PCSK9 genetic variants and risk of type 2 diabetes: a me

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Citation Report

#	Article	IF	Citations
1	Common and rare genetic variants and risk of CHD. Nature Reviews Cardiology, 2017, 14, 73-74.	13.7	8
2	Bococizumab for the treatment of hypercholesterolaemia. Expert Opinion on Biological Therapy, 2017, 17, 237-243.	3.1	20
3	PCSK9 inhibition and the global diabetes epidemic. Diabetologia, 2017, 60, 751-752.	6.3	0
4	Familial Hypercholesterolemia and Type 2 Diabetes in the Old Order Amish. Diabetes, 2017, 66, 2054-2058.	0.6	28
5	Alirocumab for the treatment of hypercholesterolaemia. Expert Review of Clinical Pharmacology, 2017, 10, 571-582.	3.1	9
6	Dyslipidaemia in type 2 diabetes mellitus. Current Opinion in Cardiology, 2017, 32, 422-429.	1.8	26
7	Physiological and therapeutic regulation of PCSK9 activity in cardiovascular disease. Basic Research in Cardiology, 2017, 112, 32.	5.9	66
8	Proprotein convertase subtilisin-kexin type 9 (PCSK9) inhibitors: Shaping the future after the further cardiovascular outcomes research with PCSK9 inhibition in subjects with elevated risk (FOURIER) trial. Metabolism: Clinical and Experimental, 2017, 74, 43-46.	3.4	19
9	PCSK9 monoclonal antibodies for the primary and secondary prevention of cardiovascular disease. The Cochrane Library, 2017, 4, CD011748.	2.8	93
10	Human genetics as a model for target validation: finding new therapies for diabetes. Diabetologia, 2017, 60, 960-970.	6.3	19
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18	Exome-wide association study of plasma lipids in >300,000 individuals. Nature Genetics, 2017, 49, 1758-1766.	21.4	470

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20	Treatment of Dyslipidemia in Diabetes: Recent Advances and Remaining Questions. Current Diabetes Reports, 2017, 17, 112.	4.2	22
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