Epigenetic Changes of Serotonin Transporter in the Patients with Alcohol Dependence: Methylation of an Serotonin Transporter Promoter CpG Island

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Objective  Psychiatric disorders such as depression, anxiety and alcohol dependence are associated with serotonin metabolism. We assessed the methylation level of the serotonin transporter (5-HTT) promoter region in control and alcohol dependent patients.

Methods  Twenty seven male patients who met the Diagnostic and Statistical Manual of Mental Disorder IV (DSM-IV) criteria for alcohol dependence were compared with fifteen controls. Polymerase chain reaction (PCR) assays of bisulfate-modified DNA were designed to amplify a part of the CpG island in the 5HTT gene. Pyrosequencing was performed and the methylation level at seven CpG islands sites was measured.

Results  We found no differences in the methylation patterns of the serotonin transporter linked promoter region (5-HTTLPR) between alcohol-dependent and control subjects.

Conclusion  Our negative finding may be because 5-HTT epigenetic variation may not affect the expression for 5-HTT or there may be other methylation site critical for its expression. To find out more conclusive result, repeating the study in more methylation sites with a larger number of samples in a well-controlled setting is needed.

Key Words  Serotonin transporter region (5-HTTLPR), Methylation, Alcohol dependence.

INTRODUCTION

Psychiatric disorders such as major depression, obsessive compulsive disorder, autism, and alcohol dependence are associated with serotonin neurotransmission.1-4 The serotonin transporter (5-HTT) is an integral membrane protein that transports the neurotransmitter serotonin from the synaptic space into presynaptic neurons and is a member of the SLC6 gene family, which reuptakes other neurotransmitters (e.g., glycine, gamma-aminobutyric acid, dopamine, and norepinephrine) across the plasma membrane of neurons and glia.5 There are many studies regarding 5-HTT genetic polymorphisms and the subsequent affects on etiology and response to treatment in mental illness.6-9

Feinn et al.4 reported a significant association with the frequency of the short (S) allele at the 5-HTT linked promoter region (5-HTTLPR) in patients with alcohol dependence. In a meta-analysis, the S allele was significantly associated with an early onset, severe subtype and a co-morbid psychiatric disorder in alcohol dependence. Decreased raphe 5-HTT, which correlated with lifetime alcohol consumption, and association of the 5-HTT gene regulatory polymorphism in patients with alcohol dependence were reported.10 Hinckers et al.11 reported that carriers of two long (L) alleles of the 5-HTT regulatory region had a lower response to alcohol than carriers of one or two S alleles among adolescents.

The CpG island that surrounds exon 1 was recently discovered; its role in psychiatric disorders is unknown. Effects of
5-HTTLPR on 5-HTT mRNA production is modest and other cis- and trans-acting elements involved in the regulation of the 5-HTT gene region, such as the CpG island, constitute the leading mechanisms that control 5-HTT activity. Although heritable components affect gene activity, gene-environment interactions may contribute to gene status by gene regulatory elements such as CpG islands.

However Philibert et al. reported relation between the methylation level of 5-HTT promoter region and well known 5-HTTLPR polymorphism. They opened the possibility that 5-HTTLPR polymorphism may be the reflection of methylation difference of 5-HTT promoter region.

We hypothesized that there was a relationship between serotonin transporter epigenetic changes and alcohol dependence; thus, we investigated the differences in quantitative CpG residue methylation of seven 5-HTT promoter CpG positions between control subjects and patients with alcohol dependence.

**METHODS**

**Subjects**

We assessed the 5-HTT promoter methylation level of the subjects. Twenty seven male patients who met the Diagnostic and Statistical Manual of Mental Disorder IV (DSM-IV) criteria for alcohol dependence were interviewed and sampled. They were all admitted to Hangang Sacred Heart Hospital. Two hospital psychiatrists determined the diagnosis in a consensus procedure using all available clinical material, including a semistructured interview based on the DSM-IV. Patients were excluded from the study if they had a history of comorbid psychiatric illness or physical disorders other than alcohol-related medical illnesses. The control subjects consisted of fifteen unrelated healthy males, mostly nondrinkers. A few of the control subjects were occasional light drinkers, as revealed by a drinking-habit questionnaire.

