Disrupted-in-Schizophrenia-1 SNPs and Susceptibility to Schizophrenia: Evidence from Malaysia

A. Talib Norlelawati1, Abdullah Kartini2, Kuzaifah Norsidah1, Musa Ramli2, Abdul Razak Tariq1, and Wan Taib Wan Rohani3

1Department of Basic Medical Sciences, Kulliyyah of Medicine, International Islamic University, Pahang, Malaysia
2Department of Psychiatry, Kulliyyah of Medicine, International Islamic University Malaysia, Pahang, Malaysia
3Faculty of Medicine, University of Sultan Zainal Abidin, Terengganu, Malaysia

Objective Even though the role of the DISCI gene as a risk factor for schizophrenia is still unclear, there is substantial evidence from functional and cell biology studies that supports the connection of the gene with schizophrenia. The studies associating the DISC1 gene with schizophrenia in Asian populations are limited to East-Asian populations. Our study examined several DISC1 markers of schizophrenia that were identified in the Caucasian and East-Asian populations in Malaysia and assessed the role of rs2509382, which is located at 11q14.3, the mutual translocation region of the famous DISC1 translocation [t (1; 11) (p42.1; q14.3)].

Methods We genotyped eleven single-nucleotide polymorphism (SNPs) within or related to DISC1 (rs821597, rs821616, rs4658971, rs1538979, rs843979, rs2812385, rs1407599, rs4658890, and rs2509382) using the PCR-RFLP methods.

Results In all, there were 575 participants (225 schizophrenic patients and 350 healthy controls) of either Malay or Chinese ethnicity. The case-control analyses found two SNPs that were associated with schizophrenia [rs4658971 (p=0.030; OR=1.43 (1.35–1.99)) and rs1538979-(p=0.036; OR=1.35 (1.02–1.80))] and rs2509382-susceptibility among the males schizophrenics [p=0.0082; OR=2.16 (1.22–3.81)]. This is similar to the meta-analysis findings for the Caucasian populations.

Conclusion The study supports the notion that the DISC1 gene is a marker of schizophrenia susceptibility and that rs2509382 in the mutual DISC1 translocation region is a susceptibility marker for schizophrenia among males in Malaysia. However, the finding of the study is limited due to possible genetic stratification and the small sample size.

Key Words Schizophrenia, SNPs, Susceptibility, DISC1.
DISC1 and Schizophrenia were found to exhibit stress-induced prefrontal cortex dysfunction, an important precondition for the development of schizophrenia.

Because genetic variations differ among populations, it is important to assess the variations of DISC1 gene role in schizophrenia in a local context before making a definitive decision regarding its role in schizophrenia. Regrettably, in Asia, there had been only few studies that have investigated the role of the DISC1 gene in Schizophrenia, and these studies were mainly performed on East-Asian populations. There were conflicting findings on the association between schizophrenia and DISC1 in Chinese and Korean populations whilst a study on DISC1/TRAX locus found significant linked of the locus in the Japanese. In addition to the DISC1 locus, the mutual translocated regions of the DISC1 gene (1; 11) (q42.1; q14.3) are also areas of interest. A recent meta-analysis of several genome-wide studies that also included publicly available data in the Genetic Association Information Network (GAIN) (www.genome.org) identified rs2509382 as being associated with schizophrenia among males. This is an interesting finding because rs2509382 is located at 11q14.3 and represents the mutual translocation region.

In the present study, a case-control association analysis was performed to examine several DISC1 markers of schizophrenia that were identified in Caucasian and East-Asian populations in terms of their relevance to Malaysian patients. The study also assessed the role of rs2509382 among the patients.

METHODS

Ethics approval
The study was registered under the National Medical Research Registry of Malaysia (NMRR-10-832-6366), and the protocol was approved by both the Medical Research and Ethics Committee, Ministry of Health, Malaysia, and the Kulliyyah of Medicine, International Islamic University, Malaysia (IIUM).

