IGF1 Gene Is Associated With Triglyceride Levels In Subjects With Family History Of Hypertension From The SAPPHIRe And TWB Projects

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Supplementary Table S1. IGF1 single-nucleotide polymorphisms (SNPs) genotyped in the Stanford Asian Pacific Program in Hypertension

SNP	Chromosome position ^a	Gene region	Allele type Minor/Major	MAF	<i>p</i> for HWE test	Number of subject successfully genotyped
rs2288377 ^b	102874762	Promoter	A/T	0.30	0.39	972 (99.3%)
rs2195239	102462924	Intron 2	G/C	0.45	0.42	973 (99.4%)
rs978458 ^b	102408461	Intron 3	A/G	0.44	0.77	956 (97.7%)
rs1520220	102402744	Intron 3	G/C	0.45	0.73	976 (99.7%)
rs6220	102400737	3'UTR	C/T	0.44	0.91	946 (96.6%)
rs6217 ^b	102400008	3'UTR	G/T	0.28	0.10	952 (97.2%)
rs6218	102399855	3'UTR	C/T	0.27	0.46	930 (95.0%)
rs6214 ^b	102399791	3'UTR	A/G	0.47	0.11	964 (98.5%)
rs6219 ^b	102396414	3'UTR	A/G	0.17	0.55	977 (99.8%)

and Insulin Resistance (SAPPHIRe) sample

^a GRCh38.p7.

^b The tag-SNPs.

MAF, minor allele frequency; HWE, Hardy-Weinberg equilibrium.

Supplementary Table S2. Comparison of genotype distribution of rs978458 in different

subsets of the Taiwan biobank (TWB) sample

Genotype	Non-hypertensive subjects without a family history (%)	Non-hypertensive subjects with a family history (%)	Hypertensive subjects without a family history (%)	Hypertensive subjects with a family history (%)	р
GG	2690 (28.59%)	335 (27.06%)	250 (29.00%)	465 (29.87%)	0.22
AG	4703 (49.92%)	642 (51.86%)	422 (48.96%)	799 (51.32%)	
AA	2025 (21.49%)	261 (21.08%)	190 (22.04%)	293 (18.82%)	

Number of hypertensive members	Number of families (%)
1	182 (45.84%)
2	120 (30.23%)
3	60 (15.11%)
4	23 (5.79%)
5	8 (2.02%)
6	3 (0.76%)
7	1 (0.25%)

Supplementary Table S3. Distribution of the number of hypertensive members in the

Stanford Asian Pacific Program in Hypertension and Insulin Resistance (SAPPHIRe) families



Supplementary Figure S1. Linkage disequilibrium (LD) structure of the IGF1 SNPs

genotyped in the SAPPHIRe sample. The pair-wise LD levels were measured by r^2 and were

estimated using the Haploview (v. 4.2) program.