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Recommended Citation

Gonzalez, S., Xu, C., Ramirez, M., Zavala, J., Armas, R., Contreras, S. A., Contreras, J., Dassori, A., Leach, R. J., Flores, D., Jerez, A., Raventós, H., Ontiveros, A., Nicolini, H., & Escamilla, M. (2013). Suggestive evidence for association between L-type voltage-gated calcium channel (CACNA1C) gene haplotypes and bipolar disorder in Latinos: a family-based association study. *Bipolar disorders*, 15(2), 206–214. <https://doi.org/10.1111/bdi.12041>

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Published in final edited form as:

Bipolar Disord. 2013 March ; 15(2): 206–214. doi:10.1111/bdi.12041.

Suggestive evidence for association between L-type voltage-gated calcium channel (CACNA1C) gene haplotypes and bipolar disorder in Latinos: a family-based association study

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Abstract

Objectives—Through recent genome-wide association studies (GWAS), several groups have reported significant association between variants in the alpha 1C subunit of the L-type voltage-gated calcium channel (CACNA1C) and bipolar disorder (BP) in European and European-American cohorts. We performed a family-based association study to determine whether CACNA1C is associated with BP in the Latino population.

Methods—This study consisted of 913 individuals from 215 Latino pedigrees recruited from the United States, Mexico, Guatemala, and Costa Rica. The Illumina GoldenGate Genotyping Assay was used to genotype 58 single-nucleotide polymorphisms (SNPs) that spanned a 602.9 kb region encompassing the *CACNA1C* gene including two SNPs (rs7297582 and rs1006737) previously

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Disclosures

The authors of this paper do not have any commercial associations that might pose a conflict of interest in connection with this manuscript.

shown to associate with BP. Individual SNP and haplotype association analyses were performed using Family-Based Association Test (version 2.0.3) and Haploview (version 4.2) software.

Results—An eight-locus haplotype block that included these two markers showed significant association with BP (global marker permuted $p = 0.0018$) in the Latino population. For individual SNPs, this sample had insufficient power (10%) to detect associations with SNPs with minor effect (odds ratio = 1.15).

Conclusions—Although we were not able to replicate findings of association between individual *CACNA1C* SNPs rs7297582 and rs1006737 and BP, we were able to replicate the GWAS signal reported for *CACNA1C* through a haplotype analysis that encompassed these previously reported significant SNPs. These results provide additional evidence that *CACNA1C* is associated with BP and provides the first evidence that variations in this gene might play a role in the pathogenesis of this disorder in the Latino population.

Keywords

bipolar disorder; calcium channels; genetic association studies; haplotypes; Hispanic Americans; L-type; pedigree; polymorphism; single nucleotide

Bipolar disorder (BP) is an often chronic and debilitating mental disorder affecting approximately 1–4% of the population (1, 2). Family, twin, and adoption studies on BP have shown that there is a substantial genetic component (3, 4), as BP has one of the highest heritability rates of all known psychiatric disorders (1). Over the past several years, genome-wide association studies (GWAS) have produced several successes in identifying genetic variants that contribute to genetically complex human disorders (5–16). Recently, a few research groups have performed independent GWAS of BP (11, 17–22), with little overlap amongst the single-nucleotide polymorphisms (SNPs) most strongly associated with BP (23). However, a meta-analysis of three GWAS of BP identified a SNP within the calcium channel, voltage-dependent, L-type, alpha 1C subunit (*CACNA1C*) gene, rs1006737, as being strongly associated with the risk of BP (21). A subsequent meta-analysis combining data from GWAS of BP and major depressive disorder (MDD) reported two SNPs in a 10.5-kb region within the *CACNA1C* gene which exceeded a genome-wide significance level (rs1006737 and rs7297582) (20).