Study populations
The study consisted of 225 unrelated schizophrenia patients and 350 healthy controls. Schizophrenia cases were collected from the Psychiatry Clinic of Tengku Ampuan Afzan Hospital, Kuantan (HTAA). The patients who attended follow-up appointments at the clinic were either from Kuantan (the state capital of Pahang) or referred from other districts in Pahang or the town Kemaman (in Terengganu). The control subjects were from the district of Kuantan. All subjects were over 18 years of age and were legitimately identified as being of Malay or Chinese ethnicity based on their personal identification card issued by the National Registry Department of Malaysia. These subjects also had grandfathers and grandmothers of similar ethnicities. The schizophrenic subjects had a consensus best-estimate DSM-IV diagnosis of schizophrenia of at least six months duration, and the symptoms of psychosis were not deemed to be secondary to substance use or neurological disorders. Mentally retarded patients were not included as subjects. In all, there were 225 (147 males and 78 females) schizophrenic patients and 350 controls (233 males and 117 females), with those of Chinese ethnicity representing only 43 and 10 of the cases and control subjects, respectively. The sample size was calculated by using OpenEpi for the unmatched case-control study. Using data from Schumacher et al. for rs4658890 (MAF=0.485), a total sample size of 994 (1:1 case-control ratio) was needed for a standard 95% two-sided confidence level and an 80% power of detection.
In our study, a total of 225 schizophrenia patients and 350 controls would give less than a 60% power of detection at a 95% confidence level.

**Genotyping and selection of SNPs**

Peripheral blood samples were collected from all 575 subjects. Genomic DNA was extracted using a Gentra® Puregene® (Qiagen, Germany) purification kit, following the manufacturer’s protocol. Four SNPs (rs821597, rs4658971, rs843979, and rs821616) that were located in the region between Intron 10 and Exon 11 of DISC1 were selected based on a previous study that was conducted on Han Chinese subjects. Another six SNPs (rs1538979, rs2812385, rs4658890, rs1407599, and rs2038636) were selected based on the meta-analysis findings of DISC1 studies performed among Caucasians in Europe. The last SNP, rs2509382, which was located at 11q14.3, represented the mutual translocation region of the famously translocated DISC1 (1q42.1) gene. The intermarker linkage disequilibrium patterns of the SNPs are shown in Figure 1.

SNPs were genotyped using PCR-RFLP methods. PCR was performed in a 15 µL reaction volume that included 0.25 µM of each primer, 200 µM of dNTPs, 1 U of One Taq™ Polymerase (New England Biolabs), 1X reaction buffer of various MgCl2 titrations, and 30 ng of DNA. The information regarding the primers and restriction enzymes used in the assay is shown in Table 1. The PCR products were digested with their respective restriction enzymes, followed by electrophoretic separation on 3.2% agarose gel. Only clear and unequivocal genotypes were included in the data analysis. Thus, the PCR-RFLP success rate of most of the SNPs studied ranged between 85% and 99%. Only one SNP (rs843979) had success rate of 73%. A minimum of 3 samples per SNP, each representing the ancestor, variant and mixed genotypes were randomly chosen based on the RFLP result, for direct nucleotide sequencing. The concordance rate between RFLP and sequencing was 100%.

| Table 1. Information of the selected SNPs and the genotyping assays |
|---|---|---|---|---|---|---|
| SNP ID | Location Chr:Kbp | Primer sequence (5'-3') | Allele | Restriction nuclease (NEB®) | Amplified restricted products | PCR condition |
|---|---|---|---|---|---|---|
| rs821597* | 1:238.26 | F-CAGGGT TAT TGT TTCCAGAGG R-CAA GTG CTC AAG AGT TC TG | A/G | EcoR1 | 432→217+215 | 4 | 60 |
| rs4658971* | 1:243.81 | F-AGAGAAGAATCACCAACAGG R-AGCCACACAAATGAGCACAC | T/C | BsrB1 | 357→208+149 | 5 | 65 |
| rs843979* | 1:266.61 | F-GAGGAGGATATAGATACCAG R-TCTTTCTTGTCTTGGCCTTCC | C/G | HinfI | 470→383+87 | 5 | 65 |
| rs821616* | 1:280.60 | F-GTATTGGGCTGCTGAGTCTG R-GACCTCTTTCTGTTCACC TCC | A/T | BsrI | 540→336+204 | 4 | 65 |
| rs1538979 | 1:23.87 | F-ACAGGAACACACAGACACTAGACG R-ATACAGGTTCTCCTACATGGCAGCCACCCAGT | C/T | MspI | 249→257+32 | 2 | 64 |
| rs16854957 | 1:48.89 | F-AGCATATCAGCATCTACAAGCTTCG R-AAATGCAAAGCTCTCCTCTT | C/A | Tsp5091 | 187→154+33 | 2 | 55 |
| rs2812385 | 1:0 | F-CACGGCTGACATCCCTTTGGTG R-GTGTCTGTAAAAGGCAATTTAGG | G/T | SspI | 220→129+91 | 2 | 65 |
| rs2038636 | 1:191.25 | F-CAGGTCAATGGGCTAGCTGAGG R-AAATGCGACGTGAGG | G/A | Tsp5091 | 108→75+33 | 2 | 55 |
| rs4658890 | 1:160.08 | F-GCAACGTACGAGGCTGCTGA | G/T | HinfI | 225→131+94 | 2 | 65 |
| rs1407599 | 1:163.91 | F-AGTCTCTGCTGTTGAAAGAGTCTCCTG R-TCTCTCTGACCCACTGCTTCCCT | G/T | Tsp5091 | 210→156+54 | 2 | 65 |
| rs2509382 | 11:NA | F-GCCCTTAGATAAAAAGGCCAGA R-AACAAATAAAAATGAAAATTTGCAACACTT | G/C | Taqα1 | 167→136+30 | 5 | 58 |

