



Genetic variation in *BDNF* is associated with antipsychotic treatment resistance in patients with schizophrenia

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ABSTRACT

Objective: Antipsychotic drugs are the mainstay of treatment for schizophrenia. However, a substantial proportion of patients are poorly responsive or resistant to first-line treatments, and clozapine treatment is often indicated. Therefore, we and others have used clozapine treatment as a proxy phenotype for antipsychotic treatment resistance in pharmacogenetic studies. In the present study, we utilized this phenotype to test previously-identified candidate genes for antipsychotic treatment response.

Method: We assessed 89 Caucasian schizophrenia patients clinically assigned to clozapine treatment versus 190 Caucasian patients that were not selected for clozapine treatment. We conducted gene-based association tests on a set of 74 relevant candidate genes nominated in the CATIE pharmacogenetic study (Need et al., 2009), using the GATES procedure (Li et al., 2011).

Results: After correcting for multiple testing in the gene-based association test, the gene for brain derived neurotrophic factor (*BDNF*) was significantly associated with treatment resistance. The top single nucleotide polymorphisms (SNPs) in *BDNF* included rs11030104 (OR=2.57), rs10501087 (OR=2.19) and rs6265 (Val66Met) (OR=2.08). These SNPs appear to be in high linkage disequilibrium with each other.

Conclusion: *BDNF* appears to have a strong association with antipsychotic treatment resistance. Future studies are needed to replicate this finding and further elucidate the biological pathways underlying the association between *BDNF* and antipsychotic drug response.

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1. Introduction

Although antipsychotic drugs have demonstrated clinical efficacy in treating schizophrenia, many patients are still partially or fully unresponsive to treatment and are classified as treatment resistant (Conley and Kelly, 2001). Treatment resistant schizophrenia is poorly understood and difficult to predict (Suzuki et al., 2011), and has significant human and economic repercussions. Although there are several definitions of treatment resistant schizophrenia (Kane et al., 1988; Meltzer, 1992; Bondolfi et al., 1998; Conley and Kelly, 2001), one clinically relevant marker of treatment resistance is the decision to prescribe clozapine. Under current guidelines, clozapine should only be used after two failed trials of other antipsychotic agents (Kane et al., 2003; Kreyenbuhl et al., 2010), due to risk of substantial side effects. Nevertheless, clozapine is the only antipsychotic drug that has been shown to be superior in treating refractory schizophrenia (Meltzer,

1997; McEvoy et al., 2006). Thus, clozapine utilization in the clinical setting may be considered as a proxy measure of antipsychotic treatment resistance. A predictor of treatment resistance would be clinically useful because currently, many patients receive multiple trials of different medications before clozapine is prescribed.

A few small pharmacogenetic studies have examined genetic predictors of antipsychotic treatment resistance (Kohlrusch et al., 2008; Jia et al., 2011). To date, the only large-scale study of antipsychotic treatment response was drawn from the Clinical Antipsychotic Trials of Intervention Effectiveness (CATIE) (Lieberman et al., 2005). In the CATIE data, 453 SNPs in 74 genes were found to be nominally associated with discontinuation from assigned antipsychotics, a proxy for treatment failure (Need et al., 2009). These included genes coding for relevant neurotransmitter systems such as dopamine (e.g., *DRD1*), serotonin (e.g., *HTR2A*), and glutamate (*GAD1*), as well as other neuronal proteins such as *BDNF*. While none of these SNPs were statistically significant after correction for multiple testing, no replication dataset was available in the report to permit the further interrogation of these candidates.

The present study utilized the results of the CATIE pharmacogenetic study as a discovery dataset for assessment of antipsychotic treatment

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resistance in our own dataset, in which clinically assigned clozapine therapy was a proxy for treatment-resistance. We conducted a gene-based association test comparing patients on clozapine therapy to non-clozapine patients. Because genes are functional units of the DNA sequence, and to reduce multiple testing burden, we chose to conduct gene-based association tests instead of SNP-based analysis. In addition, gene-based association tests are less likely to be affected by heterogeneity in linkage disequilibrium (LD) structure and allele frequencies across populations (Neale and Sham, 2004).