There has been increasing evidence in recent years that alterations in calcium signaling are involved in the pathophysiology of BP (32). The *CACNA1C* gene located on chromosome 12p13.3 encodes the major constituent of the brain L-type voltage-gated calcium channels that are critical to dendritic calcium influx in response to synaptic activity (33, 34). Reports have associated SNPs in *CACNA1C* with an increase in grey matter density in the amygdala and hypothalamus (35), total grey matter (36), and brainstem volume (37). Neuroimaging studies of healthy individuals report that the rs1006737 risk allele modulates brain function during tasks of reward and emotional processing (38, 39), verbal fluency (40), attention networks (41), and episodic and prefrontal cortex (PFC) working memory (38, 42). The *CACNA1C* rs1006737 polymorphism also impacts ventrolateral PFC activation during fear processing in BP carriers of the risk allele but not their unaffected relatives (43). In addition, studies have shown that carriers of the rs1006737 variant show a pronounced effect on measures of psychosocial functioning, including significantly higher scores on tests for depression, anxiety, interpersonal sensitivity, obsessive-compulsive thoughts, and neuroticism, even in the absence of disease (44). These studies suggest that calcium channel dysfunction may contribute in part to the genetic etiology of BP through alterations in the functional activity of brain circuitries.

Although there is mounting evidence that *CACNA1C* is involved in the genetic etiology of BP, the genetic studies performed to date have been almost exclusively centered on populations with European ancestry. The replication of significant GWAS findings in diverse populations is needed in order to corroborate initial findings from European studies and characterize their genetic contribution to BP in other ethnicities. In order to address this issue, we have designed a family-based association study to try and validate the GWAS-significant *CACNA1C* SNPs in a large set of families with Latin American (Latino) ancestry.

Materials and methods

Study samples

The sample consisted of 215 Latino pedigrees in which BP probands and extended family members were recruited from the United States (Texas, New Mexico, California), Mexico (Mexico City, Monterrey), Costa Rica, and Guatemala. Inclusion criteria required a proband with a bipolar disorder type I (BP-I) diagnosis with at least one sibling with a clinical diagnosis of BP-I or schizoaffective BP (SABP), and a minimum of two additional first-degree relatives willing to participate. Extended pedigrees were collected when possible to include other family members with a history of affective or psychotic disorders. The sample consisted of 157 case-parent trios and 258 affected subjects with one parent genotyped. All subjects reported ancestry from Mexico or Central America. Family structure based on number of affected family members and country of origin is listed in Table 1. Previous genetic structure analysis has shown that these populations are closely related, with high levels of admixture consisting of three major ancestral populations (Caucasian, Native American, and African) (45).

Subjects signed Institutional Review Board (IRB)-approved written informed consent forms prior to enrolling in the study.

The procedures were approved by the IRB of Texas Tech University Health Science Center and respective IRBs in each participating site and country, and the study was performed in accordance with the Helsinki Declaration of 1975.

All study participants were diagnosed using DSM-IV criteria (46), by a best-estimation consensus procedure using the Diagnostic Interview for Genetic Studies (DIGS) (47), Family Interview for Genetic Studies (FIGS) (48), and available psychiatric records as previously described (49). The 215 Hispanic pedigrees under study contained 929 individuals: 466 of whom were diagnosed with BP-I and 16 of whom were diagnosed with SABP. Of the study participants with a diagnosis of BP-I or SABP, 200 were male and 282 were female.

Genotyping

DNA was isolated from lymphoblastoid cell lines established and stored for each study participant at the National Institute of Mental Health (NIMH) Center for Collaborative Genetic Studies. SNP selection covering the *CACNA1C* gene was based on a tagging SNP approach ($r^2 = 0.9$) using the SNP browser™ Software version 4.0.1 based on data from the CEU (Utah residents with ancestry from northern and western Europe) HapMap population (50). Known *CACNA1C* variants rs1006737 (20, 21) and rs7297582 (20), which reached genome-wide significance in previous meta-analysis GWAS studies, were included in the panel. Additional proximal SNPs to these known variants were chosen from the National Center for Biotechnology Information (NCBI) dbSNP (51) based on criteria that SNPs were validated and had heterozygosity > 0.4 . A total of 58 SNPs were genotyped using a custom-

designed Illumina GoldenGate SNP assay according to the manufacturer's protocol (Illumina, San Diego, CA), blind to subject diagnosis and characteristics.

Genetic markers failing standard quality control criteria were excluded from further analysis: missing genotype > 0.1 , minor allele frequency < 0.01 , and exact Hardy–Weinberg equilibrium $p < 0.01$. Also excluded from the analysis were individuals with a missing genotype rate > 0.1 , or Mendelian errors > 2 . The discrepancy rate on 2% duplicate genotyping was < 0.002 . A total of 57 SNPs for 213 pedigrees (913 individuals) were retained for further statistical analysis.

