

## Supplementary material

**Table S1.** PCR-RFLP primer sequences of 7 SNPs for NNMT gene polymorphisms

SNP_I D	Primer Forward	Primer Reverse
rs694539	ACGTTGGATGGTCCTAGAGTCCTA	ACGTTGGATGCAGCCATCTCAAATG
rs225629	GAATCC	GATGC
2	ACGTTGGATGTTTAAGGTCTAGGA	ACGTTGGATGCCATGTAACAGACTT
rs230112	GAAGG	TCTGG
8	ACGTTGGATGTTATTCCCAATCCA	ACGTTGGATGTTTTTACCTTCTCCTA
rs108916	GGGTG	GACC
45	ACGTTGGATGTGATACTATCAACCT	ACGTTGGATGGAGTGTATATTGTTCA
rs215580	GCCTG	GTTG
6	ACGTTGGATGCAGCAATATTAGGT	ACGTTGGATGCGTAGATTACAGACT
rs194139	TCACCG	TTGGG
8	ACGTTGGATGCTTAGCTCCTTGAAT	ACGTTGGATGGAGGGAAAGGACAA
rs260427	GGCAC	ATTGAC
9	ACGTTGGATGCTGCTTGCTTCAGT CTGGTT	ACGTTGGATGAGCTCCTCCAAAAG GAGTC

**Table S2.** Distribution of NNMT genotypes and alleles between cases and controls

SNP	Haplotype /Allele	Cases	Controls	Chi <sup>2</sup>	P-value	OR(95%CI)
<u>rs694539</u>	AA	2(0.048)	7(0.081)			
	AG	21(0.500)	46(0.535)			
	GG	19(0.452)	33(0.384)	0.8509	0.6535	
	A	25(0.298)	60(0.349)			
	G	59(0.702)	112(0.651)	0.6675	0.4139	0.791(0.450-1.389)
<u>rs2256292</u>	CC	5(0.119)	10(0.116)			
	CG	18(0.429)	44(0.512)			
	GG	19(0.452)	32(0.372)	0.8603	0.6504	
	C	28(0.333)	64(0.372)			
	G	56(0.667)	108(0.628)	0.3683	0.5439	0.844(0.487-1.461)
<u>rs2301128</u>	AA	0(0.000)	1(0.012)			
	AG	7(0.167)	18(0.209)			
	GG	35(0.833)	67(0.779)	0.8553	0.6521	
	A	7(0.083)	20(0.116)			
	G	77(0.917)	152(0.884)	0.6493	0.4204	0.691(0.280-1.705)
<u>rs10891645</u>	AC	5(0.119)	10(0.116)			
	CC	37(0.881)	76(0.884)	0.0021	0.9635	
	A	5(0.060)	10(0.058)			
	C	79(0.940)	162(0.942)	0.002	0.9647	1.025(0.339-3.101)
<u>rs2155806</u>	CC	2(0.048)	0(0.000)			

<b>rs1941398</b>	CT	9(0.214)	23(0.267)			
	TT	31(0.738)	63(0.733)	4.4154	0.11	
	C	13(0.155)	23(0.134)			
	T	71(0.845)	149(0.866)	0.2068	0.6493	0.843(0.404-1.761)
	CC	24(0.571)	55(0.640)			
<b>rs2604279</b>	CG	14(0.333)	30(0.349)			
	GG	4(0.095)	1(0.012)	5.2819	0.0713	
	C	62(0.738)	140(0.814)			
	G	22(0.262)	32(0.186)	1.9512	0.1624	0.644(0.347~1.197)
	CC	2(0.048)	0(0.000)			
	CT	12(0.286)	21(0.244)			
	TT	28(0.667)	65(0.756)	4.5927	0.1006	
	C	16(0.190)	21(0.122)			
	T	68(0.810)	151(0.878)	2.1345	0.144	0.591(0.290-1.203)

**Table S3.** Frequency and analysis of observed haplotypes (rs2256292, rs2301128, rs10891645) in patients with SZ and in healthy controls

Haplotype	Case(freq%)	Control(freq%)	P-value	Odds Ratio (95%CI)
<b>Total</b>				
CAC	7.9	11.6	0.3579	0.651(0.259-1.636)
CGC	25.0	25.5	0.9407	0.977(0.536-1.784)
GGA	5.5	5.8	0.9268	0.948(0.304-2.958)
GGC	61.2	57.0	0.4973	1.203(0.705-2.052)
<b>Female</b>				
CAC	9.5	12.3	0.6375	0.753(0.231-2.457)
CGC	21.4	29.2	0.3343	0.660(0.283-1.540)
GGA	4.8	4.7	0.9908	1.010(0.188-5.420)
GGC	64.3	53.8	0.2445	1.547(0.740-3.236)
<b>Male</b>				
CAC	5.2	10.6	0.3424	0.474(0.099-2.278)
CGC	28.6	16.1	0.1091	2.135(0.835-5.461)
GGA	5.2	4	0.7517	1.434(0.215-8.390)
GGC	59.1	65.7	0.5676	0.791(0.354-1.769)

All those frequency less than 0.03 will be ignored in analysis