2. Methods

The sample included 89 schizophrenia patients clinically assigned to clozapine treatment and 190 patients that were not selected for clozapine treatment, as part of an earlier case-control GWAS study (Lencz et al., 2007). All patients were Caucasians. Clozapine vs. non-clozapine assignment was the phenotype used in the present study. Patient recruitment and assessment procedures have been described previously (Lencz et al., 2007). All patients were psychiatrically stable on assigned treatment at the time of study entry. All clozapine patients had failed at least two antipsychotics prior to initiating clozapine therapy. Table 1 presents the demographic characteristics of the sample.

Peripheral blood samples were taken from each patient and DNA was extracted from lymphocytes. Details of sample processing have been reported elsewhere (Lencz et al., 2007). All patients were genotyped on the Affymetrix 500k platform. After standard procedures for genotype calls and quality control, 364,961 high quality SNPs were utilized in subsequent analyses (Lencz et al., 2007).

SNP-based association tests were conducted using SVS software (Golden Helix Inc., Bozeman, MT) in an additive model (correlation-trend test), controlling for population stratification with principal component analysis. The results were then imported to the GKK program to run gene-based association tests using the GATES procedure (Li et al., 2011). At this stage of analysis, a set of 74 candidate genes, which were associated with all-cause discontinuation at the $p < 0.05$ level in the CATIE study (Need et al., 2009), was analyzed for association to clozapine assignment in our sample.

3. Results

Seventy-four genes that were nominally linked with treatment discontinuation from the CATIE study were tested in our sample using the GATES procedure. After correcting for multiple testing, only the brain-derived neurotrophic factor gene (*BDNF*) was associated with antipsychotic treatment resistance ($p = 0.0002$, combined p value at gene level from the GATES procedure). Ten out of 34 SNPs in *BDNF* were nominally ($p < 0.05$) associated with clozapine therapy. These associations likely represent a single signal due to high LD among the SNPs (Fig. 1). The top SNPs in *BDNF* included rs11030104 (OR = 2.57, 95% Confidence Interval [CI]: 1.63–4.04, $p = 0.00004$); rs10501087 (OR = 2.19, CI: 1.41–3.38, $p = 0.0006$), and rs6265 (Val66Met) (OR = 2.08, CI: 1.33–3.25, $p = 0.0008$). Fig. 2 shows the percentage of clozapine patients in each genotype for each SNP. At each of these loci, carrying the minor allele seems to confer an increased

risk of antipsychotic treatment resistance. Moreover, this increased risk appears to be dose dependent. The effect sizes seem to be large when comparing different genotype groups. Compared to the major allele homozygotes, the minor allele homozygotes were more likely to be on clozapine therapy with an odds ratio of 8.79 (CI: 2.22–34.84), 4.45 (CI: 1.34–14.82), and 4.37 (CI: 1.18–16.17) for the three SNPs, respectively.

4. Discussion

We conducted a gene-based association study on genes suggested by the CATIE study (Need et al., 2009) to be involved in treatment resistance, using clozapine therapy as a proxy for treatment resistance. *BDNF*, the gene coding for the brain-derived neurotrophic factor, appeared to be significantly associated with antipsychotic drug resistance with a moderate to large effect size. Specifically, we found that minor alleles of several *BDNF* SNPs were significantly associated with clinically-assigned clozapine therapy. There seems to be a dose-response relationship between number of minor alleles and likelihood of treatment resistance (Fig. 2). The minor allele homozygotes were 4–8 times more likely to be resistant to antipsychotic treatment than the major allele homozygotes.