All variants are located within the intron regions except for rs821616 and rs2509382 which is located within the exon of DISC1 and intergenic region in Chr18 respectively. *primers designed by Qu et al. (2007). Chr: Chromosome, Kbp: Nearest kilo-base pair distance from rs2812385, NA: not applicable, SNP: single nucleotide polymorphism.
Statistical analyses
The analyses of the allelic and genotypic differences in DISC1 gene markers between schizophrenic and control subjects were evaluated via Chi-square test, which was performed using the Shesis software program.\(^{23}\) The odds ratio (OR) for the alleles’ effect and the deviation of the genotype counts from the Hardy-Weinberg equilibrium were tested using the same software. A Bonferroni correction for multiple testing was carried out to exclude type I errors (corrected significant p of <0.005). Data from studies performed among the Chinese\(^{16,17}\) and Japanese populations\(^{24}\) were used for meta-analyses. The meta-analyses and heterogeneity tests were performed using the Comprehensive Meta-analysis Version 2 software program.\(^{25}\)

RESULTS
Ten SNPs within the DISC1 gene and rs2509382 in the DISC1 mutual translocated region of 11q14.3 were genotyped in 225 schizophrenic patients and 350 healthy controls from Malay-Malaysian and Chinese-Malaysian populations. Genotype testing on two SNPs (rs2038636 and r16854957) found they were not polymorphic in the populations studied. None of the controls and cases showed heterozygous or homozygous variants. The genotypic and allelic differences between schizophrenic patients and controls for the remaining nine SNPs are shown in Table 2. The distributions of all of the SNP genotypes conformed to Hardy-Weinberg equilibrium in the control samples.

There were significant differences in genotypes and allele

| Marker | N  | Genotype | p-value (2df) | HWE | Allele | p-value (1df) | OR (95% CI) |
|--------|----|----------|---------------|------|--------|---------------|-------------|
| rs821597 |    |          |               |      |        |               |             |
| Patients | 196 | AA       | 38 (0.20)     | 0.978| 0.138  | 0.980         | 1.003 (0.77–1.31) |
| Controls | 294 | AG       | 84 (0.43)     | 0.135| 242 (0.41)| 1.003         |             |
|         |     | GG       | 72 (0.37)     |      | 346 (0.59) |             |             |
| rs821616 |    |          |               |      |        |               |             |
| Patients | 218 | TT       | 2 (0.01)      | 0.886| 0.367  | 0.744         | 1.06 (0.73–1.54) |
| Controls | 293 | AT       | 51 (0.23)     |      | 381 (0.87) |             |             |
|         |     | AA       | 165 (0.76)    |      | 381 (0.87) |             |             |
| rs4658971 |   |          |               |      |        |               |             |
| Patients | 199 | TT       | 20 (0.10)     | 0.00067*| 0.138  | 1.44         | 1.05–2.00   |
| Controls | 303 | CT       | 80 (0.26)     | 0.0198*| 0.365  | 1.35         | 1.02–1.80   |
|         |     | CC       | 216 (0.71)    |      | 505 (0.78) |             |             |
| rs843979 |    |          |               |      |        |               |             |
| Patients | 190 | CC       | 27 (0.14)     | 0.312| 0.661  | 0.434         | 0.90–1.18   |
| Controls | 258 | CG       | 93 (0.49)     | 0.121| 213 (0.41)| 0.303        |             |
|         |     | GG       | 70 (0.37)     |      | 303 (0.59) |             |             |
| rs2812385 |   |          |               |      |        |               |             |
| Patients | 216 | GG       | 44 (0.20)     | 0.250| 0.463  | 1.00         | 1.23 (0.96–1.58) |
| Controls | 320 | GT       | 101 (0.47)    | 0.780| 247 (0.39)| 0.587        | 1.10 (0.77–1.57) |
|         |     | TT       | 71 (0.33)     |      | 391 (0.61) |             |             |
| rs1407597 |   |          |               |      |        |               |             |
| Patients | 222 | GG       | 4 (0.02)      | 0.763| 0.975  | 0.587         | 1.10 (0.77–1.57) |
| Controls | 346 | GT       | 52 (0.23)     | 0.542| 86 (0.12) | 0.60         |             |
|         |     | TT       | 166 (0.75)    |      | 607 (0.88) |             |             |
| rs4658890 |   |          |               |      |        |               |             |
| Patients | 216 | GT       | 41 (0.19)     | 0.64 | 0.0504 | 0.675         | 0.95 (0.75–1.21) |
| Controls | 336 | GC       | 122 (0.56)    | 0.268| 326 (0.48)| 0.075        | 1.50 (0.96–2.35) |
|         |     | GG       | 53 (0.24)     |      | 346 (0.51) |             |             |
| rs2509382 |   |          |               |      |        |               |             |
| Patients | 209 | GC       | 3 (0.014)     | 0.213| 0.385  | 0.075         | 1.50 (0.96–2.35) |
| Controls | 337 | GC       | 34 (0.163)    | 0.385| 378 (0.904)| 0.075       |             |
|         |     | GG       | 172 (0.823)   |      | 624 (0.934) |             |             |

