

Hypomethylation of *MB-COMT* promoter is a major risk factor for schizophrenia and bipolar disorder

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The variability in phenotypic presentations and the lack of consistency of genetic associations in mental illnesses remain a major challenge in molecular psychiatry. Recently, it has become increasingly clear that altered promoter DNA methylation could play a critical role in mediating differential regulation of genes and in facilitating short-term adaptation in response to the environment. Here, we report the investigation of the differential activity of membrane-bound catechol-*O*-methyltransferase (*MB-COMT*) due to altered promoter methylation and the nature of the contribution of *COMT* Val158Met polymorphism as risk factors for schizophrenia and bipolar disorder by analyzing 115 post-mortem brain samples from the frontal lobe. These studies are the first to reveal that the *MB-COMT* promoter DNA is frequently hypomethylated in schizophrenia and bipolar disorder patients, compared with the controls (methylation rate: 26 and 29 versus 60%; $P = 0.004$ and 0.008 , respectively), particularly in the left frontal lobes (methylation rate: 29 and 30 versus 81%; $P = 0.003$ and 0.002 , respectively). Quantitative gene-expression analyses showed a corresponding increase in transcript levels of *MB-COMT* in schizophrenia and bipolar disorder patients compared with the controls ($P = 0.02$) with an accompanying inverse correlation between *MB-COMT* and *DRD1* expression. Furthermore, there was a tendency for the enrichment of the Val allele of the *COMT* Val158Met polymorphism with *MB-COMT* hypomethylation in the patients. These findings suggest that *MB-COMT* over-expression due to promoter hypomethylation and/or hyperactive allele of *COMT* may increase dopamine degradation in the frontal lobe providing a molecular basis for the shared symptoms of schizophrenia and bipolar disorder.

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INTRODUCTION

Despite strong evidence supporting a genetic basis for schizophrenia and bipolar disorder, no specific gene with a major effect has been identified as causing these complex and devastating disorders that affect five million people in the USA and 120 million worldwide (1). Several lines of evidence indicate that an improved understanding of the etiology of psychiatric illnesses can only be obtained using integrative approaches that consider multiple factors linked to these diseases, i.e. genetic, epigenetic and/or environmental factors (1,2). Methylation of the cytosine residues in the regulatory CpG islands of genes (3,4) is one of the major molecular mechanisms for regulating differential gene expression in response to environment. There are several DNA methyltransferases that act interactively to mediate gene-specific methylation patterns in response to environmental effects as well as the physiological state of the cell (5–7). Although S-adenosyl methionine (SAM) is the major donor of methyl groups for DNA methylation, folic acid and vitamin B12 are required for homocysteine metabolism to regenerate the pool of SAM (8). Modification of the gene promoter methylation pattern in response to a variable environment can have deleterious effects similar to those derived from malfunctioning alleles. The ability of altered DNA methylation to produce a disease phenotype in response to specific environmental insults likely depends upon genetic susceptibility to a specific disease (2). Recently, we and others (9,10) showed that DNA hypermethylation-mediated hypo-expression of the *reelin* gene (*RELN*) in the frontal lobe of the brain correlated with schizophrenia pathogenesis. These findings led us to hypothesize that alterations in the expression patterns of other neuronal network genes involved in the pathogenesis of schizophrenia and bipolar disorder may be mediated by epigenetic modifications such as altered DNA methylation of regulatory regions.

Catechol-*O*-methyltransferase (*COMT*), one of the most intensively investigated genes in psychiatric illnesses (2), regulates the homeostatic levels of neurotransmitter dopamine in the synapses. This gene is located in chromosome 22q11.21, a region linked by several studies to schizophrenia and bipolar disorder (11–14). *COMT* has two known isoforms: membrane-bound *COMT* (*MB-COMT*) and soluble *COMT* (*S-COMT*), each with its own promoter (15). *MB-COMT* is the predominant form involved in the degradation of synaptic dopamine in the human brain (15,16). *MB-COMT* exhibits a functional polymorphism at codon 158 where an adenine (A) substitution for guanine (G) in the gene sequence produces a methionine (Met) substitution for valine (Val) in the *COMT* protein. The Val-coding allele of the Val158Met polymorphism codes for an enzyme with approximately three times higher physiological activity (17). Several studies showed that increased dopamine degradations arising from *COMT* hyperactivity are associated with disturbances in attention, executive cognition and working memory performance in normal populations and schizophrenic patients (18–22) and that these effects may even be progressive over time (21). In contrast, the hypoactive allele (Met) of the gene is associated with less variability in reaction time and greater stability in performance (23).

Despite the fact that several studies reported an association between the Val allele and the risk of schizophrenia and bipolar disorder (16,24,25), meta-analyses of family-based and case–control association studies were inconsistent (26–28). Furthermore, the *COMT* Val158Met polymorphism is linked to early onset major depressive disorder (29) and suicide (30), suggesting that depressive symptoms may also be associated with *COMT* functions. The variability in genetic associations between the *COMT* Val158Met polymorphism and major psychiatric disorders suggests that, in addition to a genetic basis for *COMT* activity, environmental factors may also be involved in modulating the effects of the risk allele. For example, besides folic acid and vitamin B12, butyrate (31), tea polyphenols (32) and alcohol (33–35) influence genes promoter DNA methylation and may also affect expression level and modulate the phenotypic outcome of risk alleles. There is precedence in the field of cancer research supporting epigenetic modulation of *COMT* function, as DNA hypermethylation of the CpG-rich islands in the promoter of *MB-COMT* has been correlated with hypo-expression of the gene in some types of cancer (36). Here, our overall goal is to analyze the combined genetic and epigenetic data in correlation with the role of gene–environmental interactions in an integrated model. In the present study, we provide evidence for over-activity of the *MB-COMT* in the frontal lobe of patients with schizophrenia and bipolar disorder due to over-expression of the gene resulting from epigenetic promoter DNA hypomethylation and, preliminarily to a lesser extent, the presence of the hyperactive allele of *COMT* Val158Met polymorphism.

RESULTS

The differential promoter DNA methylation patterns associated with altered gene expression are tissue-specific (3–5). Frontal lobe is one of the most commonly observed dysfunctional regions of the brain in schizophrenia and bipolar disorder (37,38). Recent studies indicate that epigenetic modifications may be involved in the dysfunction of prefrontal cortex in schizophrenia (9,10,39). Furthermore, schizophrenia-associated problems in attention, cognitive processes and working memory have been linked to frontal lobe dopamine deficiency (40). Because of the relative scarcity of dopamine transporters in the frontal lobe of the brain, the turnover of dopamine is likely to be primarily dependent on the catabolism of dopamine by the action of *COMT* in this brain region (41). Therefore, we compared the status of promoter DNA methylation and the corresponding transcript levels of the *MB-COMT* in the frontal lobe of post-mortem brains of schizophrenia and bipolar disorder patients against the control subjects. In the same samples, the allele and genotype frequencies of the *COMT* Val158Met polymorphism were also determined as an alternate or an additional mechanism for regulating dopamine levels due to the hyperactivity associated with the Val allele.