Statistical analyses

Analyses of individual SNPs were completed with the Family-Based Association Test (FBAT) version 2.0.3 (52), which was used for testing association under the null hypothesis of 'no association and no linkage' using an additive genetic model. To thoroughly test the locus identified by the two individual SNPs previously reported to be in association with BP, we also identified a haplotype block which included those SNPs and analyzed the haplotypes for association. Haploview version 4.2 (53) was used to visualize linkage disequilibrium (LD) relationships between all genotyped variants (57 SNPs) within and surrounding the *CACNA1C* region (54) and to construct LD blocks following the D method described by Gabriel et al. (55). Haplotype analyses were performed using the Haplotype-Based Association Testing (HBAT) assessment in the FBAT program. Haplotype-specific and global-permuted p-values were calculated for individual haplotype tests conducted under *biallelic* mode in haplotype FBAT, assuming an additive genetic model. In order to maximize power in the statistical analyses, permutation procedures were implemented to calculate the empirical p-values derived from 10,000 permutations. The significance level for all statistical tests was two-tailed ($p < 0.05$).

Power analyses were estimated using the transmission disequilibrium test (TDT) for discrete traits of the Genetic Power Calculator (<http://pngu.mgh.harvard.edu/~purcell/gpc/>) with $\alpha = 0.05$ and D of 0.9. The current family study design ($n = 215$) yielded an estimate of 0.65 [odds ratio (OR) = 1.75] and 0.82 (OR = 2.00) for a gene of moderate effect and a power estimate of 0.40 (OR = 1.5) and 0.10 (OR = 1.15) for a gene with minor effect. We calculated that our sample had reasonable power to detect a common risk variant, but would be insufficient to detect a susceptibility locus with a minor effect as seen in genetic studies of the *CACNA1C* gene for BP. To make optimal use of our data, we performed haplotype-based tests to increase the power of the association study (55).

Results

The primary aim of this study was to test to see if findings of association of SNPs rs1006737 and rs7297582 in the *CACNA1C* gene with BP, as reported in a European ancestry cohort by Liu et al. (20), were valid in other populations, specifically in the Latino population. When tested individually, we found no significant association between these two SNPs and BP (rs7297582, $Z = 0.306$, $p = 0.760$; rs1006737, $Z = 0.725$, $p = 0.469$) in the Latino population studied.

Given the negative findings of association for SNPs rs1006737 and rs7297582, we wished to test whether SNPs in LD with these variants would associate with BP in our Latino cohort. We used a tagged-SNP approach to cover the *CACNA1C* gene in addition to several SNPs in close proximity to rs1006737 and rs7297582, assuming that these SNPs were not functional SNPs but rather variants in LD with true causal mutation. Twelve LD blocks were identified in our sample (Fig. 1). Haploblock 4 encompasses both rs1006737 and rs7297582 SNPs, previously shown to associate with BP. Allele frequencies and the results of FBAT

for single SNP analysis for the eight SNPs comprising Haploblock 4 are shown in Table 2. Univariate (single-marker) FBAT demonstrated no preferential transmission for any of the variant alleles tested (Table 2).

Since the use of haplotypes may provide greater statistical power in association studies of common genetic variation than can be obtained with individual SNPs (55), we conducted haplotype analyses focusing on Haploblock 4 which contained SNPs previously associated with BP. To determine whether any specific haplotype would confer a higher risk for BP, the specific and global-haplotype FBAT tests of association were performed. One common and three rare haplotypes displayed significant association under an additive model (Table 3). The p-values of the whole haplotype permutation test were also statistically significant (χ^2 sum, $p = 0.0018$) (Table 3). Of note, among these four disease-associated haplotypes, three (GGGCGGA, GGGCGAGA, and AAAGAGGA) showed protective effect with negative Z-scores and one showed a susceptibility effect with a positive Z-score (GAAGGAAG). Interestingly, the specific alleles previously shown to be associated with BP on SNPs rs1006737 (A) and rs7297582 (A), were also seen on the GAAGGAAG haplotype showing increased risk ($Z = 2.236$, $p = 0.031$). The common haplotype significantly associated with BP was present across all countries in our cohort, while the other three haplotypes were very rare.

