

Table S1. Genome-wide significant SNPs in the GWASs reported by Yue et al.¹ and Shi et al.²

CHR	SNP	Allele	Frequency	Case	Control	P-value	OR	Ref.
6	rs1233710	T/C	0.350	4,773	7,207	4.76×10 ⁻¹¹	0.79	1
6	rs1635	A/C	0.350	4,773	7,207	6.91×10 ⁻¹²	0.78	1
6	rs2142731	G/A	0.242	4,773	7,207	5.14×10 ⁻¹⁰	0.79	1
11	rs11038167	A/C	0.375	4,773	7,207	1.09×10 ⁻¹¹	1.29	1
11	rs11038172	A/G	0.399	4,773	7,207	7.21×10 ⁻¹⁰	1.25	1
11	rs835784	A/G	0.249	4,773	7,207	2.73×10 ⁻¹¹	1.27	1
1	rs10489202	T/G	0.131	8,133	11,007	9.50×10 ⁻⁰⁹	1.23	2
8	rs16887244	G/A	0.326	8,133	11,007	1.27×10 ⁻¹⁰	0.84	2
8	rs1488935	A/G	0.320	8,133	11,007	5.06×10 ⁻⁰⁹	0.85	2

The frequency is based on the first allele.

Abbreviation:

CHR, chromosome; SNP, single nucleotide polymorphism; OR, odds ratio; Ref., references.

Table S2. Genome-wide significant SNPs in the GWASs reported by Li et al.³ and Yu et al.⁴

CHR	SNP	Allele	Frequency	Case	Control	P-value	OR	Ref.
2	rs1518395	A/G	0.308	12,083	24,097	3.78×10 ⁻¹³	0.862	3
2	rs78681500	T/C	0.063	12,083	24,097	3.08×10 ⁻¹⁰	0.751	3
3	rs2073499	A/G	0.312	12,083	24,097	2.61×10 ⁻⁰⁸	0.899	3
6	rs111782145	C/T	0.037	12,083	24,097	1.80×10 ⁻⁰⁸	0.813	3
6	rs7757969	C/T	0.269	12,083	24,097	4.82×10 ⁻⁰⁸	1.110	3
6	rs4479915	G/C	0.254	12,083	24,097	4.82×10 ⁻⁰⁹	0.876	3
7	rs11534004	G/A	0.334	12,083	24,097	1.71×10 ⁻⁰⁸	0.890	3
2	rs1051061	G/A	0.412	8,723	12,813	1.14×10 ⁻¹⁰	1.17	4
6	rs115070292	G/A	0.136	8,723	12,813	4.96×10 ⁻¹⁰	0.77	4
10	rs10883795	C/T	0.496	8,723	12,813	7.94×10 ⁻¹⁰	0.87	4
10	rs10883765	C/T	0.449	8,723	12,813	3.06×10 ⁻⁰⁹	0.87	4

The frequency is based on the first allele.

Abbreviation:

CHR, chromosome; SNP, single nucleotide polymorphism; OR, odds ratio; Ref., references.

