S1 Table. Association of rs6511720 genotype and CHD risk in CARDIoGRAM and C4D

SNP(allele)	log(OR) with CHD per-allele	SE (log(OR)	Sample numbers
rs6511720 (T~minor allele) CARDIoGRAM	- 0.1253	0.03321	8948cases 4747 controls
rs6511720 (T~minor allele) C4D	- 0.1082	0.0287	15393 cases 15036 controls
fixed effects meta-analysis	- 0.1155	0.0217	24341 cases 19783 controls