\(^*\)-test, statistical significant at \(*p<0.05\). All genotypes distribution in controls conformed to Hardy Weinberg Equilibrium (HWE). SNP: single nucleotide polymorphism
frequencies between patients and controls for two SNPs, rs4658971 and rs1538971. Those having the T minor allele instead of the C major allele for rs4658971 \( p=0.0303, OR=1.44 (1.03–2.00) \) and rs1538971 \( p=0.036, OR=1.35 (1.02–1.80) \) are more susceptible to schizophrenia. Except for significant genotypic differences for rs4658971 \( \chi^2=14.66, p=0.00067 \), the genotypic and allelic differences were reversed after a Bonferroni correction for multiple testing. The nine SNPs were further tested for a relationship with disease based on the gender. The results of the genotype and allele frequencies in male participants for SNP rs2509382 \( p=0.0082, OR=2.16 (1.22–3.81); \) corrected \( p=0.090 \).

The meta-analyses results for rs821616, rs821597, rs843979, and rs4658971 are shown in Figure 2. There was a significant association between rs821957 and schizophrenia in the combined data \( p=0.031, OR=1.14 (1.01–1.28); \) heterogeneity test \( I^2=62.6, p=0.069 \). In contrast, the meta-analysis of rs4658971 found no significant association of this SNP with schizophrenia.

**DISCUSSION**

There are conflicting findings regarding the association of the DISC1 gene with schizophrenia. This inconsistency could be due to the limited selection of polymorphic markers that were tested or the inadequacy of the sample's representation. Based on the summary of all published association studies of the DISC1 gene and schizophrenia that were accessed in the Schizophrenia Research Forum (www.szgene.org), there were more than 25 studies performed on Caucasian popula-

