

ESM Table 1: Established HDL-C SNPs considered for the study and their MAF in Pima Indians

SNP ID	Position ^a	Proxy SNP (r2) ^b	A1	A2	Locus ^c	MAF_Pima	MAF_EUR ^d	Primary Trait ^e
rs4660293	chr1:40028180	rs10127775 (r2=1)	A	G	<i>PABPC4</i>	0.03	0.23	HDL-C
rs1689800	chr1:182168885		A	G	<i>ZNF648</i>	0.16	0.35	HDL-C
rs4846914	chr1:230295691		G	A	<i>GALNT2</i>	0.44	0.4	HDL-C
rs1042034	chr2:21225281		C	T	<i>APOB</i>	0.21	0.22	TG
rs12328675	chr2:165540800		C	T	<i>COBLL1</i>	0	0.13	HDL-C
rs2972146	chr2:227100698		G	T	<i>IRS1</i>	0.001	0.37	HDL-C
rs13107325	chr4:103188709		C	T	<i>SLC39A8</i>	0.003	0.07	HDL-C
rs6450176	chr5:53298025		G	A	<i>ARL15</i>	0.23	0.26	HDL-C
rs2814944	chr6:34552797		G	A	<i>C6ORF106</i>	0.003	0.16	HDL-C
rs605066	chr6:139829666		C	T	<i>CITED2</i>	0.27	0.42	HDL-C
rs1084651	chr6:161089817		A	G	<i>LPA</i>	0.003	0.16	HDL-C
rs17145738	chr7:72982874		C	T	<i>MLXIPL</i>	0.11	0.12	TG
rs4731702	chr7:130433384		C	T	<i>KLF14</i>	0.39	0.48	HDL-C
rs9987289	chr8:9183358		A	G	<i>PPP1R3B</i>	0.24	0.09	HDL-C
rs7016529 ^f	chr8:19806631		C	T	<i>LPL</i>	0	0.01	TG
rs12678919	chr8:19844222	G	A	<i>LPL</i>	0.001	0.12	TG	
rs2293889	chr8:116599199	T	G	<i>TRPS1</i>	0.24	0.41	HDL-C	
rs10808546	chr8:126495818	C	T	<i>TRIB1</i>	0.33	0.44	TG	
rs643531	chr9:15296034	C	A	<i>TTC39B</i>	0.004	0.14	HDL-C	
rs11789603 ^f	chr9:107647019	C	T	<i>ABCA1</i>	0	0.09	HDL-C	
rs1883025	chr9:1076664301	C	T	<i>ABCA1</i>	0.39	0.25	HDL-C	
rs2923084	chr11:10388782	A	G	<i>AMPD3</i>	0.32	0.17	HDL-C	
rs3136441	chr11:46743247	rs3136457 (r2=1)	T	C	<i>LRP4</i>	0.36	0.15	HDL-C
rs174601	chr11:61623140		C	T	<i>FADS1-2-3</i>	0.009	0.36	TG
rs964184	chr11:116648917		G	C	<i>APOA1-C3-A4-A5</i>	0.42	0.13	TG
rs12225230 ^f	chr11:116728630		G	C	<i>APOA1-C3-A4-A5</i>	0.26	0.14	TG
rs7115089	chr11:122530591		C	G	<i>UBASH3B</i>	0.36	0.37	TC
rs7134375	chr12:20473758		C	A	<i>PDE3A</i>	0.46	0.42	HDL-C

SNP ID	Position ^a	Proxy SNP (r2) ^b	A1	A2	Locus ^c	MAF_Pima	MAF_EUR ^d	Primary Trait ^e
rs3741414	chr12:57844049		C	T	<i>LRP1</i>	0.39	0.24	TG
rs7134594	chr12:110000193		C	T	<i>MVK</i>	0.27	0.47	HDL-C
rs11057244 ^f	chr12:123773263		G	C	<i>ZNF664</i>	0.05	0.03	HDL-C
rs4759375	chr12:123796238		T	C	<i>SBNO1</i>	0	0.06	HDL-C
rs4765127	chr12:124460167		G	T	<i>ZNF664</i>	0.2	0.34	HDL-C
rs838880	chr12:125261593		C	T	<i>SCARB1</i>	0.42	0.31	HDL-C
rs1532085	chr15:58683366		A	G	<i>LIPC</i>	0.37	0.39	HDL-C
rs2070895 ^f	chr15:58723939	rs1077834 (r2=1)	G	A	<i>LIPC</i>	0.14	0.21	HDL-C
rs2652834	chr15:63396867		A	G	<i>LACTB</i>	0	0.2	HDL-C
rs3764261	chr16:56993324	rs17231506 (r2=0.98)	C	A	<i>CETP</i>	0.29	0.32	HDL-C
rs9939224 ^f	chr16:57002732	rs12720922 (r2=0.94)	G	T	<i>CETP</i>	0.35	0.76	HDL-C
rs16942887	chr16:67928042		G	A	<i>LCAT</i>	0.16	0.12	HDL-C
rs2925979	chr16:81534790		T	C	<i>CMIP</i>	0.05	0.3	HDL-C
rs4148008	chr17:6687294		C	G	<i>ABCA8</i>	0.25	0.32	HDL-C
rs11869286	chr17:37813856		G	C	<i>STARD3</i>	0.44	0.34	HDL-C
rs4129767	chr17:76403984		G	A	<i>PGS1</i>	0.42	0.49	HDL-C
rs7241918	chr18:47160953		G	T	<i>LIPG</i>	0	0.17	HDL-C
rs2040293 ^e	chr18:47278345		G	A	<i>LIPG</i>	0.32	0.26	HDL-C
rs12967135	chr18:57849023		G	A	<i>MC4R</i>	0.005	0.23	HDL-C
rs7255436	chr19:8433196		C	A	<i>ANGPTL4</i>	0.42	0.45	HDL-C
rs737337 ^f	chr19:11347493	rs56322906 (r2=1)	T	C	<i>LOC55908</i>	0.49	0.08	HDL-C
rs4420638	chr19:45422946		A	G	<i>APOE</i>	0.01	0.17	LDL-C
rs5167 ^e	chr19:45448465		T	G	<i>APOE-C1-C2</i>	0.49	0.37	LDL-C
rs386000	chr19:54792761		G	C	<i>LILRA3</i>	0.21	0.2	HDL-C
rs1800961 ^g	chr20:43042634		C	T	<i>HNF4A</i>	0.04	0.03	HDL-C
rs6065906	chr20:44554015		T	C	<i>PLTP</i>	0.005	0.18	HDL-C
rs181362	chr22:21932068		C	T	<i>UBE2L3</i>	0.27	0.2	HDL-C

MAF_Pima: minor allele frequency as determined by previously analyzed whole genome sequence data in 335 Pima Indians, MAF_EUR: MAF in subjects of European ancestry, A1: Reference allele, A2: Alternate allele.

^a Position is based on genome build 37.

^b Proxy SNP used for the actual analysis and r^2 with the GWAS index SNP.

^c Locus based on Teslovich et al. [7].

^d Minor allele frequency in Europeans from teslovich et al [7]

^e Primary lipid trait at the locus based on Teslovich et al. [7].

^f Minor allele frequency in Europeans from 1000 genome data.

^g rs1800961 is a missense variant and was included for further analysis even though it had a MAF=0.04.