Discussion

In the present study, we attempted to validate, in nearly 1,000 Latinos, the significant CACNA1C variants associated with BP identified in populations of predominantly European ancestry (11, 18, 20, 21). When analyzed individually, we were unable to replicate association of SNPs rs7297582 and rs1006737 with BP in the Latino population. However, these two previously identified SNPs have not been shown to be functionally significant within the CACNA1C gene and may only be in LD with functional variants at this locus. Moreover, allele frequencies of genotyped markers in strong LD with the two previously associated SNPs are highly variable across ancestral populations (hapmap.ncbi.nlm.nih.gov), suggesting the likelihood of a differential profile of CACNA1C variants with ethnicity (Table 4). Given the negative findings of our single locus tests compared to previous associations found in the Caucasian population, combined with the finding that haplotypes at this locus are associated with BP in the Latino population, the possibility remains that there might be additional functional variants in both the European and Latino ancestry populations that might still be in strong LD with rs7297582 and rs1006737.

It is known that in the presence of multiple tightly linked markers a haplotype test may be more powerful to detect association than corresponding single-locus tests (56). Haplotype-based analysis of the haploblock encompassing the two previously reported SNPs associated with BP provided significant evidence for association with BP in the Latino population. Our complementary LD block-based analysis revealed that an eight-locus haploblock (consisting of rs769087, rs1006737, rs2159100, rs4765905, rs2370413, rs7297582, rs758170, and rs1860002) provided stronger evidence of association than single-marker analysis. In affected individuals, three distinct haplotypes (GGGCGGA, GGGCGAGA, and AAAGAGGA) were under-represented and one haplotype (GAAGGAAG) was over-represented, giving rise to a significant permuted whole marker haplotypic association (χ^2 sum, $p = 0.0018$). We performed a conditional analysis to see if the association was driven by the two previously reported risk variants. We removed the two variants and performed the family-based haplotype association test with the remaining six variants under identical parameters outlined in the methods section. These results were very similar to our initial findings, in which one common and three rare haplotypes were significantly associated (see

Supplementary Table 1). The whole marker permutation test was also significant, indicating that the association is not driven by these two previously reported SNPs.

Since the fruition of this study, The Psychiatric GWAS Consortium (PGC) BP Working Group published their results of the most comprehensive GWAS of BP to date, including 16,731 samples and a replication sample of 46,918 individuals. They report a genome-wide significant SNP, rs4765913, in *CACNA1C* associated with BP (57). A combined analysis of PGC BP and PGC schizophrenia showed stronger association of this SNP compared to the BP GWAS alone (26, 57). The PGC significant variant falls between two haploblocks identified in our Latino population (Fig. 2). This SNP is approximately 6 kb downstream from the proximal marker of the associated haploblock and approximately 16 kb upstream of the next closest marker. Although we did not test SNP rs4765913 directly, we believe it is very likely, due to its proximity, to be in strong LD with the haploblock associated with BP in our Latino cohort, thus providing additional support for our findings.

Another recently published study reported a significant association of rs4765905 located within *CACNA1C* with BP and schizophrenic patients ($n = 16,373$) compared to control subjects ($n = 14,044$) (27). SNP rs4765905 was also genotyped in our current study and is nested in the LD block used in this association study. Taken as a whole, these results provide compelling evidence that this region of the *CACNA1C* gene plays a role in the pathogenesis of BP.

There are several limitations of the present study. First, compared to the previous studies in European ancestry populations, this was a smaller sample size. Larger sample sizes might be required in order to detect statistically significant association for single, common alleles in the Latino population. Secondly, this study only utilized published SNPs and did not use a sequencing approach to detect other potential SNPs or genetic variants within the gene which might be more directly causal for the disease. Finally, the combination of families from several countries may introduce heterogeneity into the analysis. However, if that were to be the case, this would diminish the chances of identifying association of gene variants. There are also some strengths to the present study: family-based association analyses were used, which prevent potential false positive results due to stratification in case-control analyses, and a narrow phenotype of carefully diagnosed BP-I and SABP cases were utilized as the affected phenotype of interest, drawn from families with multiplex BP cases.