The DNA and RNA samples were derived from Brodmann's area 46 (dorso-lateral frontal cortex dissected from post-mortem brains) obtained from the Stanley Medical Research Institute (SMRI). In addition, the Harvard Brain

Tissue Resource Center (HBTRC) provided 10 well-matched samples of brain tissue from the frontal lobe (Brodmann's area 9 and 10) of schizophrenia patients and normal controls. Demographics of these subjects are summarized in Table 1. The ethnicity of most of the samples was white (96.6%), except for two samples from blacks: one from a Hispanic and one from a Native American. Although only the SMRI data were used for matching samples for methylation analysis (because of the lack of information on all the variables for the HBTRC samples), the patterns of methylation were similar in the two subgroups. Gene-expression analysis was performed using total RNA obtained from the SMRI and RNA extracted from the cortical area of the HBTRC brain samples.

There were no significant differences between the case and control groups with respect to age ($t = 0.86$, $P = 0.39$), post-mortem interval ($t = -0.59$, $P = 0.55$) and brain laterality ($\chi^2 = 0.06$, $P = 0.81$). Further details of the SMRI brain samples are provided at: http://www.stanleyresearch.org/programs/brain_collection.asp. Although there was a trend for greater brain pH in controls compared with patients (mean = 6.6 versus 6.45, two-sample median test $Z = -1.79$, $P = 0.07$), there was no effect of pH on any of the variables (data not shown).

MB-COMT promoter methylation status of schizophrenia and bipolar disorder patients

The methylation status of the promoter region of *MB-COMT* was analyzed using methylation-specific PCR (MSP), and representative examples are shown in Figure 1. At least two MSP trials were carried out for each sample. Although there was ~90% consistency between two trials (the same unmethylated or methylated signals in both MSP trials), a third trial was also undertaken to define the predominant methylation status of any ambiguous samples. In addition, bisulfite sequencing (as described under Materials and Methods) was undertaken for further clarification and confirmation. A summary of the analyses of 35 schizophrenia patients, 35 bipolar disorder patients and 35 controls is shown in Table 2. In general, it appeared that *MB-COMT* promoter was predominantly unmethylated in the frontal lobe of human post-mortem brain tissue from both patients and controls. However, careful examination of these brain samples revealed a significantly higher frequency of methylation in control subjects (60%), compared with either schizophrenia (26%; $\chi^2 = 7.01$; $P = 0.004$) or bipolar disorder patients (29%; $\chi^2 = 7.89$; $P = 0.008$) (Table 2). Although the frequency of methylation in the left brain of the controls was significantly higher than in the right brain (81 versus 42% methylated, respectively), this difference was less pronounced in schizophrenia (29 versus 22% methylated) and bipolar disorder patients (30 versus 27% methylated) (Fig. 2). These results suggested that the group differences in total methylation derived primarily from differences inherent to the left brain between controls and patients with schizophrenia ($P = 0.003$) or bipolar disorder ($P = 0.002$). As shown in Figure 1, a representative example for the left brain of the control subjects (Fig. 1B, M) indicates that the degree of CpG methylation was relatively high in the promoter region corresponding to base pairs -150 to +100 relative to the start point of the coding

Table 1. Sample characteristics

Source	Diagnosis	Number of cases	Sex M/F	Age mean (SD)	Laterality L/R
SMRI	Schizophrenia	35	26/9	42.5 (8.47)	17/18
SMRI	Bipolar disorder	35	18/17	45.2 (10.54)	20/15
SMRI	Control	35	26/9	44.2 (7.63)	16/19
HBTRC	Schizophrenia	5	5/0	46.0 (2.74)	3/2
HBTRC	Control	5	5/0	45.4 (2.61)	3/2

M, male; F, female; L, left; R, right.

region, as determined by bisulfite sequencing and detailed in the Materials and Methods section. On the contrary, in schizophrenia and bipolar disorder patients, the target cytosines of the promoter region were hypomethylated and these changes were found to extend even into the coding region (Fig. 1B, U). These differences were not attributable to the effects of age or gender (Table 3).

MB-COMT promoter methylation and alcoholism

The history and degree of life-time alcohol consumption for all of the samples (until death) were provided by the SMRI. As recent studies reported an association between alcoholism and promoter DNA hypermethylation of several genes (33–35), the relationship between alcohol abuse and the frequency of *MB-COMT* methylation was examined. Among schizophrenia patients, *MB-COMT* promoter methylation was remarkably more frequent in current heavy (4/9) and moderate (2/3) alcohol users (total 6/12 methylated = 50%) than patients with no history of alcohol use (1/9) or social drinking (1/7) (total 2/16 methylated = 12.5%, $P = 0.04$, Fisher's exact test). Such a pattern of alcohol abuse was rare in controls (i.e. among the 35 control cases examined, there were only two current-moderate abusers with an unmethylated and two previous-heavy abusers with a methylated *MB-COMT* promoter).

Overall, only nine out of 35 schizophrenic patients and 10 out of 35 bipolar patients of the SMRI samples exhibited *MB-COMT* promoter methylation. Interestingly, most of these schizophrenic patients were moderate/heavy alcohol abusers at the time of their death (6/9, 66%). Similarly, 30% of these bipolar patients were either previously or at the time of their death heavy alcohol abuser. Thus, overall, our data suggest that a portion of the observed methylation could be due to alcohol abuse and hence the actual *MB-COMT* methylation rate may be even lower than what has been observed in schizophrenia and bipolar patients. Despite the fact that these preliminary results may be informative, the data should be interpreted cautiously because of our small sample size.

MB-COMT promoter methylation and drug use

We also examined the link between antipsychotic and mood stabilizer drugs and *MB-COMT* promoter methylation by comparing the frequency of methylation in bipolar disorder patients who were either treated or untreated with such drugs. In the bipolar disorder samples from the SMRI, 12 patients had used antipsychotic drugs at the time of death, whereas 16 patients