### Table 3. Association of genotypes and alleles distribution of nine DISC1 related SNPs with schizophrenia in males

| Marker       | Genotype | p-value (2df) | HWE | Allele   | p-value (1df) | OR (95% CI) |
|--------------|----------|---------------|-----|----------|---------------|-------------|
| rs821597     | AA       | 24 (0.20)     | 0.891 | 0.401     | 0.711          | 1.06 (0.77–1.47) |
| Controls     |          | 34 (0.18)     |      |           |               |             |
| rs821616     | TT       | 2 (0.01)      | 0.867 | 0.822     | 0.613          | 0.89 (0.57–1.40) |
| Controls     |          | 4 (0.02)      |      |           |               |             |
| rs4658971    | TT       | 14 (0.11)     | 0.0014* | <0.0001  | 0.103         | 1.40 (0.93–2.08) |
| Patients     |          | 26 (0.20)     |      |           |               |             |
| Controls     |          | 4 (0.02)      |      |           |               |             |
| rs1538979    | TT       | 6 (0.04)      | 0.109 | 0.062     | 0.226          | 1.24 (0.88–1.75) |
| Patients     |          | 63 (0.46)     |      |           |               |             |
| Controls     |          | 13 (0.06)     |      |           |               |             |
| rs843979     | CC       | 17 (0.14)     | 0.746 | 0.403     | 0.738          | 1.03 (0.77)  |
| Patients     |          | 62 (0.52)     |      |           |               |             |
| Controls     |          | 30 (0.17)     |      |           |               |             |
| rs2812385    | GG       | 30 (0.21)     | 0.271 | 0.760     | 0.103          | 1.29 (0.95–1.74) |
| Patients     |          | 69 (0.48)     |      |           |               |             |
| Controls     |          | 34 (0.16)     |      |           |               |             |
| rs1407599    | TT       | 2 (0.014)     | 0.959 | 0.867     | 0.826          | 0.95 (0.61–1.48) |
| Patients     |          | 32 (0.22)     |      |           |               |             |
| Controls     |          | 3 (0.013)     |      |           |               |             |
| rs4658900    | TT       | 26 (0.18)     | 0.543 | 0.040     | 0.553          | 0.91 (0.68–1.23) |
| Patients     |          | 83 (0.58)     |      |           |               |             |
| Controls     |          | 52 (0.23)     |      |           |               |             |
| rs2509382    | CC       | 2 (0.015)     | 0.0304* | 0.674  | 0.0082*       | 2.16 (1.22–3.81) |
| Patients     |          | 25 (0.18)     |      |           |               |             |
| Controls     |          | 1 (0.005)     |      |           |               |             |

\( \chi^2 \)-test, statistical significant at \(^*\) \( p<0.05 \). SNP: single nucleotide polymorphism
DISC1 and Schizophrenia

In Asia, there have been studies that showed an encouraging association between the DISC1 gene and schizophrenia among the Han Chinese.\textsuperscript{16,17} A study among the Japanese\textsuperscript{18} however, showed that DISC1 gene variants had a negative role in schizophrenia. Nevertheless, a study of the Korean population also indicated that DISC1 variation is associated with the poor-concentration phenotype of schizophrenia.\textsuperscript{19} The two SNPs (rs821616 and rs821597) that were reportedly associated with schizophrenia\textsuperscript{17} showed no such association in our cohort of patients. However, this study found two SNPs (rs4658971 and rs1538979) to be significantly associated with schizophrenia in terms of both genotypes and allelic distribution. The effect of these variations in DISC1 gene expression are unknown because both variations are located within introns of the DISC1 gene. There is no evidence suggesting the effects of these two SNP on gene splicing. Interestingly, a meta-analysis of more than five case-control association studies performed in Europe found that rs1538979 was a susceptibility marker for schizophrenia.\textsuperscript{20} However, an upgrade meta-analysis studies\textsuperscript{3} has refuted the association of the common variants of DISC1 with schizophrenia.

A recent meta-analysis of several genome-wide studies, including the publicly available data of the Genetic Association Information Network (GAIN) (www.genome.org), identified rs2509382 as having an association with schizophrenia among males.\textsuperscript{21} Our study found that rs2509382 caused a similarly significant risk of schizophrenia among the males \(p=0.0082, \text{OR}=2.16 (0.12–3.81)\). This finding supported the possibility of a sex-specific genetic predisposition toward the develop-
ment of schizophrenia. Interestingly, rs2509382 is located at 11q14.3 near the non-coding and brain expressed DISC1Fusion Partner 1 (DISC1FP1) gene\(^27\) that represents the partner of translocation region of the famously described DISC1 (1q42.1) translocation. The DISC1FP1 region association with schizophrenia is not well explored although a third party analysis on the Psychiatric Genetics Consortium Genome-Wide Association Study (GWAS) data has reported a weak association.\(^28\)

The exploration of the association of other variants within the 11q14.3 location with schizophrenia may provide further information regarding the relationship between this genetic location and schizophrenia.

Our study analyzed both Malay and Chinese subjects as one cohort of participants. This was based on the assumption that the genetic variations among the Malay and Chinese populations were relatively similar, as suggested by the Singapore Genome Variation Project,\(^29\) which was performed in a neighbouring country, Singapore. Additionally, evidence from a study on the genetic structure of populations found that all sub-ethnic groups of the Peninsular Malay are clustered together with the Chinese population and that these clusters are quite separated from the Yoruba, Indians, and Malaysian Indigenous Negritos.\(^30\)

A meta-analysis was performed to improve the statistical power of this project by combining data from several studies performed on the Chinese population. Combining allele frequency results from three studies, we found that there was no significant evidence of an association between SNPs rs821616, rs843979, and rs4658971 and schizophrenia. However, the combined data showed that rs821597 carries the risk of susceptibility to schizophrenia. The current genetic evidence further supports the important role of the DISC1 gene in the development of schizophrenia, and the association seems to affect both Caucasian and Asian populations.

