

Table S1 Primer sequence of the single nucleotide polymorphisms used in the study

SNP	Allele change	Primer sequence
rs10983755	G>A	F,5'-ATTGGAAGTGCTTGGAGGA-3' R,5'-TTGTAAAGCTTTTAGGACAGTGTCT-3'
rs11536879	A>G	F,5'-CATGTTCTCTGCAATGGTTTG-3' R,5'-GCAGCAAGGGTTGAAAAACT-3'
rs4986790	A>G	F,5'-AGCATACTTAGACTACTACCTCCATG-3' R,5'-GAGAGATTTGAGTTTCAATGTGGG-3'
rs4986791	C>T	F,5'-GGTTGCTGTTCTCAAAGTGATTTTGGGAGAA-3' R,5'-GGAAATCCAGATGTTCTAGTTGTTCTAAGCC-3'
rs1927907	G>A	F,5'-TTCAACCCTTGCTGCTTTCT-3' R,5'-CGAAAAGGCAAAGGATGTCT-3'
rs7873784	G>C	F,5'-ACGTTGGATGATGAGAGGTACCCTCTTAAC-3' R,5'-ACGTTGGATGGCTCTAAAGATCAGCTGTAT-3'
rs115336889	G>C	F,5'-GTCATTCCAAAGTTATTGCCTA-3' R,5'-CTGTGGTCATATTTCCAGTTTT-3'

Table S2 Association of haplotypes composed of rs10983755, rs4986791, rs1927907 and rs115336889 with susceptibility to benign prostatic hyperplasia.

Haplotype	Case		Control		Odds ratio	Lower limit	Upper limit	P-value
	Frequency	Number	Frequency	Number				
GCGG	0.032	16	0.111	107	0.26	0.15	0.45	<0.001
GCGC	0.138	69	0.190	183	0.68	0.50	0.92	0.012
GTGG	0.108	54	0.126	121	0.84	0.60	1.18	0.321
GTGC	0.461	231	0.214	206	3.15	2.49	3.98	<0.001
GTAC	0.069	35	0.053	52	1.32	0.85	2.05	0.221
ATGC	0.081	41	0.053	52	1.56	1.02	2.39	0.038

Table S3 Comparison of haplotypes composed of rs10983755, rs4986791 and rs1927907 between aggressive and non-aggressive benign prostatic hyperplasia populations

Haplotype	Aggressive		Non-aggressive		Odds ratio	Lower limit	Upper limit	P-value
	Frequency	Number	Frequency	Number				
GCG	0.119	26	0.202	58	0.54	0.32	0.88	0.014
GTG	0.673	145	0.495	141	2.09	1.45	3.01	<0.001
GTA	0.075	16	0.087	25	0.83	0.43	1.6	0.581
ATG	0.092	20	0.109	31	0.84	0.46	1.51	0.553