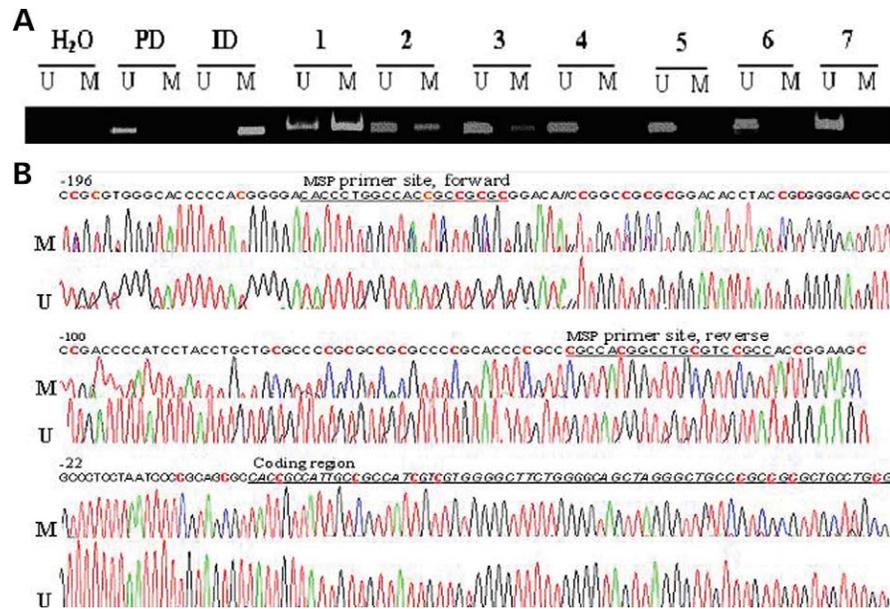


Figure 1. Representative examples of the MSP and sequence analyses of *MB-COMT* promoter. Brain samples from schizophrenia and bipolar disorder patients and control subjects were analyzed using MSP as outlined under Materials and Methods. (A) MSP: lanes U and M indicate the presence of unmethylated and methylated templates, respectively. Placental DNA (PD) and *in vitro* methylated DNA (ID) served as negative and positive controls, respectively. A no-template sample (H₂O) was used to detect DNA contamination in the PCR reactions. Samples 1–3 indicate the presence of a methylated promoter DNA and samples 4–7 represent an unmethylated promoter. (B) Bisulfite sequencing of the *MB-COMT* promoter of an un-methylated template (sample 4) correlates with conversion of cytosine (C) to thymine (T) in the DNA sequence traces in the promoter region corresponding to CpGs (U). However, the samples exhibiting methylation (sample 1) contained sequence traces for Cs in addition to Ts at several CpGs (which include two SP1 binding sites), indicating that the template is a mixture of unmethylated and methylated DNA (M). The color codes for the nucleotides are as follows: red, thymine; green, adenine; black, guanine and blue, cytosine. The original DNA sequence is indicated at the top of the trace; red Cs are within CpG sequences and are targets for methylation.

Table 2. Frequency of *MB-COMT* promoter methylation in the SMRI samples as determined by MSP

	Methylated/total (%)		
Frontal lobe hemisphere	Left	Right	Total
Schizophrenia	5/17 (29)	4/18 (22)	9/35 (26)
Bipolar	6/20 (30)	4/15 (27)	10/35 (29)
Control	13/16 (81)	8/19 (42)	21/35 (60)
Comparison	OR (95% CI); <i>P</i>	OR (95% CI); <i>P</i>	OR (95% CI); <i>P</i>
Control versus schizophrenia	0.10 (0.01–0.61); 0.003	0.39 (0.10–1.59); 0.197	0.23 (0.07–0.71); 0.004
Control versus bipolar disorder	0.10 (0.02–0.46); 0.002	0.50, (0.12–2.07); 0.350	0.27 (0.09–0.80); 0.008

did not. The frequency of *MB-COMT* promoter methylation was the same in these two subgroups (no antipsychotic drugs: 37% methylated versus antipsychotic drugs: 33% methylated; $\chi^2 = 0.05$; $P = 0.82$). Among the bipolar disorder patients, 20 used either antipsychotic or mood stabilizer drugs at the time of death, whereas eight had used neither class of drugs (we did not have drug use information for seven of the patients). Only three patients with schizophrenia were drug-free at the time of death. Altogether, the frequency of *MB-COMT* promoter methylation was not significantly different in these two sub-groups as well (neither drug: 45% methylated versus either drug: 30% methylated, $P = 0.4$). Note that half of the bipolar patients who used neither drug and exhibited methylation of *MB-COMT* promoter were heavy alcohol abusers at the time of death. By excluding them from this analysis, 33% of non-drug users show methylation of *MB-COMT* promoter. Interestingly, the

use of histone deacetylase/DNA methylation-inhibitor valproate also did not specifically alter methylation status (29% exhibited methylation out of 17 valproate users). These results support the idea that antipsychotic or mood-stabilizing drugs were not responsible for the observed methylation differences. Furthermore, our analyses indicate that smoking, brain pH, post-mortem interval and age of onset did not affect the frequency of *MB-COMT* methylation (data not shown).

Expression analysis of *MB-COMT*

It has been well established in the literature that hypermethylation of the promoter DNA generally correlates with silencing of gene expression (3,5). We treated a cancer cell line, H441, which exhibited partial *MB-COMT* promoter DNA methylation, with 5-azacytidine (a demethylating agent) as described

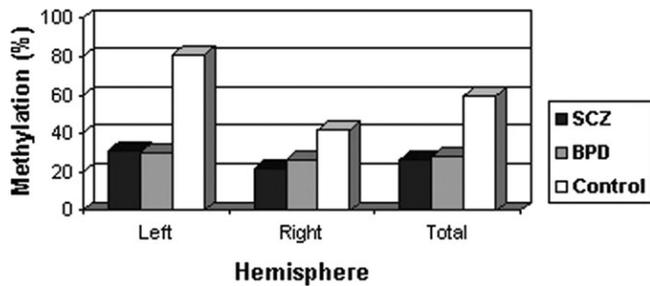


Figure 2. The effect of laterality of brain in *MB-COMT* promoter DNA methylation. The frequency of methylated *MB-COMT* promoter in the left and right hemispheres of the brains of patients with schizophrenia (SCZ) or bipolar disorder (BPD) and normal controls.

Table 3. DNA methylation status as a function of gender

	Methylated/total (%)		Gender (χ^2 , <i>P</i>)
	Male	Female	
Total sample	27/69 (39)	13/36 (36)	0.13 (0.72)
Schizophrenia	7/26 (27)	2/9 (22)	0.11 (0.74)
Bipolar disorder	4/17 (23)	6/18 (33)	0.41 (0.52)
Control	16/26 (62.)	5/9 (56)	0.10 (0.75)

elsewhere (42). We found that there was a prominent induction of *MB-COMT* expression (data not shown) confirming a similar observation made in an independent study (36). Therefore, we examined the relationship between *MB-COMT* promoter DNA methylation status and gene expression in the HBTRC and SMRI samples.

The expression analysis of *MB-COMT* at the transcript level was undertaken for the frontal cortical area of the brain tissues provided by the HBTRC using semi-quantitative reverse transcriptase PCR (RT-PCR) and was subsequently confirmed by quantitative real-time PCR (qRT-PCR). To minimize the effects of confounding variables, we dissected the brain cortex of the HBTRC samples who were male with a mean age of 46 and a narrow range of age (SD = 2.7). Pure gray matter was excised and DNA and RNA were extracted from the same sections using TRIzol. Semi-quantitative RT-PCR analysis of these samples showed a distinct pattern of over-expression of *MB-COMT* in schizophrenia patients, compared with control subjects (Fig. 3A). In order to further clarify and confirm the differences in *MB-COMT* expression in patients and controls, we re-evaluated these samples by qRT-PCR (Table 4). Overall, we found that there was 2.7 times higher expression of *MB-COMT* in schizophrenia patients than in controls using the samples from the HBTRC (Table 4). Furthermore, qRT-PCR analysis of the SMRI samples also showed that the transcript level was significantly ($t = -2.37$, $P = 0.02$) higher in patients with schizophrenia and bipolar disorder compared with control subjects (Fig. 3B). However, the magnitude of this difference was less than what was observed in the HBTRC samples. One possibility for this difference may be derived from the nature of the tissues used for the extraction of RNA. We used the same brain section from the gray matter for the isolation of

both DNA and RNA for the HBTRC samples, whereas homogenized brain tissues dissected from adjacent cortical brain regions were used to extract either DNA or RNA for the SMRI samples, due to an established sample collection procedure of the provider. Nevertheless, as expected, both semi-quantitative RT-PCR (Fig. 3A) and qRT-PCR data (Fig. 3B, Table 4) were consistent in revealing an inverse correlation between *MB-COMT* promoter methylation and expression (Fig. 3C). Furthermore, the level of *MB-COMT* expression of combined samples exhibiting unmethylated *MB-COMT* promoters was 56% more than the controls that exhibited methylation in the left brain. This was even more prominent among 11 schizophrenic and bipolar patients who were drug-free at the time of death. Excluding the two patients who were severely alcoholic, the expression of *MB-COMT* among these patients with an unmethylated promoter was more than 2-fold higher than controls with a methylated promoter in the left frontal lobe of the brain. These studies also showed that the expression levels of *MB-COMT* with Met- or Val-coding alleles were not drastically different (data not shown).