These data are consistent with prior studies implicating *BDNF* and antipsychotic treatment response. An early study (Krebs et al., 2000) found that the short allele of the *BDNF* dinucleotide repeat polymorphism (168 bp) was more prevalent among patients refractory to several different trials of neuroleptics compared to patients who responded to neuroleptics. In contrast, the long alleles (172–176 bp) were more prevalent among antipsychotic responders (OR = 2.7). This polymorphism is located 1040 bp upstream of the transcription initiation site, and may play a role in regulating *BDNF* gene expression. A more recent study also reported that the short allele was associated with poor response to risperidone (Xu et al., 2010), further implicating *BDNF* in antipsychotic response. Interestingly, this microsatellite and the Val66Met polymorphisms are separated by only 1280 bps and the two markers are in linkage disequilibrium ($D' > 0.70$) (Neves-Pereira et al., 2002).

BDNF interacts with multiple neurotransmitters, including dopamine and serotonin, that are major targets for antipsychotic drugs (Buckley et al., 2011). The Val66Met polymorphism (rs6265) is a frequently studied SNP, with the G to A polymorphism resulting in a valine to methionine substitution at codon 66. This alters the intracellular trafficking and packaging of pro*BDNF*, leading to reduced synaptic plasticity (Egan et al., 2003). The Met allele is associated with decreased performance on memory tests (Ho et al., 2007; Goldberg et al., 2008) and smaller hippocampal volume (Szeszko et al., 2005). Because antipsychotics work partially by enhancing dentate gyrus glutamate transmission or through modulating long-term potentiation in other areas of hippocampus (Miyamoto et al., 2005; Tammimga et al., 2010), it would be plausible that the Met allele-associated smaller hippocampus volume leaves little room for this mechanism and therefore results in treatment resistance.

There are limitations in the present study. The phenotype, clozapine assignment, differed from that used in the CATIE study (discontinuation); neither is a perfect indicator of antipsychotic resistance. However, there is no consensus on the best definition of treatment resistance. Clozapine therapy and treatment discontinuation in chronic patients are clinically meaningful representations of treatment failure. Secondly, the significant *BDNF* SNPs found in CATIE were not directly genotyped in our data due to different genotyping platforms. Moreover, the CATIE sample was heterogeneous and included several racial groups. In contrast, our sample consisted of all Caucasians. Both of these issues were partially resolved, however, by our use of gene-based association tests. This approach is less affected by heterogeneity in LD structure and allele frequencies across populations. Finally, our sample size was relatively small. Therefore, although the finding was consistent with previous evidence, it needs to be replicated in larger samples.

Table 1
Demographic characteristics of the study sample.

	Clozapine (n = 89)	Non-clozapine (n = 190)	Total (n = 279)	p
Age (mean ± SD)	37.59 ± 9.95	39.31 ± 10.52	38.8 ± 10.4	0.20
% Female	29.20%	37.60%	35.50%	0.17
Age of onset (mean ± SD)	20.16 ± 5.06	22.76 ± 10.07	21.87 ± 8.72	0.03
% Positive family history	28.40%	26.92%	23.7%	0.81