Mean age for the control subjects and alcohol dependent patients was 44.93±6.47 and 46.67±13.74 years, respectively. The two groups did not differ significantly in age (t=-0.46, df=17.5, p=0.65). The institutional review board of Hangang Sacred Heart Hospital approved the study and all subjects provided informed consent after a complete description of the present study.

**Procedures**

Alcohol dependent patients consumed alcohol almost continually before admission. Ten mL of fasting blood was drawn into an EDTA tube on the day after admission and the blood was immediately centrifuged at 3,800 rpm for 10 minutes. Plasma was stored at -70°C until it was thawed for assay.

**Bisulfite modification**

Discrimination of methylated and unmethylated site of CpG island were done Bisulfite modification. Bisulfite treatment changes unmethylated cytosine residues to uracil, but leaves methylated 5-methylcytosine unaffected. Bisulfite treatment of 200 ng of each sample was accomplished using the EZ DNA Methylation-Gold kit (Zymo Research, Orange, CA, USA) according to the manufacturer’s instructions. The bisulfite converted DNA was eluted in 20 ul of elution buffer (Zymo Research, Orange, CA, USA). DNA samples were immediately stored at -20°C until use.

**Primer design and pyrosequencing methylation analysis**

Pyrosequencing were used to identify sequence change after bisulfate modification. Pyrogram (Figure 1) shows DNA sequence representation with sequential addition of dNTP detected by luciferase in one sample. Methylation Index was described above gray area which represents each CpG sites.

Polymerase chain reaction (PCR) assays were designed to

| Gene | GeneBank No. | Primer | Size (bp) |
|------|-------------|--------|-----------|
| 5HTT | *182138     | Forward| 5’-GTAAAGTAAAGAGTAGGAAAGTTAGG-3’ |
|      |             | Biotinylated-reverse| 5’-AACCCTCAATCTGAGTACAT-3’ |
|      |             | Sequencing   | 5’-TTTTGGTTTTGGGT-3’ |

*Figure 1. Representative pyrogram of the 5HTT gene promoter. The seven targeted cytosines are enclosed in gray squares. The Methylation Index (MtI) was calculated as the average rate of G incorporation at each CpG.*
amplify a part of the CpG island in the 5HTT gene. Primers were designed using the PSQ assay design program (Biotage, Charlotte, NC, USA). The primer sequences are listed in Table 1. The PCR reaction was carried out in a volume of 50 µl with 20 ng or less converted gDNA, 5 µl of 10X Taq buffer, 5 units Hot Start Taq polymerase (Enzynomics, Korea), 4 µl of 2.5mM dNTP mixture, and 2 µl of 10 pmole/ul. PCR primers. Amplification was carried out according to the general guidelines suggested by pyrosequencing: denaturating at 95°C for 5 min, followed by 45 cycles at 95°C for 40 sec, 60°C for 45 sec, and 72°C for 40 sec and a final extension at 72°C for 5 min. Confirmation of PCR product quality was established on a 1.5% agarose gel stained with ethidium bromide. Pyrosequencing was performed using the PSQ96MA System (Biotage, Charlotte, NC, USA) according to the manufacturer’s instructions, including single strand binding protein (Pyro Gold reagents; Qiagen Inc., Valencia, CA, USA). PCR primer sequences, PCR conditions and sequencing primer sequences are given in Table 1.

Data analysis

The methylation index (MtI) at the promoter and for the samples was calculated as the average value of methylated cytosine/(methylated cytosine+unmethylated cytosine) for all examined CpGs in the gene. All data were analyzed using SPSS for Windows software (version 12.0; SPSS Inc., Chicago, IL, USA). All analyses were set at a two-tailed significance level of p<0.05.

