

Haz R Iddaa Excel Program _TOP_

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Genetic variation in serotonin transporter gene associated with re-emerging affective disorders. Serotonin (5-HT) dysfunction has been implicated in the development of affective disorders, and single-nucleotide polymorphism (SNP) G/A-1438G>A in the serotonin transporter gene (5-HTT) is believed to correlate with the transcriptional activity of the gene. In the current study, we investigated whether the occurrence of affective disorders is affected by G/A-1438G>A and examined its interaction with other five SNPs in the 5-HTT gene in a Korean population. A total of 229 Korean first-episode affective disorder patients and 153 age-, sex-, and clinical status-matched healthy controls were genotyped. The occurrence of affective disorders was examined in relation to the five SNPs, and the interaction of these SNPs on the occurrence of affective disorders was assessed using haplotype-based logistic regression analysis. The 5-HTT genotype frequencies were analyzed by both logistic and log-linear regression models to examine the risk of affective disorders in relation to G/A-1438G>A. A genetic comparison using log-linear models with multiple comparisons adjusted by the Bonferroni correction revealed that G/A-1438G>A was significantly associated with affective disorders only in the cases ($P=0.007$). Haplotype-based analyses indicated that the frequency of haplotype G/G was significantly lower among affective disorder cases, compared with controls (PA polymorphism in the 5-HTT gene in a Korean population).

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