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#	Paper	IF	Citations
21	Cholesterol as an Endogenous ERR[Agonist: A New Perspective to Cancer Treatment. <i>Frontiers in Endocrinology</i> , <b>2018</b> , 9, 525	5.7	24
20	Familial combined hyperlipidemia: An overview of the underlying molecular mechanisms and therapeutic strategies. <i>IUBMB Life</i> , <b>2019</b> , 71, 1221-1229	4.7	15
19	PCSK9 inhibitor improves cardiac function and reduces infarct size in rats with ischaemia/reperfusion injury: Benefits beyond lipid-lowering effects. <i>Journal of Cellular and Molecular Medicine</i> , <b>2019</b> , 23, 7310-7319	5.6	19
18	Scavenger Receptors as Biomarkers and Therapeutic Targets in Cardiovascular Disease. <i>Cells</i> , <b>2020</b> , 9,	7.9	1
17	Series of Novel and Highly Potent Cyclic Peptide PCSK9 Inhibitors Derived from an mRNA Display Screen and Optimized via Structure-Based Design. <i>Journal of Medicinal Chemistry</i> , <b>2020</b> , 63, 13796-1382	8.3	15
16	Pharmacogenetics of Statin-Induced Myotoxicity. Frontiers in Genetics, 2020, 11, 575678	4.5	15
15	Novel Treatment Strategies for Secondary Prevention of Cardiovascular Disease: A Systematic Review of Cost-Effectiveness. <i>Pharmacoeconomics</i> , <b>2020</b> , 38, 1095-1113	4.4	11
14	Lipid-based gene delivery to macrophage mitochondria for atherosclerosis therapy. <i>Pharmacology Research and Perspectives</i> , <b>2020</b> , 8, e00584	3.1	9
13	Potent Lys Patch-Containing Stapled Peptides Targeting PCSK9. <i>Journal of Medicinal Chemistry</i> , <b>2021</b> , 64, 10834-10848	8.3	3
12	Pseurotin A as a novel suppressor of hormone dependent breast cancer progression and recurrence by inhibiting PCSK9 secretion and interaction with LDL receptor. <i>Pharmacological Research</i> , <b>2020</b> , 158, 104847	10.2	16
11	Non-alcoholic fatty liver disease: a metabolic burden promoting atherosclerosis. <i>Clinical Science</i> , <b>2020</b> , 134, 1775-1799	6.5	9
10	Assessing drug target suitability using TargetMine. F1000Research, 2019, 8, 233	3.6	2
9	Assessing drug target suitability using TargetMine. F1000Research, 2019, 8, 233	3.6	1
8	A Series of Novel, Highly Potent, and Orally Bioavailable Next-Generation Tricyclic Peptide PCSK9 Inhibitors. <i>Journal of Medicinal Chemistry</i> , <b>2021</b> , 64, 16770-16800	8.3	6
7	Involvement of LDL and ox-LDL in Cancer Development and Its Therapeutical Potential <i>Frontiers in Oncology</i> , <b>2022</b> , 12, 803473	5.3	6
6	Analysis of rare genetic variation underlying cardiometabolic diseases and traits among 200,000 individuals in the UK Biobank <i>Nature Genetics</i> , <b>2022</b> ,	36.3	4
5	Antitumor activity and molecular mechanism of proprotein convertase subtilisin/kexin type 9 (PCSK9) inhibition <i>Naunyn-Schmiedebergus Archives of Pharmacology</i> , <b>2022</b> , 1	3.4	O

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4	Adverse event profiles of PCSK9 inhibitors alirocumab and evolocumab: data mining of the FDA Adverse Event Reporting System. <i>British Journal of Clinical Pharmacology</i> ,	3.8	1
3	Effects of Loading-Dose Statins Combined with PCSK9 Inhibitor Pre-Treatment before Primary Percutaneous Coronary Intervention on the Short-Term Prognosis in Patients with ST-Segment Elevation Myocardial Infarction. <b>2022</b> ,		O
2	Interactions between PCSK9 and NLRP3 inflammasome signaling in atherosclerosis. 14,		O
1	Patient adherence to fully reimbursed proprotein convertase subtilisin/kexin type 9 inhibitor (PCSK9i) treatment.		O