

Table S4 continued. Associations with previously studied candidate variants.

Pathway	Locus	Gene	SNP	Variant	Position	Allele	Europeans				South Asians				r ² with best imputed proxy	Tagging level
							Freq	OR	95% CI	P	Freq	OR	95% CI	P		
Hemostasis																
	5q35.3	<i>F12</i>	rs1801020	C46T	176,769,138	A	0.24	1.07 (1.02-1.12)	2.0E-03	0.42	1.02 (0.96-1.08)	5.4E-01	1	1		
	6p25.1	<i>F13A1</i>	rs5985	V34L	6,263,794	A	0.25	0.98 (0.94-1.02)	3.8E-01	0.12	0.98 (0.90-1.08)	7.4E-01	1	2		
	1q24.2	<i>F5</i>	rs6025	R506Q	167,785,673	T	0.03	0.90 (0.80-1.01)	6.7E-02	0.01	0.97 (0.75-1.26)	8.4E-01	0.27	1		
	13q34	<i>F7</i>	rs6046	R353Q	112,821,160	A	0.11	1.00 (0.94-1.06)	9.8E-01	0.27	0.95 (0.89-1.02)	1.3E-01	1	1		
	17p13.2	<i>GP1BA</i>	rs2243093	T-5C	4,776,675	C	0.13	1.03 (0.97-1.08)	3.6E-01	0.12	1.03 (0.94-1.13)	5.3E-01	1	2		
	5q11.2	<i>ITGA2</i>	rs1062535 ^d	T275T	52,387,170	A	0.39	1.01 (0.9701-0.5)	6.9E-01	0.36	0.99 (0.93-1.05)	6.7E-01	1	1		
	17q21.32	<i>ITGB3</i>	rs5918	L33P	42,715,729	C	0.15	0.99 (0.94-1.05)	8.2E-01	0.09	0.89 (0.80-1.00)	4.0E-02	1	1		
Metabolic																
	5q33.1	<i>ADRB2</i>	rs1042713	G16R	148,186,633	A	0.37	1.04 (1.00-1.08)	4.1E-02	0.44	0.99 (0.94-1.06)	8.7E-01	1	1		
	5q33.1	<i>ADRB2</i>	rs1042714	E27Q	148,186,666	G	0.43	0.95 (0.91-0.98)	3.0E-03	0.22	1.03 (0.96-1.11)	3.9E-01	1	1		
	5q33.1	<i>ADRB2</i>	rs1800888	I164T	148,187,078	T	0.01	0.94 (0.80-1.10)	4.4E-01	<0.01	-	-	0.16	1		
	8p12	<i>ADRB3</i>	rs4994	W64R	37,942,955	G	0.07	0.99 (0.92-1.06)	6.7E-01	0.13	0.97 (0.89-1.06)	5.4E-01	0.43	1		
	16q24.3	<i>CYBA</i>	rs4673	Y72H	87,240,737	A	0.34	1.04 (1.00-1.08)	6.9E-02	0.37	0.99 (0.93-1.05)	6.8E-01	1	1		
	8q24.3	<i>CYP11B2</i>	rs1799998	T-344C	143,996,602	G	0.45	1.01 (0.97-1.04)	7.7E-01	0.44	0.99 (0.93-1.05)	7.0E-01	1	2		
	10q23.33	<i>CYP2C9</i>	rs1057910	I359L	96,731,043	C	0.07	1.02 (0.95-1.10)	6.0E-01	0.11	0.99 (0.90-1.08)	7.7E-01	1	2		
	10q23.33	<i>CYP2C9</i>	rs17443251 ^b		96,697,880	C	0.13	1.02 (0.96-1.07)	5.2E-01	0.05	0.87 (0.76-0.99)	4.0E-02	1	2		
	6q23.2	<i>ENPP1</i>	rs1044498	K121Q	132,214,061	C	0.14	1.03 (0.98-1.09)	2.8E-01	0.17	0.98 (0.91-1.06)	6.6E-01	1	2		
	2q36.3	<i>IRS1</i>	rs1801278	G971R	227,368,788	T	0.07	1.09 (0.97-1.22)	1.4E-01	0.04	1.02 (0.83-1.25)	8.7E-01	0.20	1		
	1p31.3	<i>LEPR</i>	rs1805096 ^e	P1019P	65,874,845	A	0.38	0.99 (0.96-1.03)	7.2E-01	0.45	0.99 (0.93-1.05)	7.1E-01	1	1		
	1p36.22	<i>MTHFR</i>	rs1801133	C677T	11,778,965	A	0.35	1.04 (1.00-1.08)	6.1E-02	0.16	1.04 (0.96-1.13)	2.9E-01	1	1		
	1q43	<i>MTR</i>	rs1805087	D919G	235,115,123	G	0.19	1.01 (0.96-1.06)	7.7E-01	0.31	0.98 (0.91-1.04)	4.4E-01	1	2		
	3p25.1-2	<i>PPARG</i>	rs1801282	Pro12Ala	12,368,125	G	0.12	1.01 (0.95-1.07)	7.2E-01	0.13	1.08 (1.00-1.18)	9.9E-02	1	1		
	2p11.2	<i>VAMP8</i>	rs1010		85,662,493	C	0.42	1.06 (1.03-1.10)	8.7E-04	0.38	1.04 (0.97-1.10)	2.5E-01 #	1	2		
Mineralcorticoid																
	17q23.3	<i>ACE</i>	rs4343 ^a	Ins/Del	58,919,763	A	0.47	0.97 (0.93-1.01)	9.2E-02	0.41	1.05 (0.99-1.11)	1.3E-01	1	1		
	1q42.2	<i>AGT</i>	rs699	M235T	228,912,417	G	0.41	1.03 (0.99-1.06)	1.9E-01	0.39	1.03 (0.97-1.10)	2.9E-01	1	1		
	3q24	<i>AGTR1</i>	rs5186	A1166C	149,942,678	C	0.30	1.00 (0.96-1.04)	8.6E-01	0.09	1.01 (0.91-1.12)	8.0E-01	0.96	1		
	14q32.3	<i>BDKRB2</i>	rs1799722	C-58T	95,740,892	T	0.49	0.99 (0.95-1.03)	6.3E-01	0.37	0.98 (0.92-1.04)	4.4E-01	0.18	2		
Other																
	4p16.3	<i>ADD1</i>	rs4961	G460W	2,876,505	T	0.19	0.99 (0.95-1.04)	7.6E-01	0.17	1.02 (0.94-1.10)	6.8E-01	1	1		
	9p21.3	<i>CDKN2A/B</i>	rs1333049		22,115,503	C	0.52	1.28 (1.23-1.33)	2.7E-34	0.51	1.12 (1.05-1.19)	3.3E-04	1	GWAS		
	22q11.21	<i>COMT</i>	rs4680	V158M	18,331,271	G	0.48	0.99 (0.95-1.02)	5.2E-01	0.46	0.97 (0.91-1.03)	3.3E-01	1	2		
	6q25.1	<i>ESR1</i>	rs2234693	T-401C	152,205,028	C	0.46	0.98 (0.95-1.02)	3.0E-01	0.42	1.02 (0.96-1.08)	5.6E-01	1	1		
	6p22.1	<i>HFE</i>	rs1799945	H63D	26,199,158	G	0.15	0.97 (0.92-1.02)	2.4E-01	0.07	0.88 (0.79-0.99)	3.1E-02	1	2		
	6p22.1	<i>HFE</i>	rs1800562	C282Y	26,201,120	A	0.06	1.03 (0.96-1.12)	3.9E-01	<0.01	-	-	1	2		
	13q14.2	<i>HTR2A</i>	rs9526246 ^c		46,347,862	A	0.42	1.05 (1.01-1.08)	1.9E-02	0.43	1.03 (0.97-1.10)	2.9E-01	1	2		
	12p12.3	<i>MGP</i>	rs1049897 ^f		14,925,788	A	0.39	1.03 (0.99-1.06)	1.9E-01	0.39	1.02 (0.96-1.08)	5.1E-01	1	2		
	1q41	<i>MIA3</i>	rs17465637		220,890,152	C	0.73	1.08 (1.02-1.14)	3.8E-03	0.64	1.09 (1.00-1.18)	5.2E-02	1	GWAS		
	1p36.22	<i>NPPA</i>	rs5065	R29X	11,906,068	G	0.15	0.97 (0.92-1.02)	2.9E-01	0.16	0.96 (0.89-1.04)	3.5E-01	1	1		
	5q31.3	<i>NR3C1</i>	rs6195	N363S	142,759,510	C	0.03	0.96 (0.87-1.07)	4.8E-01	<0.01			1	2		
	3q27.1	<i>THPO</i>	rs6141	A5713G	185,572,960	C	0.46	1.03 (0.99-1.07)	1.3E-01	0.47	0.97 (0.92-1.03)	3.6E-01	1	2		