RESULTS

An example of a bisulfite methylation profile is shown in Figure 1. There was no difference in MtI between patients with alcohol dependence and controls at any of the seven CpG residues (Table 2). We considered low-level methylation (0-5%) to represent background ‘noise’ with questionable significance.

| Position | Alcohol dependence (N=27) | Control (N=15) | p |
|----------|---------------------------|----------------|----|
|          | Mean±SD                   | Mean±SD        |    |
| Position 1 | 1.31±2.43                 | 0.89±3.46      | 0.17 |
| Position 2 | 2.68±2.70                 | 2.35±2.95      | 0.62 |
| Position 3 | 1.12±1.85                 | 2.56±3.32      | 0.16 |
| Position 4 | 4.44±3.48                 | 3.72±5.87      | 0.18 |
| Position 5 | 3.24±2.90                 | 3.81±3.57      | 0.70 |
| Position 6 | 2.76±2.76                 | 3.64±2.51      | 0.34 |
| Position 7 | 1.56±2.30                 | 3.63±2.51      | 0.14 |

Statistical significance was determined by the Mann-Whitney U test

DISCUSSION

Complex epigenetic mechanisms that regulate gene activity without altering DNA code produce long-lasting changes in gene expression essential to development, cellular differentiation, and adaptation to environmental changes. These mechanisms that include DNA methylation, post-translational covalent modifications of histones, nucleosome sliding and nucleosome and histone substitution, cause modifications of chromatin conformation, which, in turn, regulate gene expression. For example, the amount of DNA methylation in promoter regions correlates with gene inactivation. Currently, with regard to alcohol dependence, alpha synuclein (SNCA), alcohol dehydrogenase and HTT genes are often studied in epigenetics. This study examined serotonin transporter epigenetic changes in alcohol dependent patients. However, we found no differences in the methylation patterns of the serotonin transporter promoter region between alcohol dependent and control subjects.

Currently, with regard to alcohol intoxication, there are few studies regarding methylation patterns of the serotonin transporter promoter. Philibert et al. found that females had more CpG methylation in the 5-HTT gene and less mRNA expression. They discovered a trend of increased overall methylation in people with a lifetime history of major depression. However, there was higher 5-HTT mRNA expression in patients with a lifetime history of alcohol dependence.

Serotonin transporter mRNA expression is affected by 5-HTTLPR genotype. However, there was no relationship with overall CpG methylation level in the present study. This result is in accordance with Philiert et al., who didn’t found the main effect of total methylation and 5-HTT mRNA level. Although increased average methylation only after 5HTTLPR genotype (l/l or s/s) was taken into account; effects of CpG methylation were then evident.

The epigenetics field and its methodologies are relatively new and limited. To identify epigenetic changes associated with alcohol dependence, the ideal experiment would prospectively investigate the dynamics of genome-wide epigenetic changes in the brains of individuals who eventually become affected with alcohol dependence, as compared to unaffected control individuals. At the present time, however, limitations with our current range of epigenetic methodologies render such an approach impossible. In addition, because there have been no thorough epigenetic studies with regard to alcohol dependence to date, it is not known how large a sample would need to be investigated. At this stage, the degree of any potential epigenetic differences between cases and controls is un-
known, although the effects observed are likely to be small.

This study has an important limitation. Peripheral blood-based assays will progress after properly controlling different confounding variables that have an effect on DNA methylation, including medications, plasma homocysteine level, and underlying disease. However, the use of peripheral cells would also allow the longitudinal study of epigenetic changes throughout the life course of patients with alcohol dependence, providing information about the epigenetic factors associated with the development, remission and relapse of the disorder.

Conclusively, in this research, there were no differences of 5-HTTLPR epigenetic DNA methylation between patients with alcohol dependence and control subjects. It is thought that 5-HTT epigenetic variation may not affect the expression for 5-HTT or there may be other methylation site critical for its expression.

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