Several functional and cell biology studies have supported the role of DISC1 in schizophrenia. DISC1 product interacts with other proteins implicated in the central nervous system development.\(^10,31\) In-\textit{vitro}, DISC1 has been shown to have a multifunctional protein whose truncation may lead to the disruption of intracellular transport, neurite growth, and neuronal migration.\(^28\) In addition to neurodevelopment, other studies also revealed the functional roles of DISC1 in tran-

---

| Study name | Odds ratio | Lower limit | Upper limit | Z-value | p-value |
|------------|------------|-------------|-------------|---------|---------|
| Qu et al.\(^{17}\) (2007) | 1.420 | 1.138 | 1.774 | 3.085 | 0.002 |
| Hotta et al.\(^{24}\) (2011) | 1.062 | 0.898 | 1.255 | 0.703 | 0.482 |
| Current | 1.003 | 0.773 | 1.302 | 0.025 | 0.980 |
| Qu et al.\(^{17}\) (2007) | 1.140 | 1.012 | 1.284 | 2.159 | 0.031 |

---

| Study name | Odds ratio | Lower limit | Upper limit | Z-value | p-value |
|------------|------------|-------------|-------------|---------|---------|
| Qu et al.\(^{17}\) (2007) | 1.608 | 1.150 | 2.248 | 2.776 | 0.006 |
| Chen et al.\(^{16}\) (2007) | 1.104 | 0.871 | 1.399 | 0.820 | 0.412 |
| Current | 0.940 | 0.650 | 1.361 | -0.327 | 0.744 |
| Current | 1.040 | 0.903 | 1.199 | 0.550 | 0.582 |

---

| Study name | Odds ratio | Lower limit | Upper limit | Z-value | p-value |
|------------|------------|-------------|-------------|---------|---------|
| Qu et al.\(^{17}\) (2007) | 1.010 | 0.764 | 1.335 | 0.067 | 0.946 |
| Current | 1.435 | 1.035 | 1.990 | 2.166 | 0.030 |
| Current | 1.171 | 0.947 | 1.448 | 1.458 | 0.145 |

---

| Study name | Odds ratio | Lower limit | Upper limit | Z-value | p-value |
|------------|------------|-------------|-------------|---------|---------|
| Qu et al.\(^{17}\) (2007) | 1.255 | 0.995 | 1.582 | 1.915 | 0.056 |
| Current | 1.114 | 0.850 | 1.461 | 0.783 | 0.434 |
| Current | 1.193 | 1.000 | 1.423 | 1.963 | 0.050 |

---

Figure 2. Meta-analysis findings of rs821597 (A), rs821616 (B), rs4658971 (C) and rs843979 (D).
scripion and cAMP signalling\textsuperscript{12} and in the presynaptic activity of glutamate release.\textsuperscript{13} The positive findings regarding genetic predisposition found in the current study, combined with the implicated proteomic pathways and functional roles of DISC1 that were reported by various studies, give additional support to the importance of DISC1 in the pathology of schizophrenia.

Our study has several limitations. The case-control approach used in this study is prone to false positive and false negative associations, especially due to undeclared population stratifications. The Malay ethnic group is loosely defined based on Article 160 of the Constitution of Malaysia. Therefore, a Malay is defined as a Malaysian citizen born to a Malaysian citizen who professes to be a Muslim, habitually speaks the Malay language, and adheres to Malay customs. To reduce this bias, the Malay participants were identified based on the ethnic identification of the previous two generations. Because schizophrenia is a disease with complex clinical presentations, a general association based on either having or not having the disease, as was used by the current study, may be too simplistic to unravel the complex nature of the disease with regard to genetic predisposition.

In conclusion, the current study found two SNPs within the DISC1 gene that predispose individuals to schizophrenia within the Malaysian population. Additionally, we found an association of rs2509382 with schizophrenia among male participants, thus reinforcing previous genome-wide meta-analysis findings regarding this SNP in a male Caucasian population.\textsuperscript{13} Additional studies that investigate the interaction between genetic predisposition and the clinical and social features of schizophrenia could elucidate the role of this gene in the clinical pathology of schizophrenia.