Inverse correlation between *MB-COMT* and *DRD1* expression

As aberrant *MB-COMT* promoter methylation and corresponding differential expression patterns could influence the level of synaptic dopamine and the expression of downstream genes, we analyzed the expression of *DRD1* at the level of transcription in the same brain samples using qRT-PCR, as described under Materials and Methods. In general, there was an overall hypo-expression of *DRD1* in the patients compared with the control subjects, with a clear inverse correlation between *MB-COMT* and *DRD1* expression both in controls and in patients (Fig. 4). In order to quantify the correlation between *MB-COMT* and *DRD1* expression, we classified the total samples, controls and patients groups into two subgroups (low and high *MB-COMT* expressive) as sorted by *MB-COMT* expression levels. There was a significant degree of inverse correlation between *MB-COMT* and *DRD1* expression in the 52 low and 52 high *MB-COMT* expressive groups of the total samples ($P = 0.001$; two-tailed *t*-test). The expression of *DRD1* in 17 schizophrenic and bipolar disorder patients with high levels of *MB-COMT* was also significantly reduced compared with the same group with low levels of *MB-COMT* ($P = 0.045$ and 0.025 , respectively; two-tailed *t*-test) (Fig. 4). A trend for such relationship in the control subjects was also detected ($P = 0.058$). It is also noteworthy that the expression of *DRD1*, in high *MB-COMT* expressive schizophrenic and bipolar disorder patients, was significantly less than the control subjects ($P = 0.006$ and 0.03 , respectively).

COMT Val158Met polymorphism in schizophrenia and bipolar disorder

The hyperactive Val allele of the Val158Met polymorphism was linked to schizophrenia and bipolar disorder (25,26). Therefore, to determine the relative contributions of either genetic or epigenetic alterations in the pathogenesis of

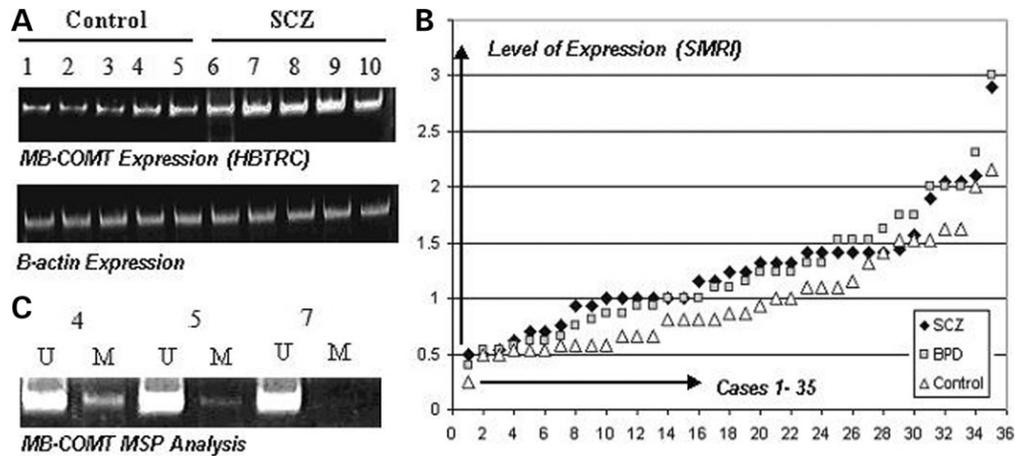


Figure 3. *MB-COMT* expression profiles of schizophrenia and bipolar disorder and the relationship between methylation status and *MB-COMT* expression. (A) Semi-quantitative RT-PCR of schizophrenia and control samples was performed as described under Materials and Methods. Control samples were loaded in lanes 1–5 and samples from schizophrenia patients were loaded in lanes 6–10. β Actin was used as an internal standard to normalize the abundance of the RT-PCR product derived from the *MB-COMT* gene. (B) qRT-PCR analysis of the patients and control subjects was performed as described under Materials and Methods. For each sample (1–35, X-axis), the relative quantitation of the *MB-COMT* expression compared with the mean of the controls is sorted from minimum to maximum level (Y-axis). As indicated earlier, most of the schizophrenic (SCZ) and bipolar disorder patients (BPD) show higher levels of expression of the *MB-COMT* in the frontal lobe compared with the controls. (C) MSP analysis showing levels of methylation in representative samples from (A).

Table 4. *MB-COMT* expression analysis of HTBRC samples by qRT-PCR

Samples	1	2	3	4	5	6	7	8	9	10
Brain laterality	R	R	L	L	L	L	R	L	R	L
Methylation status	M	M	M	M	U	U	U	U	U	M
Relative expression of <i>MB-COMT</i>	0.84	1.15	0.5	0.87	2.45	2.3	4.6	2.3	4.6	1.7
Relative fold change	Controls = 1.16 (SD = 0.76)					SCZ = 3.1 (SD = 1.4)				
Patients versus controls	$(t = -2.74, P = 0.026)$									

R, right; L, left; M, methylated; U, unmethylated.

Samples 1–5 are from controls and 6–10 from schizophrenic patients. CT value was used for relative quantitation of *MB-COMT* expression (normalized to *GADPH* gene expression as an internal control). Fold changes was calculated as $2^{-\Delta\Delta CT}$.

schizophrenia and bipolar disorder, we surveyed the potential contribution of this polymorphism in the same samples used for the evaluation of *MB-COMT* promoter methylation status (Fig. 5). Although the data should be interpreted cautiously because of the small sample size (i.e. 115), we found that the frequency of the Val allele was significantly low in controls (37.5%) compared with the patients with schizophrenia (55.0%, $P = 0.026$) in the SMRI and HBTRC samples combined. In addition, controls were less frequently homozygous for the Val allele (10%) compared with the patients with bipolar disorder (28.5%, $P = 0.040$) (Table 5). The frequency of Met homozygosity also exhibited a corresponding increase in controls compared with the schizophrenics (35 versus 15%, $P = 0.04$), but not bipolar disorder patients. Additionally, the overall age of onset of schizophrenia in patients homozygous or heterozygous for the Val allele was 7 years younger compared with the patients homozygous for Met (20.2 versus 27.4 years, $t = 2.60$, $P = 0.014$). Thus, the presence of the Val allele exhibits a significant correlation ($P = 0.01$) with an earlier onset of schizophrenia. Furthermore, we found a correlation between *COMT* Val homozygosity and DNA hypermethylation of the *RELN* promoter in a cAMP-response element as determined using a previously described method

(9). All of the schizophrenics and control subjects with a Val/Val genotype exhibited a hyper-methylated *RELN* promoter (14/14), whereas it was only 50% (10/20) in Met/Met homozygotes ($P = 0.01$) and 75.5% in Val/Met heterozygotes.