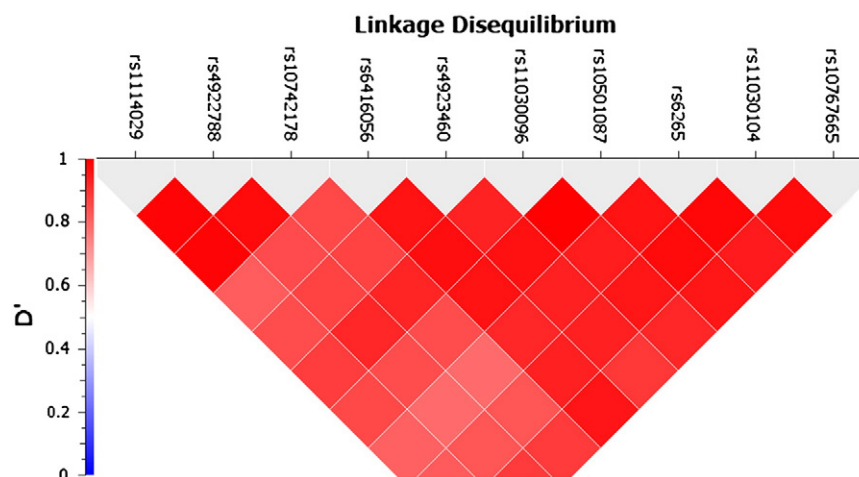


Fig. 1. Linkage disequilibrium plot (D') for the significant SNPs in *BDNF*. The plot was generated using PLINK (v1.07) software using HapMap release 3 data. Bright red color indicates high D' , which represents high correlation among different SNPs.

In summary, with a gene-based association test approach, *BDNF* was associated with antipsychotic treatment resistance using clinically assigned clozapine therapy as a proxy. Future studies are needed to replicate the finding and further elucidate the biological pathways

underlying the association between *BDNF* and antipsychotic drug response.

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Contributors

Dr. Jian-Ping Zhang designed the study, conducted data analysis, managed literature search, and wrote a draft of the manuscript.

Dr. Todd Lencz designed the study, participated in data analysis, and revised the manuscript.

Dr. Stephen Geisler recruited subjects and provided clinical data.

Dr. Pamela DeRosse conducted clinical interview and managed the database.

Dr. Evelyn J. Bromet recruited subjects and provided clinical data.

Dr. Anil K. Malhotra provided the general hypothesis, designed the study, and revised the manuscript.

All authors contributed to and have approved the final manuscript.

Conflict of interest

Dr. Zhang, Dr. Geisler, Dr. DeRosse, and Dr. Bromet reported no actual or potential conflict of interest.

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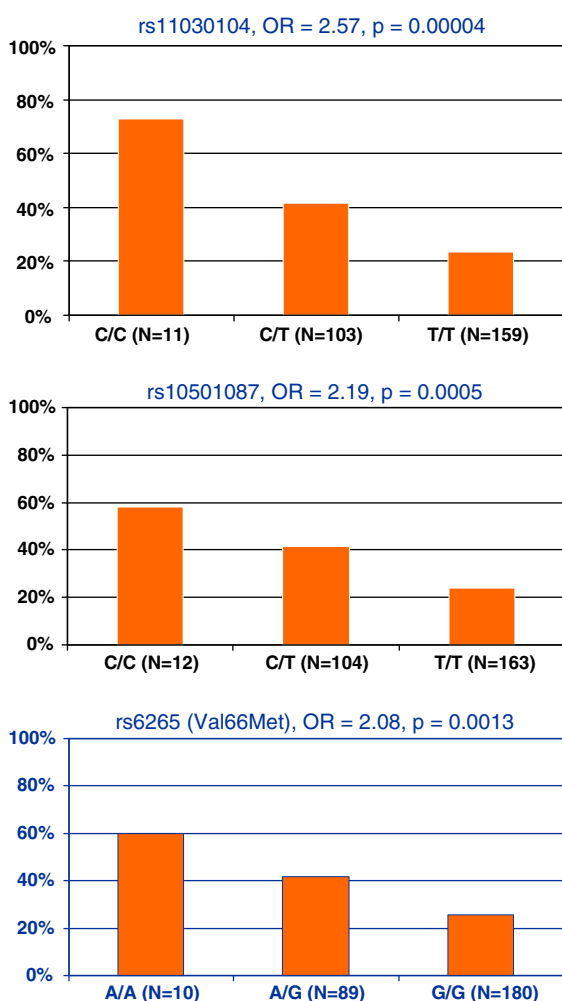


Fig. 2. Percentage of patients on clozapine therapy in each genotype group for three *BDNF* SNPs (odds ratios (OR) are for each additional minor allele, compared to no minor allele).

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