

Supplementary Table 1. Genetic instruments of *HMGCR* and *PCSK9* variants

Gene	SNP	Effect allele frequency	Effect allele	Other allele	Beta	Standard error	p-value
<i>HMGCR</i>	rs12916	0.57	T	C	-0.06061	0.003	7.79E-78
	rs17238484	0.75	G	T	-0.05184	0.005	1.35E-21
	rs5909	0.90	G	A	-0.05102	0.007	4.93E-13
	rs2303152	0.88	G	A	-0.03498	0.005	1.04E-09
	rs10066707	0.58	G	A	-0.04110	0.005	2.97E-19
	rs2006760	0.81	C	G	-0.04407	0.006	1.67E-13
<i>PCSK9</i>	rs11206510	0.15	C	T	-0.06871	0.001	2.38E-53
	rs2479409	0.67	A	G	-0.05309	0.001	2.52E-50
	rs2149041	0.84	C	G	-0.05259	0.001	1.44E-35
	rs2479394	0.72	A	G	-0.03192	0.001	1.58E-19
	rs10888897	0.40	T	C	-0.04192	0.001	8.43E-31
	rs7552841	0.63	C	T	-0.04589	0.001	5.40E-15
	rs562556	0.19	G	A	-0.05292	0.002	6.16E-21

The presented original effect sizes were toward LDL cholesterol (continuous increase, mmol/L) from the genetic instruments introduced from previous studies (PMID: 27959767, 33886544).

After calculation of the causal estimates, the effect sizes were transformed toward eGFR % change by 50 mg/dL decrease in LDL cholesterol, calculated by $[\text{exponential}(\text{beta} \times -50/38.7) - 1] \times 100 (\%)$.

eGFR, estimated glomerular filtration rate; LDL, low-density lipoprotein; SNP, single nucleotide polymorphism.