Our analyses reported here provide a clear trend that needs to be further validated in a larger population in the future studies. Indeed, given our small sample size, the observed association between Val allele and these mental diseases should be considered cautiously, particularly due to the lower frequency of the Val allele in our control subjects compared with larger studies in Caucasians (e.g. 49% in reference 26 and 42% in alfred.med.yale.edu). Note that within our data set, the data on allele frequencies are consistent with the Hardy–Weinberg equilibrium. The observed frequencies of *COMT* genotypes did not significantly deviate from those expected under Hardy–Weinberg equilibrium for controls [$\chi^2_{(1)} = 1.20$, $P = 0.273$], schizophrenia patients [$\chi^2_{(1)} = 1.80$, $P = 0.180$] or bipolar disorder patients [$\chi^2_{(1)} = 0.25$, $P = 0.615$].

As mentioned earlier, our objective in studying this polymorphism was to undertake an integrated approach to examine both genetic and epigenetic factors involved in the pathogenesis of major mental disorders. In our sample set,

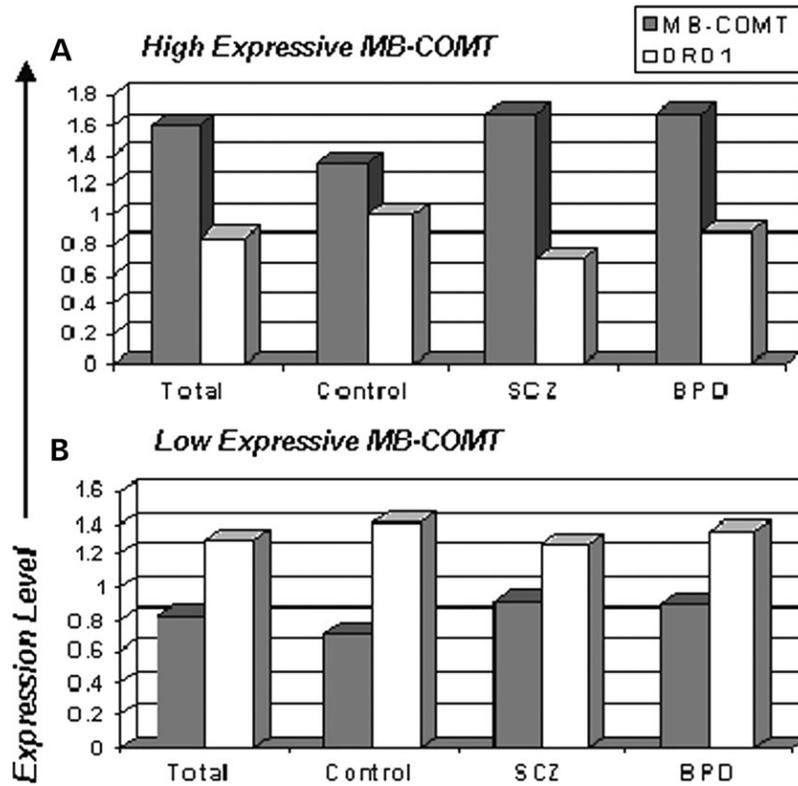


Figure 4. Relationship between the expression of *MB-COMT* and *DRD1*. The upper (A) and lower (B) panels represent the gene expression profiles of high and low expressive *MB-COMT* groups, respectively, in total samples, control subjects, schizophrenia (SCZ) and bipolar disorder patients (BPD). As it is shown, when *MB-COMT* expression is high, the expression of *DRD1* is low and vice versa, both in control subjects and in the patient group implying that, in general, hyper-expression of *MB-COMT* is associated with hypo-expression of *DRD1*.



Figure 5. *COMT* Val158Met polymorphism in schizophrenia and bipolar disorder. The *COMT* Val158Met polymorphism analysis was performed as described under Materials and Methods. Briefly, *COMT* gene-specific PCR products encompassing the polymorphic region were cleaved by the *Nla*III restriction enzyme to an 88 bp fragment and two variable fragments 96 and 114 bp in length, which indicated the Met and Val alleles, respectively. The first lane contains a 50 bp DNA ladder used as the marker. Representative examples are shown for heterozygosity (lanes 2, 4, 5 and 7), Met homozygosity (lanes 3 and 9) and Val homozygosity (lanes 6, 8 and 10).

differences in frequency of *MB-COMT* hypomethylation in the left brain (51%) and differences in Val homozygosity between all cases and controls (16.5%) were found to explain 60% of the variance in risk for either schizophrenia or bipolar disorder (51% plus 16.5% minus overlaps). Interestingly, among 14 schizophrenic and bipolar patients of the SMRI samples with Met homozygosity, the only patient who was diagnosed as schizo-affective and one bipolar patient with severe alcoholism had a methylated *MB-COMT* promoter (13%, both of them in right hemisphere), whereas nine of 13 controls with such genotype had a methylated *MB-COMT* promoter (69%, $\chi^2 = 10.83, P = 0.001$).

The effect of *COMT* Val158Met polymorphism and promoter hypomethylation on suicide

Val homozygosity and *MB-COMT* hypomethylation were also examined in bipolar patients who had committed suicide. Among 15 bipolar patients who died due to suicide, 12 had *MB-COMT* promoter hypomethylation (80%), whereas two of the three who had a methylated *COMT* promoter were Val homozygous. The rate of suicide was 60% (6/10) and 33% (3/9) among Val and Met homozygotes, respectively. It is also noteworthy that all of the Met homozygotes in the suicidal group had a hypomethylated *MB-COMT* promoter. After excluding the three patients from the non-suicidal group who died due to drowning which may or may not reflect a suicidal attempt, the frequency of either *MB-COMT* hypomethylation or Val homozygosity was 93% (14/15) and 65% (11/17) in the suicidal and non-suicidal bipolar groups, respectively ($\chi^2 = 3.82, P = 0.05$). Despite no conclusive pattern of promoter methylation or Val158Met *COMT* polymorphism in the seven schizophrenic patients who died by suicide, there were no cases exhibiting Met homozygosity with a methylated *MB-COMT* promoter among all suicidal patients, including schizophrenics. Interestingly, the average level of *MB-COMT* expression in the suicidal patients with Met homozygosity was 55% more than other patients.

