variant classification. <i>Genetics in Medicine</i> , 2022, 24, 293-306.	1.1	53
7	Six years'™ experience with LipidSeq: clinical and research learnings from a hybrid, targeted sequencing panel for dyslipidemias. <i>BMC Medical Genomics</i> , 2020, 13, 23.	0.7	52
8	Whole-Genome Duplication of PCSK9 as a Novel Genetic Mechanism for Severe Familial Hypercholesterolemia. <i>Canadian Journal of Cardiology</i> , 2018, 34, 1316-1324.	0.8	34
9	ClinGen Variant Curation Interface: a variant classification platform for the application of evidence criteria from ACMG/AMP guidelines. <i>Genome Medicine</i> , 2022, 14, 6.	3.6	34
10	Role of DNA copy number variation in dyslipidemias. <i>Current Opinion in Lipidology</i> , 2018, 29, 125-132.	1.2	33
11	Large-scale deletions of the ABCA1 gene in patients with hypoalphalipoproteinemia. <i>Journal of Lipid Research</i> , 2018, 59, 1529-1535.	2.0	22
12	Progress in finding pathogenic DNA copy number variations in dyslipidemia. <i>Current Opinion in Lipidology</i> , 2019, 30, 63-70.	1.2	18
13	Genetic Determinants of Myocardial Infarction Risk in Familial Hypercholesterolemia. <i>CJC Open</i> , 2019, 1, 225-230.	0.7	10
14	Efficacy of Evolocumab in Monogenic vs Polygenic Hypercholesterolemia. <i>CJC Open</i> , 2019, 1, 115-118.	0.7	8