In conclusion, these results provide the first evidence that genetic variations in *CACNA1C* might play a role in the pathogenesis of BP in the Latino population. We provide further support to previous case-control studies in Caucasian populations, strengthening the notion that genetic variation of *CACNA1C* influences the susceptibility to BP. Further investigations of the *CACNA1C* gene are thus warranted to increase our understanding of its role in the pathogenesis of BP. Careful replication with appropriate designs in independent sample sets from the Latino population is needed as advocated for stringent confirmation of genetic associations. Furthermore, comprehensive fine-mapping and resequencing will be required to characterize the disease gene architecture of this region in the Latino population.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

We thank all the patients and their families for their support and participation. This study was funded in part by the National Institutes of Mental Health (RO1-MH0698567) and by the Center of Excellence in Neurosciences at the Paul L. Foster School of Medicine. We utilized the Illumina genotyping system of the Institutional Genomic

Resource Core at the University of Texas Health Science Center at San Antonio under the Cancer Center Support Grant (P30 CA54174) through the gracious help of Teresa Johnson-Pais, Mandy Rolando, and Korri Weldon. We also acknowledge the help of Carolina Livi in preparing the illustrations.

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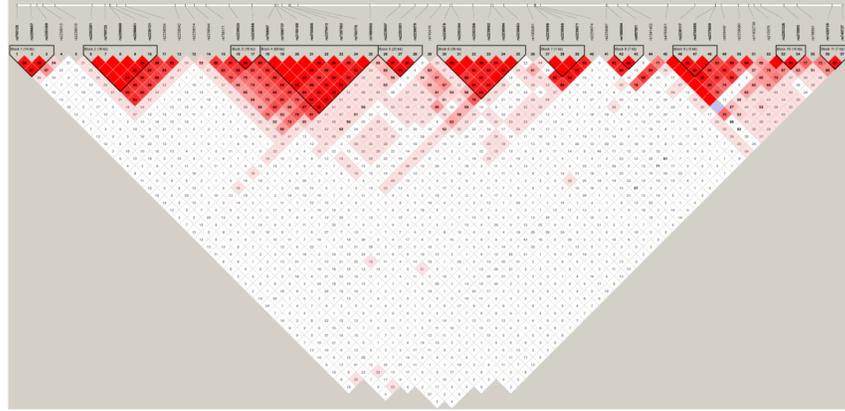


Fig. 1.

Calcium channel, voltage-dependent, L-type, alpha 1C subunit (*CACNA1C*) haplotype blocks in the Latino population. Linkage disequilibrium plot of the *CACNA1C* gene (shown as a black line) and adjacent regions with respect to the 57 single-nucleotide polymorphisms (SNPs) with minor allele frequency > 10%. The data visualized here using Haploview V4.2 (53) are based on the current genotyping data from the Latino population. Squares indicate pair-wise r^2 values on a red-scale with $D = 1$ (red) through to $D = 0$ (white). Linkage disequilibrium blocks, as defined in Haploview, are represented by the triangular lines. Order of Blocks and SNPs (SNPs within Blocks in bold) from left to right: **Block 1 (14 kb): rs765125, rs2299657, rs2283269;** rs2238013; rs2238019; **Block 2 (18 kb): rs2283281, rs758723, rs2299660, rs2299661, rs2239131;** rs2238034; rs2238042; rs2239014; rs2238044; rs758171; **Block 3 (16 kb): rs2239024, rs2238056;** **Block 4 (69 kb): rs769087, rs1006737, rs2159100, rs4765905, rs2370413, rs7297582, rs758170, rs1860002;** **Block 5 (20 kb): rs2239047, rs2283301, rs2238070;** rs740418; **Block 6 (31 kb): rs2238078, rs2283304, rs2283306, rs2239062, rs2238084;** rs2239063; rs1009281; **Block 7 (1 kb): rs2239066, rs2239068, rs2239071;** rs2239074; rs2239087; **Block 8 (7 kb): rs1860094, rs887001;** rs1541452; rs4765951; **Block 9 (14 kb): rs2239117, rs4765959;** rs2370600; rs929492; **Block 10 (7 kb): rs2238090, rs11832738;** rs215976; **Block 11 (16 kb): rs2283326, rs215993;** rs758561; **Block 12 (7 kb): rs1420725, rs740727.**