Table 5. Frequency of Val and Met alleles of the Val158Met polymorphism of *COMT* in schizophrenia, bipolar disorder and control subjects

Diagnosis	Genotype frequency			Allele frequency	
	Val/Val, N (%)	Val/Met, N (%)	Met/Met, N (%)	Val, N (%)	Met, N (%)
Control	4/40 (10.0)	22/40 (55.0)	14/40 (35.0)	30 (37.5)	50 (62.5)
Schizophrenia	10/40 (25.0)	24/40 (60.0)	6/40 (15.0)	44 (55.0)	36 (45.0)
OR (95% CI); <i>P</i>	3.00 (0.76–14.27); 0.078	1.23 (0.46–3.28); 0.651	0.33 (0.09–1.08); 0.039	2.04 (1.03–4.03); 0.026	
Bipolar disorder	10/35 (28.6)	16/35 (45.7)	9/35 (25.7)	36 (51.4)	34 (48.6)
OR (95% CI); <i>P</i>	3.60 (1.06–12.08); 0.040	0.62 (0.23–1.68); 0.303	1.90 (0.62–5.83); 0.270	0.86 (0.45–1.64); 0.652	

ORs and test statistics are for each patient group versus the control group. Genotype comparisons are computed in relation to the sum of the other genotypes.

These data suggest that Met homozygosity and *MB-COMT* promoter hypermethylation may be protective against schizophrenia, bipolar disorder and suicide.

***COMT* polymorphism and promoter methylation in cardiac disease**

Because most of our control subjects died due to cardiac diseases, we compared *MB-COMT* methylation and polymorphism status in patients and controls with the same cause of death to exclude the possibility that higher frequencies of *MB-COMT* methylation and Met homozygosity may be attributed to cardiac diseases. We were particularly interested in the fact that the frequency of the Met allele in our control samples (62.5%) was slightly more than the average in populations of European descent (58%) (alfred.med.yale.edu). However, our analysis excluded this possibility as the frequency of *MB-COMT* methylation was the same in 15 schizophrenic patients who died by cardiac disease (26.5%) and in 20 bipolar patients who died by cardiac and other diseases (35%).

DISCUSSION

Our studies provide the first comprehensive molecular explanation at the level of genetic and epigenetic alterations for a direct relationship between the risk for schizophrenia and bipolar disorder and hyperactivity of *MB-COMT*. *MB-COMT* over-expression or hyperactivity observed in schizophrenia and bipolar disorder (Fig 3 and Tables 4–5) could lead to a significant increase in the rate of degradation of dopamine in the synaptic cleft resulting in a hypodopaminergic state and frontal lobe hypoactivity. Both schizophrenia and bipolar disorder patients exhibit hypofrontality, executive dysfunctions and frontal lobe lesions with associated unresponsiveness and failures to suppress inappropriate responses (38,43). Additionally, negative/depressive symptoms and recognized schizophrenia-associated problems in attention, hedonic activities, cognitive processes, working memory and social functioning have also been linked to dopamine dysfunction in the frontal lobe of the brain (1,40).

There have been contradictory data on the differences in expression of *COMT* in schizophrenia and mood disorders in the literature. One study suggested that there is no difference in the levels of *COMT* transcripts between cases and controls (44), whereas another study reported a reduction

in expression in superficial cortical layers and a corresponding increase in deep layers in schizophrenia (45). Although one possibility is that the differential laminar distribution of *COMT* transcripts could explain the lack of difference in the overall expression levels noted in the first study, our examination of the PCR primers/probe sequences used for the analysis of the *COMT* transcripts suggests that the primers/probes used in these studies could amplify or hybridize with both *S-COMT* and *MB-COMT* isoforms. A recent study reported that promoter DNA methylation of the *S-COMT* was not involved in the pathogenesis of schizophrenia (46). In this study, we evaluated the *MB-COMT*, which is regulated by a distinct promoter and is the primary mediator of the catabolism of synaptic dopamine in the human brain (15).

Recent studies indicate that a *DRD1*-mediated dopaminergic mechanism is responsible for working memory function (47) and synaptic plasticity (48) in the prefrontal cortex. Here, an inverse correlation between *MB-COMT* and *DRD1* expression was observed both in the controls and in the patients. This implies that frontal lobe dopamine deficiency may cause under-stimulation of dopamine receptors and hence a developmentally regulated hypo-expression of *DRD1* gene. Thus, *MB-COMT* over-expression in schizophrenic and bipolar patients may aggravate the end-effects in the patients leading to long-term downstream defects in the developing brain. In the promoter of several neuronal genes [e.g. brain-derived neurotrophic factor (BDNF)], the cAMP response elements (CREs) are regulated indirectly by *DRD1*-like receptors through a cascade of events mediated by cAMP, protein kinase-A and cAMP responsive element modulator (CREM). Although promoter DNA de-methylation leads to gene expression, neuronal inactivation could lead to *BDNF* promoter methylation and lack of gene expression (49). The promoter sequence of *RELN*, a gene hypo-expressed in schizophrenia and bipolar disorder patients (10,50–53), also harbors a *CRE* element that could be subject to regulation by dopamine through *DRD1* receptors (9). The observed inverse correlation between *MB-COMT* and *DRD1* expression and the association between *MB-COMT* over-activity and hyper-methylation of *CRE* in the *RELN* promoter in schizophrenia imply that the modulatory effects of *DRD1* on *RELN* expression could be compromised by epigenetically and/or genetically determined *MB-COMT* over-activity. Considering the roles of *RELN* in cell signaling and synapse formation, the resultant *RELN* hypo-expression should influence the brain neuronal network development and contribute

to the diseases (10,50–53). However, a hypoactive Met allele or promoter methylation and hypo-expression of *MB-COMT* could lead to a higher synaptic dopamine level that could illicit protective effects. Recent findings showed that activation of different forms of dopamine receptors in the frontal lobe depends on the synaptic dopamine level (54). Low levels of dopamine are associated with DRD1 activation, whereas higher levels are associated with activations of both DRD1 and DRD2 receptors that co-operate to provide the basis for the functionality of working memory. Interestingly, higher levels of synaptic dopamine delivered by DRD2 antagonists could also augment DRD1-mediated neuronal functions (54). It is also noteworthy that our analysis of other dopamine-related genes in the same samples revealed that in control subjects, when *MB-COMT* promoter was hypomethylated, almost always the *DRD2* promoter was also hypomethylated (data not shown). Thus, it appears that the synaptic dopamine deficiency associated with *MB-COMT* hypomethylation could result in secondary *DRD2* promoter hypomethylation in order to fine tune the dopamine transmission in the frontal lobe. On the contrary, in schizophrenic and bipolar patients, the level of compensatory *DRD2* hypomethylation was significantly low compared with the extent observed in the control subjects (data not shown).

The Met allele of the *COMT* gene is associated with greater stability in performance in normal individuals (21), whereas the Val allele is correlated with susceptibility for schizophrenia, particularly in cannabis abusers (55). Also a recent study indicates that children with 22q11 deletion, who are hemizygous for *COMT* Met allele, have higher IQ and achievement scores and perform better on measures of prefrontal cognition (e.g. Continuous Performance Task) compared with those with the Val allele (56). Schizophrenic patients homozygote for the Val also exhibit a significant degree of reduction in brain volumes particularly on the left side (anterior cingulate cortex, left amygdala-uncus and left thalamus), compared with Met homozygotes (57). Hence, the observed epigenetic *MB-COMT* hypomethylation/over-expression effects may also contribute to worsening of these brain abnormalities through reduction of *RELN* expression or other mechanisms.