Table S3. “Leave-one-out” sensitivity analysis for meta-analysis

Excluded study	Remaining sample size		Heterogeneity	Random effect model			Fixed effect model		
	Case	Control		OR	95%CIs	Meta P value	OR	95%CIs	Meta P value
Shi, 2011a	10,590	13,881	p=0.250, $I^2=21.7\%$	1.124	1.053-1.198	4.00×10^{-4}	1.132	1.074-1.194	4.03×10^{-6}
Shi, 2011b	9,957	15,810	p=0.092, $I^2=41.4\%$	1.133	1.048-1.226	1.81×10^{-3}	1.148	1.086-1.215	1.37×10^{-6}
Ma, 2012	13,364	19,306	p=0.089, $I^2=41.7\%$	1.149	1.071-1.233	1.13×10^{-4}	1.165	1.111-1.223	4.97×10^{-10}
Jin, 2013	13,247	19,327	p=0.743, $I^2=0$	1.185	1.129-1.244	6.42×10^{-12}	1.185	1.129-1.244	6.42×10^{-12}
Wong, 2013	13,842	18,324	p=0.089, $I^2=41.7\%$	1.149	1.073-1.231	7.19×10^{-5}	1.164	1.110-1.221	3.39×10^{-10}
Saito, 2014	13,308	19,356	p=0.093, $I^2=41.1\%$	1.136	1.060-1.218	3.31×10^{-4}	1.156	1.102-1.213	3.64×10^{-9}
Guan, 2015	12,869	18,821	p=0.111, $I^2=38.6\%$	1.154	1.076-1.238	6.33×10^{-5}	1.171	1.115-1.230	3.39×10^{-10}
Li, 2016a	14,040	20,049	p=0.085, $I^2=42.4\%$	1.141	1.066-1.221	1.33×10^{-4}	1.159	1.106-1.215	7.75×10^{-10}
Li, 2016b	13,940	19,949	p=0.081, $I^2=43.0\%$	1.146	1.070-1.227	1.04×10^{-4}	1.162	1.108-1.218	4.66×10^{-10}
Current	13,903	18,318	p=0.081, $I^2=43.0\%$	1.140	1.062-1.223	2.83×10^{-4}	1.159	1.105-1.216	1.74×10^{-9}

Table S4. Association of rs10489202 with nearby gene expression in the lymphoblastoid cell lines from 85 East Asian individuals⁵

	Beta estimate	Standard error	T-value	P-value
CD247	-0.00302	0.0361	-0.0837	0.934
CREG1	-0.0306	0.0730	-0.419	0.676
RCSD1	0.0232	0.0803	0.289	0.774
MPZL1	0.00714	0.0541	0.132	0.895
ADCY10	0.000448	0.0158	0.0283	0.977
MPC2	-0.0232	0.0671	-0.346	0.730
DCAF6	0.0505	0.0632	0.797	0.427
GPR161	-0.0278	0.0192	-1.443	0.153
TIPRL	-0.313	0.0868	-3.605	0.000567
SFT2D2	-0.00534	0.0264	-0.202	0.840
TBX19	0.0486	0.0278	1.747	0.0848

References

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- Shi Y, Li Z, Xu Q, Wang T, Li T, Shen J *et al.* Common variants on 8p12 and 1q24.2 confer risk of schizophrenia. *Nat Genet* 2011; **43**(12): 1224-7.
- Li Z, Chen J, Yu H, He L, Xu Y, Zhang D *et al.* Genome-wide association analysis identifies 30 new susceptibility loci for schizophrenia. *Nat Genet* 2017; **49**(11): 1576-83.
- Yu H, Yan H, Li J, Li Z, Zhang X, Ma Y *et al.* Common variants on 2p16.1, 6p22.1 and 10q24.32 are associated with schizophrenia in Han Chinese population. *Mol Psychiatry* 2017; **22**(7): 954-60.
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Table S5. Functional prediction analysis of the SNPs in high LD with rs10489202 ($r^2 > 0.8$) using the Gwava dataset (http://www.sanger.ac.uk/sanger/StatGen_Gwava)

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21
#																					
ID	rs2420	rs17556487	rs10489202	rs1846416	rs4656564	rs7535138	rs2272911	rs718885	rs149912	rs12404574	rs4657727	rs12727036	rs1060041	rs7538581	rs12408205	rs17557135	rs17557162	rs11587594	rs66950781	rs17485889	rs4657730
ZEB1	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
ZNF143	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
ZNF263	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
ZNF274	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
ZZZ3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
bound_motifs	-	-	-	-	3	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
cpg_island	-	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
dnase_fps	1	-	-	-	29	-	-	-	1	-	-	-	-	-	-	-	-	-	-	-	-
in_cpg	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
pwm	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
repeat	1	-	-	-	-	1	1	-	1	-	1	-	-	-	-	-	-	1	-	-	-
seq_A	1	-	-	-	-	-	-	1	1	1	1	1	1	1	1	1	1	-	-	1	-
seq_C	-	-	-	-	-	-	-	1	-	-	-	-	-	-	-	-	-	1	-	-	-
seq_G	-	-	-	1	1	-	-	-	-	-	-	-	-	-	-	-	-	-	1	-	-
seq_T	-	-	1	-	-	1	1	-	-	-	-	-	-	-	-	-	1	-	-	-	-