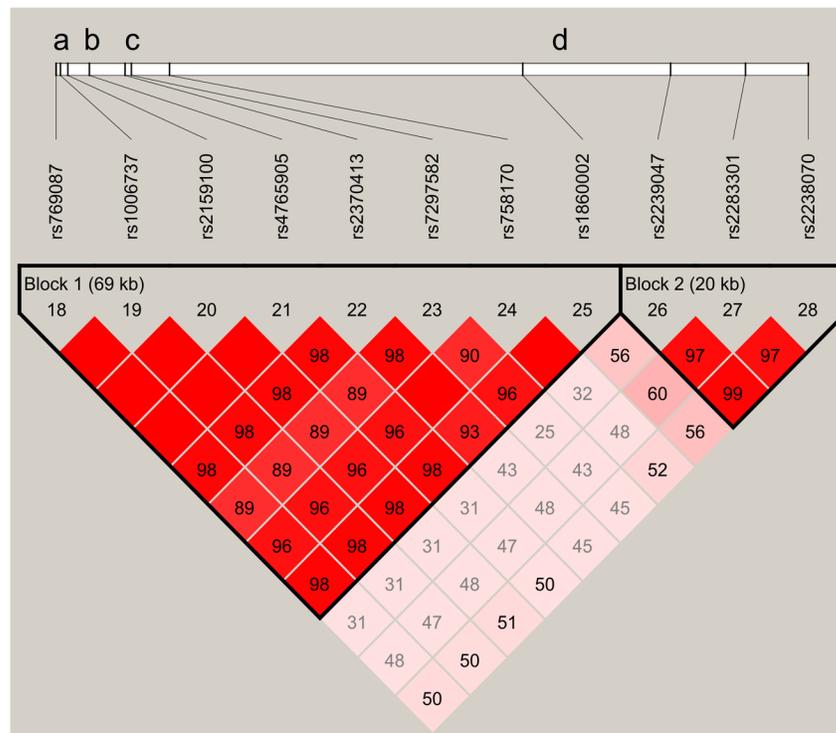


Fig. 2. Calcium channel, voltage-dependent, L-type, alpha 1C subunit (*CACNA1C*) haploblocks in the Latino population in relation to single-nucleotide polymorphisms (SNPs) previously associated with bipolar disorder. Haploview V4.2 (53) was used to define linkage disequilibrium (LD) blocks represented by the triangular lines, based on the current genotyping data from the Latino population. The 111.5 kb *CACNA1C* gene region (shown as a black line) is depicted with respect to 11 genotyped SNPs forming the two haploblocks. Block 1 consists of 8 SNPs in strong LD including rs1006737^a and rs7297582^c, SNPs that have previously shown to associate with bipolar disorder (20, 21). Ripke et al. (27) reported rs4765905^b associated with bipolar disorder and schizophrenia. Subsequent to this study, Sklar et al. (57) reported a bipolar disorder associated SNP (rs4765913^d), which is positioned between Blocks 1 and 2. Squares indicate pair-wise r^2 values on a red-scale with $D = 1$ (red) through to $D = 0$ (white).

Table 1
 Distribution of families per country of origin based on corresponding number of affected individuals^a

Affected individuals	United States		Mexico		Costa Rica		Guatemala	
	Families	Mean (range)	Families	Mean (range)	Families	Mean (range)	Families	Mean (range)
1	16	3.56 (2-6)	22	3.05 (3-4)	5	3.40 (3-5)	2	3.00 (3)
2	29	3.69 (2-7)	45	3.84 (3-8)	12	4.67 (3-9)	9	3.56 (3-5)
3	14	4.00 (3-8)	22	5.05 (3-13)	13	5.85 (4-10)	3	4.00 (4)
4	2	4.00 (4)	7	5.86 (4-8)	5	7.80 (5-14)	2	6.00 (6)
5	1	8.00 (8)			2	8.50 (8-9)	1	8.00 (8)
7					1	10.00 (10)		

Affected individuals = number of family members with a diagnosis of bipolar disorder type I and/or schizoaffective bipolar disorder; families = number of families per country of origin; mean = average number of individuals per family; range = range of individuals per family.

^aThere is no data presented for families with six affected individuals

Table 2

Family-Based Association Test univariate results under additive model for eight SNPs with linkage disequilibrium in *CACNA1C* gene

Marker	Allele	Allelic frequency	Families	Z	p-value
rs769087	G	0.715	98	-0.372	0.710
	A	0.285	98	0.372	0.710
rs1006737 ^a	G	0.712	100	-0.725	0.469
	A	0.288	100	0.725	0.469
rs2159100	G	0.711	99	-0.588	0.557
	A	0.289	99	0.588	0.557
rs4765905	C	0.710	101	-0.585	0.559
	G	0.290	101	0.585	0.559
rs2370413	G	0.520	130	-0.323	0.746
	A	0.480	130	0.323	0.746
rs7297582 ^a	G	0.697	108	-0.306	0.760
	A	0.303	108	0.306	0.760
rs758170	G	0.713	102	-0.639	0.523
	A	0.287	102	0.639	0.523
rs1860002	A	0.561	123	-1.161	0.246
	G	0.439	123	1.161	0.246

SNP = single-nucleotide polymorphism; CACNA1C = calcium channel, voltage-dependent, L type, alpha 1C subunit; families = number of informative families.