Despite the fact that the *MB-COMT* promoter region is often hypomethylated in schizophrenia and bipolar disorder patients, there was a predominant unilateral partial methylation in the control subjects. These observations may indicate that differential expression of *MB-COMT* in the left frontal lobe may occur in response to the effects of other neuronal pathways and genes involved in brain laterality. Alternatively, *MB-COMT* promoter methylation in controls could occur as a physiological feedback response to environmentally mediated *MB-COMT* over-expression. Accordingly, a steady-state level of dopamine, required for specific neurons, is maintained to ensure the consolidation of the left brain dominance in defined end effects. In contrast, the observed promoter hypomethylation in the left brain of schizophrenia and bipolar disorder patients is likely a consequence of a failure to adhere to the predestined methylation pattern during a critical period for the establishment of brain laterality or a failure of physiological feedback modulation of the *MB-COMT*. This idea may also provide a plausible explanation for the loss of

brain lateralization in schizophrenia and bipolar disorders noted in several studies (58–61). One of the potential molecular mechanisms responsible for these defects could be loss of imprinting, as there is some evidence for linkage to maternally (but not paternally) transmitted alleles at the chromosome 22q locus in mental disorders (13).

Note that a systemic pathogenic insult such as folic acid deficiency during a critical developmental period can influence the extent of epigenetic modifications, including loss of imprinting and/or aberrant promoter DNA methylation patterns (62–64). Several lines of evidence indicate that folic acid deficiency and polymorphism of genes related to folate metabolism are associated with schizophrenia and mood disorders (65–67) as reviewed elsewhere (68). Furthermore, even the style of maternal care is known to alter offspring's genetic methylation patterns, as observed for the glucocorticoid receptor in the hippocampus (69).

Overall, these observations suggest that it may be possible to trace the environmental influences on the modification of methylation patterns of genes, which may affect susceptibility to schizophrenia, bipolar disorder and possibly other mental disorders. Accordingly, the concept of the roles of different methylomes as the means of short-term/long-term adaptation in response to the influences of the environment requires further extensive studies to understand the molecular basis of the major psychiatric disorders. It should be noted that the analyses reported here were conducted using brain samples corresponding to the left and right lobes from different individuals rather than derived from the same persons due to the lack of availability of the latter type of samples. Although we expect that the overall findings reported here should remain the same as pooling of the data derived with samples from several individuals should cancel out inter-individual variations, these investigations should be repeated in the future using left and right lobe-derived brain samples from the same individuals for further validation.

Despite the unavailability of an antibody to measure the COMT protein and the lack of the ideal samples for research due to practical issues, the establishment of an association between promoter DNA hypomethylation and the Val158Met polymorphism of *MB-COMT* in schizophrenia and bipolar disorder would suggest new treatment modalities in psychiatry targeting the specific dysfunctional genes. Animal studies have demonstrated that COMT inhibitor drugs prevent stress-induced anhedonic state, improve prefrontal cortex performance and potentiate clozapine-induced extra-cellular dopamine release (70,71). In human, these drugs (e.g. tolcapone) were effective in treating major depressive disorders (72,73). These observations along with the results presented here argue that COMT inhibitor drugs may be of benefit in the treatment of other major mental diseases. It is interesting to note that other drugs with similar end effects (e.g. amphetamine) improve prefrontal cortex functions affecting working memory tasks in subjects with the Val/Val genotype of *COMT* (74). Mirtazapine, an antidepressant with noradrenergic property, is also more effective than specific serotonin re-uptake inhibitors in depressed patients with the *COMT* Val allele (75). In schizophrenia, apomorphine, a dopamine agonist, was also known to improve psychotic symptoms (76). A recent meta-analysis of 700 patients from

30 studies showed that the concurrent administration of L-dopa with antipsychotic drugs could further improve the symptoms of schizophrenia in half of the cases (77). While generally, improvement of cognitive symptoms by atypical antipsychotic drugs is related to prefrontal cortex dopamine release (78), even atypical anti-psychotics such as olanzapine are less effective in aiding working memory performance and prefrontal physiology in schizophrenic patients with the *COMT* Val allele (79). These findings along with the observed efficacy of aripiprazole, a partial agonist of dopamine receptors in psychotic disorders, imply that negative and cognitive symptoms of schizophrenia as well as depressive symptoms of bipolar and unipolar disorders could be targeted by COMT-inhibitor drugs, particularly in patients with a hypomethylated *MB-COMT* promoter or the Val/Val genotype, as we and others discussed elsewhere (2,80,81).

Conclusion

Promoter hypomethylation-associated over-expression of *MB-COMT* in the left frontal lobe and or the presence of the hyperactive Val allele of the Val158Met polymorphism may play a significant role in the pathogenesis of major psychiatric disorders. The connection established between *MB-COMT* over-activity and pathogenesis in this report strongly suggests that it is worthwhile to consider preventive strategies such as nutrition- or drug-based modulation of *COMT* promoter DNA methylation or the use of COMT-inhibitor drugs to prevent or improve the symptoms of schizophrenia and bipolar disorder.

MATERIALS AND METHODS

DNA and RNA extraction

Total RNA and DNA were prepared from the same dissected cortex of the HBTRC brain samples, using TRIzol (Life Technologies, Inc., Grand Island, NY, USA) according to manufacturer's instructions. The SMRI extracted the total RNA and DNA using TRIzol and standard phenol-chloroform protocol, respectively, from the adjacent cortical areas of the post-mortem brains. cDNA synthesis and PCR amplification were performed using standard protocols as previously described (42).

Bisulfite treatment

Bisulfite converts unmethylated cytosine to uracil, whereas methylated cytosine remains unchanged (82). Then, a subsequent NaOH treatment converts uracil to thymine. Approximately 1 µg of genomic DNA was denatured by treatment with 2 M NaOH and then treated with 10 mM hydroquinone and 3 M sodium bisulfite (pH 5) at 50°C for 16 h. The bisulfite-treated DNA was purified using Wizard DNA purification resin (Promega) and treated with 3 M NaOH, ethanol precipitated with ammonium acetate and glycogen and re-suspended in 20 µl of water.

MSP and bisulfite sequencing

MSP is known as one of the most sensitive screening methods to distinguish between methylated and unmethylated DNA at

CpG sites. Although the sensitivity of direct sequencing of the bisulfite-treated DNA or cloned DNA is ~10–20% (83,84), the sensitivity of MSP is known to be >1% (83). This high sensitivity places the MSP as a highly favored method for screening of the methylated CpG sites within the CpG islands (83,85,86). Our MSP analysis of the *COMT* gene was carried out using primer sets described by Sasaki *et al.* (36). The two sets of primers corresponding to a CpG-rich area located in the *MB-COMT* gene promoter were used to amplify methylated and unmethylated strands in separate PCR reactions. The primer sequences used for the MSP analysis are as follows: methylated forward (MF) primer: 5'-TATTTTGGTTATCGTCGCGC-3'; methylated reverse (MR) primer: 5'-AACGAACGCAAACCGTAACG-3'; unmethylated forward (UF) primer: 5'-TATTTTGGTTATTGTTGTGT-3' and unmethylated reverse (UR) primer: 5'-AACAAACACAAA CCATAACA-3'.