Acknowledgments

Staff and patients of Psychiatry Clinic, Tengku Ampuan Afzan Hospital, Kuantan, Pahang & staff of Molecular Research Laboratory, Kulliyyah of Medicine, IUM. This study was supported by ERGS Grant (ERGS-13-027-0060) by the Ministry of Education, Malaysia. The authors have no conflicts of interest that are directly relevant to the content of this manuscript. The study is funded by the ERGS Grant (13-027-0060) provided by the Ministry of Education, Malaysia.

REFERENCES

1. St Clair D, Blackwood D, Muir W, Carothers A, Walker M, Spowart G, et al. Association within a family of a balanced autosomal translocation with major mental illness. Lancet 1998;336:13-16.
2. Schumacher J, Laje G, Abos Jamra R, Becker T, Muhleisen TW, Vasiilescu G, et al. The DISC locus and schizophrenia: evidence from a association study in a central European sample and from a meta-analysis across different European populations. Hum Mol Genet 2009;18:2719-2727.
3. Mathieson I, Munafo MR Flint J. Meta-analysis indicates that common variants at the DISC1 locus are not associated with schizophrenia. Mol Psychiatry 2012;17:634-641.
4. Ekelund J, Howatta I, Parker A, Paunio T, Varilo T, Martin R, et al. Chromosome 1 loci in Finnish schizophrenia families. Hum Mol Genet 2001;10:1611-1617.
5. Saetre P, Agartz I, De Francisci A, Lundmark P, Diurowoc S, Kahlwer A, et al. Association between a disrupted-in-schizophrenia 1 (DISC1) single nucleotide polymorphism and schizophrenia in a combined Scandinavian case-control sample. Schizophr Res 2008;106:237-241.
6. Hodgkinson CA, Goldman D, Jaeger I, Persaud S, Kane JM, Lipsky RH, et al. Disrupted in schizophrenia 1 (DISC1): association with schizophrenia, schizoaffective disorder, and bipolar disorder. Am J Hum Genet 2004;75:862-872.
7. Sullivan PF. Questions about DISC1 as a genetic risk factor for schizophrenia. Mol Psychiatry 2013;18:1050-1052.
8. Chubb JE, Bradshaw NJ, Soares DC, Porteous DJ, Millar JK. The DISC locus in psychotic illness. Mol Psychiatry 2008;13:36-64.
9. Porteous DJ, Thomson P, Brandon NJ, Millar JK. The genetics and biology of DISC1—an emerging role in psychosis and cognition. Biol Psychiatry 2006;60:123-131.
10. Ishizuka K, Paek M, Kamiya A, Sawa A. A review of Disrupted-In-Schizophrenia-1 (DISC1): neurodevelopment, cognition, and mental conditions. Biol Psychiatry 2006;59:1189-1197.
11. Hayashi-Takagi A, Takaki M, Graziane N, Seshadri S, Murdoch H, Dunlop AJ, et al. Disrupted-in-Schizophrenia-1 (DISC1) regulates spines of the glutamate synapse via Rac1. Nat Neurosci 2010;13:327-332.
12. Lee FH, Fadel MP, Preston-Maher K, Cordes SP, Lapcote SJ, Price DJ, et al. Disc1 point mutations in mice affect development of the cerebral cortex. J Neurosci 2011;31:3197-3206.
13. Carless MA, Glahn DC, Johnson MP, Curran JE, Bozaoglu K, Dyer TD, et al. Impact of DISC1 variation on neuroanatomical and neurocognitive phenotypes. Mol Psychiatry 2011;16:1096-1104, 1063.
14. Cannon TD, Hennah W, van Erp TG, Thompson PM, Lonnqvist J, Huttunen M, et al. Association of DISC1/TRAX haplotypes with schizophrenia, reduced prefrontal gray matter, and impaired short- and long-term memory. Arch Gen Psychiatry 2005;62:1205-1213.
15. Gamo NJ, Duque A, Paspadis CD, Kata A, Fine R, Boven L, et al. Role of disrupted in schizophrenia 1 (DISC1) in stress-induced prefrontal cognitive dysfunction. Transl Psychiatry 2013;3:e328.