^aSNPs associated with bipolar disorder in previous studies.

Table 3
Haplotype analysis for eight SNPs with linkage disequilibrium in *CACNA1C* gene

Haplotype	Frequency	Families	S-E(S)	Var(S)	Z	p-value	rank(S_obs)	P_2side
GGGCAGGA	0.463	119	8.497	60.133	1.096	0.273	7432.5	0.283
AAAGGAAG	0.255	93	3.057	46.619	0.448	0.654	17063.5	0.651
GGGCGGGG	0.102	59	5.412	23.034	1.128	0.259	7647	0.290
GGGCGGGA	0.093	52	-13.865	22.972	-2.893	0.004 ^a	52692.5	0.004 ^a
GGGCGAGG	0.023	16	0.305	6.082	0.124	0.902	22174	0.837
AAAGGGAG	0.017	15	0.185	4.574	0.087	0.931	27104.5	0.977
GGGCAAGG	0.012	11	-2.418	3.209	-1.350	0.177	45600	0.269
GGGCGAGA	0.008	5	-2.528	1.218	-2.291	0.022 ^a	52591	0.009 ^a
AAAGGGGG	0.008	5	0.694	1.934	0.499	0.618	17009.5	0.644
GGGCGGAG	0.004	4	0.075	1.419	0.063	0.950	25296	0.952
AAAGGAAA	0.003	3	-0.498	1.171	-0.460	0.646	29224	0.895
GGGCAAGA	0.002	1	1	0.5	1.414	0.157	6667	0.248
AAAGAGGA	0.002	1	-1.917	0.543	-2.601	0.009 ^a	52076.5	0.028 ^a
<u>GA</u> AGG <u>AA</u> G ^b	0.002	2	2.5	1.25	2.236	0.025 ^a	847	0.031 ^a

Whole marker permutation test (χ^2 -sum, $p = 0.0018$)

Haplotypes comprised of single-nucleotide polymorphisms (SNPs): rs769087, rs1006737, rs2159100, rs4765905, rs2370413, rs7297582, rs758170, and rs1860002. Haplotypes with frequency 0.001 not shown. CACNA1C = calcium channel, voltage-dependent, L type, alpha 1C subunit; Frequency = frequency of the haplotype; S = test statistics for the observed number of transmitted alleles; E(S) = expected value of S under the null hypothesis (i.e., no linkage or association); P_2side = two-sided p-values using permutation test based on 10,000 permutations.

^a p-values < 0.05.

^b Bipolar risk alleles are bolded and underlined.

Table 4

Allele frequencies among different populations

Marker	Allele	Current study	CEU	JPT + CHB	YRI
rs769087	G	0.715	0.642	0.936	0.364
	A	0.285	0.358	0.064	0.636
rs1006737 ^a	G	0.712	0.627	0.933	0.383
	A	0.288	0.373	0.067	0.617
rs2159100	G	0.711	0.642	0.928	0.383
	A	0.289	0.358	0.072	0.617
rs4765905	C	0.710	0.636	0.928	0.400
	G	0.290	0.364	0.072	0.600
rs2370413	G	0.520	0.508	0.372	0.950
	A	0.480	0.492	0.628	0.050
rs7297582 ^a	G	0.697	0.575	0.950	0.958
	A	0.303	0.425	0.050	0.042
rs758170	G	0.713	0.642	0.930	0.527
	A	0.287	0.358	0.070	0.473
rs1860002	A	0.561	0.517	0.652	0.058
	G	0.439	0.483	0.348	0.942

CEU, CEPH = Utah residents with ancestry from northern and western Europe; JPT + CHB = Japanese in Tokyo and Han Chinese in Beijing; YRI = Yoruba in Ibadan, Nigeria.

^aSingle-nucleotide polymorphisms associated with bipolar disorder in previous genome-wide association studies of European ancestry.