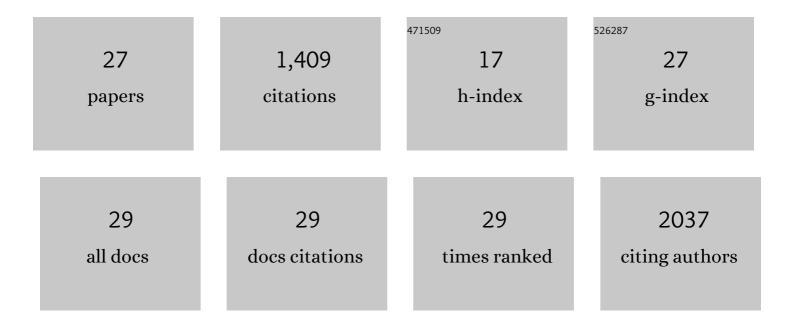
## Jacqueline S Dron

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3854020/publications.pdf

Version: 2024-02-01



#	Article	IF	CITATIONS
1	Improving reporting standards for polygenic scores in risk prediction studies. Nature, 2021, 591, 211-219.	27.8	265
2	Polygenic Versus Monogenic Causes of Hypercholesterolemia Ascertained Clinically. Arteriosclerosis, Thrombosis, and Vascular Biology, 2016, 36, 2439-2445.	2.4	174
3	Severe hypertriglyceridemia is primarily polygenic. Journal of Clinical Lipidology, 2019, 13, 80-88.	1.5	136
4	Genetics of Hypertriglyceridemia. Frontiers in Endocrinology, 2020, 11, 455.	3.5	100
5	Genetics of Triglycerides and the Risk of Atherosclerosis. Current Atherosclerosis Reports, 2017, 19, 31.	4.8	89
6	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
7	PCSK9: Regulation and Target for Drug Development for Dyslipidemia. Annual Review of Pharmacology and Toxicology, 2017, 57, 223-244.	9.4	58
8	Complexity of mechanisms among human proprotein convertase subtilisin–kexin type 9 variants. Current Opinion in Lipidology, 2017, 28, 161-169.	2.7	57
9	Targeted next generation sequencing as a tool for precision medicine. BMC Medical Genomics, 2019, 12, 81.	1.5	54
10	Six years' experience with LipidSeq: clinical and research learnings from a hybrid, targeted sequencing panel for dyslipidemias. BMC Medical Genomics, 2020, 13, 23.	1.5	52
11	Polygenic influences on dyslipidemias. Current Opinion in Lipidology, 2018, 29, 133-143.	2.7	51
12	Polygenic determinants in extremes of high-density lipoprotein cholesterol. Journal of Lipid Research, 2017, 58, 2162-2170.	4.2	49
13	The evolution of genetic-based risk scores for lipids and cardiovascular disease. Current Opinion in Lipidology, 2019, 30, 71-81.	2.7	49
14	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
15	The polygenic nature of mild-to-moderate hypertriglyceridemia. Journal of Clinical Lipidology, 2020, 14, 28-34.e2.	1.5	32
16	Large-scale deletions of the ABCA1 gene in patients with hypoalphalipoproteinemia. Journal of Lipid Research, 2018, 59, 1529-1535.	4.2	22
17	Combined hyperlipidemia is genetically similar to isolated hypertriglyceridemia. Journal of Clinical Lipidology, 2021, 15, 79-87.	1.5	20
18	Targeted sequencing reveals expanded genetic diversity of human transfer RNAs. RNA Biology, 2019, 16, 1574-1585	3.1	19

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#	Article	IF	CITATIONS
19	Progress in finding pathogenic DNA copy number variations in dyslipidemia. Current Opinion in Lipidology, 2019, 30, 63-70.	2.7	18
20	2019 George Lyman Duff Memorial Lecture. Arteriosclerosis, Thrombosis, and Vascular Biology, 2020, 40, 1970-1981.	2.4	16
21	Partial LPL deletions: rare copy-number variants contributing towards severe hypertriglyceridemia. Journal of Lipid Research, 2019, 60, 1953-1958.	4.2	12
22	Genetic Predictor to Identify Individuals With High Lipoprotein(a) Concentrations. Circulation Genomic and Precision Medicine, 2021, 14, e003182.	3.6	10
23	Human variant of scavenger receptor BI (R174C) exhibits impaired cholesterol transport functions. Journal of Lipid Research, 2021, 62, 100045.	4.2	8
24	Recent Advances in the Genetics of Atherothrombotic Disease and Its Determinants. Arteriosclerosis, Thrombosis, and Vascular Biology, 2017, 37, e158-e166.	2.4	7
25	Role of Common Genetic Variation in Lone Atrial Fibrillation. Circulation Genomic and Precision Medicine, 2021, 14, e003179.	3.6	5
26	Recent Highlights of ATVB. Arteriosclerosis, Thrombosis, and Vascular Biology, 2018, 38, e185-e197.	2.4	3
27	A tip of the CAP1 to cholesterol metabolism. European Heart Journal, 2019, 41, 253-254.	2.2	3