For the MSP analysis, unmethylated (placental) DNA and *in vitro* methylated DNA were used as negative and positive controls, respectively, whereas a no DNA control (water) was used to detect any PCR contaminations. Various combinations of PCR cycles and annealing temperatures were examined to optimize PCR conditions, so that the unmethylated and methylated template-specific primers only generated PCR products that represent the unmethylated and methylated conditions, respectively. The high level of sensitivity of MSP allows the efficient high throughput examination of the DNA methylation status of test samples.

For bisulfite sequencing, the following primers were used to amplify a 421 bp fragment. Forward primer: 5'-GTAAGATT AGATTAAGAGGT-3' (36); reverse primer: 5'-CAATATT CCACCCTAAATCTAAAA-3' and a nested primer (5'-GAT ATTTTAC(T)GAGGATATT-3') or the reverse primer was used as the sequencing primer.

PCR conditions

Master mixes were prepared and used in the PCR amplifications. A typical reaction of a total volume of 25 µl consisted of 2.5 µl of 10× standard buffer, 0.4 µl of 10 µM dNTPs, 1.5 µl of DMSO, 2.5 U of platinum *Taq* DNA polymerase (Invitrogen) and 25 pmol of each of methylated or unmethylated specific primers. Approximately 30 ng bisulfite-modified genomic DNA was used as template. PCR amplification was done with an initial incubation at 94°C for 2 min, followed by 31 cycles of 94°C for 30 s, 57°C for 40 s, 72°C for 40 s, followed by a final extension for 10 min at 72°C. Ten microliters of the MSP products were run on 6% polyacrylamide gel by electrophoresis, stained with ethidium bromide and visualized under UV light.

Approach for determination of *MB-COMT* promoter methylation status

There were two MSP trials for each sample. As indicated earlier, for each MSP trial, two sets of primers were used to amplify methylated or unmethylated strands in separate PCR reactions. Approximately, 90% of the samples showed the same results for both MSP trials. Samples were classified as unmethylated if no methylation signal was detected in the

two MSP trials. Similarly, a sample was designated as methylated when a methylated signal was also detected in both MSP trials. Representative examples of samples exhibiting unmethylated or methylated status were subject to bisulfite sequencing (Fig. 1). Our results indicated that samples designated unmethylated did not exhibit traces corresponding to Cs due to complete conversion, whereas those determined to be methylated contained sequence traces of Cs indicating the presence of methylated Cs resistant to conversion by bisulfite treatment (see Fig. 1B for example). When the two initial MSP trials were not consistent (i.e. unmethylated in the first reaction and methylated in the second reaction), or whenever a weak methylated signal was observed, a third trial was also undertaken to define the methylation. Additionally, bisulfite sequencing was performed to make the final determination of the methylation status for these ambiguous samples. As these 'ambiguous' cases were distributed almost equally in different groups (three to four in each group, 11 in total) and excluding them from the analysis did not change the results of our statistical analyses in general, and for the observed findings as reported, we included them in the final analysis. Please also note that to avoid any bias with experiments, the investigators were blinded for the diagnosis of the samples until the MSP trials were completed. After the determination of the methylation status of the samples, the identity of samples was declared for data analysis.

Primers for *MB-COMT* expression analysis using semi-quantitative RT-PCR and qRT-PCR

The gene-specific primer pair used in the analysis of *MB-COMT* expression was designed from different exons to avoid any genomic DNA amplification and by placing them in locations which would exclusively amplify the *MB-COMT* isoform. The forward and reverse primers for the semi-quantitative RT-PCR were 5'-CTGCTTTGCTGCCGAGCTCAGAGGAGAC-3' and 5'-TTCTTGTCGCCACG TTCATG-3'. The qRT-PCR primers that solely amplify a 110 bp fragment were 5'-CTGCTTTGCTGCCGAGCTCAGAGGAGAC-3' and 5'-GCCCAGCAACACAGCTGCCAACAG-3'.

Semi-quantitative RT-PCR of the β actin gene and real-time PCR of the GAPDH and β actin genes were used as an internal standard to normalize the abundance of the semi-quantitative RT-PCR and qRT-PCR products derived from the *MB-COMT* gene for the HBTRC samples. The β -actin primer sequences for the semi-quantitative RT-PCR were 5'-ACACTGTGCCCATCTACGAGG-3' and 5'-AGGGGCCGGACTCGTCATACT-3' and for the qRT-PCR were 5'-C GAGCACAGAGCCTCGCCTTTGCC-3' and 5'-TGTCGACGACGAGCGCGGATAT-3'. The GAPDH primers were 5'-CACAAAGAGGAAGAGAGAGACCCTC-3' and 5'-TGAGTGTGGCAGGGACTC C-3'.

qRT-PCR analysis was done twice for the HBTRC samples, using ABI PRISM 7700 following several test trials with 1/2, 1/10, 1/100, 1/1000 diluted and no template controls with varying amount of primers to optimize for the best conditions. The qRT-PCR was set up in a reaction volume of 50 μ l using components supplied in a Sybr-green PCR mix (BIO-RAD, Hercules, CA, USA). Each reaction also contained 100 nm

each amplification primer and 2 μ l of 1/10 diluted DNA. DNA amplifications were carried out in a 96-well reaction plate (MicroAmp, Foster City, CA, USA). Thermal cycling was initiated with a first denaturation step of 15 min at 95°C. The subsequent thermal cycling profile was 95°C for 15 s and 60°C for 45 s. Data obtained following 40 cycles of amplification were analyzed using the SDS 1.9.1 software.

The qRT-PCR for the SMRI samples was carried out twice in a 384-well plate (Applied Biosystem, Foster City, CA, USA) in a reaction volume of 10 μ l using ABI PRISM 7900HT. Gene expression was measured with good reproducibility between duplicates and without the formation of primer dimers as evidenced by the appearance of a single product with a T_m of 82°C in the dissociation curve (data not shown). Cycle threshold (CT) value was used for relative quantitation of *MB-COMT* expression. Fold changes was calculated as $2^{-\Delta\Delta CT}$.

Primers for *DRDI* expression analysis

qRT-PCR was carried out using the same condition as described earlier for the *MB-COMT* analysis using ABI PRISM 7900HT. The primers were, forward primer: 5'-CTGCGACGAATAATGCCATAGAGA-3' and reverse primer: 5'-ATTGCACTCCTTGGAGATGGAGC-3'.

COMT gene Val158Met polymorphism

The *COMT* region was PCR amplified using a previously described primer set to evaluate the existence of the Val158Met polymorphism (87). The primers, 5'-TCGTGGACGCCGATTCAGG-3' and 5'-ACAACGGGTCAGGCA TGC-3', were used to amplify a 207 bp fragment after 35 cycles using the previously described conditions. The PCR products were further analyzed by restriction digestion by targeting the 177th base of the fragment with the *Nla*III restriction enzyme to distinguish between the Val and Met alleles (Fig. 5).

Data analysis

Univariate, bivariate and multivariate statistical tests were used for the analysis of the data. Parametric tests were used wherever possible. Large sample approximations (e.g. χ^2) were used wherever possible; otherwise, for small samples, exact tests were used (e.g. Fisher's exact test). Differences between cases (either schizophrenic or bipolar) and controls were examined. All statistical analyses were conducted using SAS software version 8.2. Note that the data from SMRI and HBTRC samples were pooled only for the polymorphism analysis.

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