Variants ordered by biological pathway, then gene. Per-allele odds ratios are presented for the effect allele, which is the minor allele in European populations.

r² with best imputed proxy was estimated using ~2.5M directly genotyped or HapMap-imputed SNPs in the CARDIoGRAM Consortium.

Tagging levels are 1 (r²>0.8 with all HapMap/Seattle SNPs of MAF≥0.02), 2 (r²>0.5 with all HapMap/Seattle SNPs of MAF≥0.05), 3 (only non-synonymous and known functional variants of MAF>0.01) and GWAS (specific SNPs previously identified in recent GWAS).

^a rs4343 has r²=1 with the insertion/deletion polymorphism in the *ACE* gene in CEU HapMap 2 population.

^b rs17443251 has r²=0.75 with the more commonly studied R144C variant (rs1799853) in the *CYP2C9* gene in CEU HapMap 2 population.

^c rs9526246 has r²=0.97 with the more commonly studied T102C variant (rs6313) in the *HTR2A* gene in CEU HapMap 2 population.

^d rs1062535 has r²=0.97 with the more commonly studied C807T variant (rs1126643) in the *ITGA2* gene in CEU HapMap 2 population.

^e rs1805096 has r²=0.89 with the more commonly studied rs6700896 variant in the *LEPR* gene in CEU HapMap 2 population.

^f rs1049897 has r²=1 with the more commonly studied A102T variant (rs4236) in the *MGP* gene in CEU HapMap 2 population.

^g rs4968624 has r²=0.97 with the more commonly studied L125V variant (rs668) in the *PECAM1* gene in CEU HapMap 2 population.

^h rs12944077 has r²=1 with the more commonly studied S563N variant (rs12953) in the *PECAM1* gene in CEU HapMap 2 population.