16. Chen QY, Chen Q, Feng GT, Lindpaintner K, Wang LJ, Chen ZX, et al. Case-control association study of Disrupted-in-Schizophrenia-1 (DISC1) gene and schizophrenia in the Chinese population. J Psychiatr Res 2007;41:428-434.
17. Qu M, Tang F, Yue W, Ruan Y, Lu T, Liu Z, et al. Positive association of the Disrupted-in-Schizophrenia-1 gene (DISC1) with schizophrenia in the Chinese Han population. Am J Med Genet B Neuropsychiatr Genet 2007;144B:266-270.
18. Bae JS, Kim JY, Park BL, Cheong HS, Kim JH, Shin IG, et al. Lack of association between DISC1 polymorphisms and risk of schizophrenia in a Korean population. Psychiatry Res 2013;208:189-190.
19. Kim HJ, Park HJ, Jung KH, Ban JY, Ra J, Kim JW, et al. Association study of polymorphisms between DISC1 and schizophrenia in a Korean population. Neurosci Lett 2008;430:60-63.
20. Zhang X, Toguchi M, Ohashi J, Maeda K, Kato T, Okazaki Y, et al. Association study of the DISC1/TRAX locus with schizophrenia in a Japanese population. Schizophr Res 2005;79:175-180.
21. Debono R, Topless R, Markie D, Black MA Merriman TR. Analysis of the DISC1 translocation partner (11q14.3) in genetic risk of schizophrenia. J Med Genet 2001;38:77-81.
22. Porteous DJ, Thomson P, Brandon NJ, Millar JK. The genetics and biology of DISC1—an emerging role in psychosis and cognition. Biol Psychiatry 2006;60:123-131.
23. Chen QY, Chen Q, Feng GT, Lindpaintner K, Wang LJ, Chen ZX, et al. Case-control association study of Disrupted-in-Schizophrenia-1 (DISC1) gene and schizophrenia in the Chinese population. J Psychiatr Res 2007;41:428-434.
24. Qu M, Tang F, Yue W, Ruan Y, Lu T, Liu Z, et al. Positive association of the Disrupted-in-Schizophrenia-1 gene (DISC1) with schizophrenia in the Chinese Han population. Am J Med Genet B Neuropsychiatr Genet 2007;144B:266-270.
and Japanese patients with treatment-resistant schizophrenia (TRS). Prog Neuropsychopharmacol Biol Psychiatry 2011;35:636-639.
25. Borenstein M, Hedges L, Higgins J, Rothstein H. Comprehensive Meta-analysis, version 2. Englewood NJ: Biostat Inc.; 2005.
26. Allen NC, Bagade S, McQueen MB, Ioannidis JPA, Kavvoura FK, Khoury MJ, et al. Systematic Meta-Analyses and Field Synopsis of Genetic Association Studies in Schizophrenia: The SzGene Database. Nat Genet 2008;40:827-834.
27. Zhou X, Chen Q, Schaukowitch K, Kelsey JR Geyer MA. Insoluble DISC1-Boyman fusion proteins generated by DISC1 translocation. Mol Psychiatry 2010;15:669-672.
28. Thomson PA, Malavasi E, Grunevald E, Soares DC, Borkowska M, Millar JK. DISC1 genetics, biology and psychiatric illness. Front Biol (Beijing) 2013;8:1-31.
29. Teo YY, Sim X, Ong RT, Tan AK, Chen J, Tantoso E, et al. Singapore Genome Variation Project: a haplotype map of three Southeast Asian populations. Genome Res 2009;19:2154-2162.
30. Hatin WI, Nur-Shafawati AR, Zahri MK, Xu S, Jin L, Tan SG, et al. Population genetic structure of peninsular Malaysia Malay sub-ethnic groups. PLoS One 2011;6:e18312.
31. Morris JA, Kandpal G, Ma L, Austin CP. DISC1 (Disrupted-In-Schizophrenia 1) is a centrosome-associated protein that interacts with MAP1A, MIPT3, ATF4/5 and NUDHEL: regulation and loss of interaction with mutation. Hum Mol Genet 2003;12:1591-1608.
32. Soda T, Frank C, Ishizuka K, Baccarella A, Park YU, Flood Z, et al. DISC1-ATF4 transcriptional repression complex: dual regulation of the cAMP-PDE4 cascade by DISC1. Mol Psychiatry 2013;18:898-908.
33. Maher BJ, LoTurco JJ. Disrupted-in-schizophrenia (DISC1) functions presynaptically at glutamatergic synapses. PLoS One 2012;7:e34053.