Research paper

Investigation of betaine as a novel psychotherapeutic for schizophrenia

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Abstract

Background: Betaine is known to act against various biological stresses and its levels were reported to be decreased in schizophrenia patients. We aimed to test the role of betaine in schizophrenia pathophysiology, and to evaluate its potential as a novel psychotherapeutic.

Methods: Using Chdh (a gene for betaine synthesis)-deficient mice and betaine-supplemented inbred mice, we assessed the role of betaine in psychiatric pathophysiology, and its potential as a novel psychotherapeutic, by leveraging metabolomics, behavioral-, transcriptomics and DNA methylation analyses.

Findings: The Chdh-deficient mice revealed remnants of psychiatric behaviors along with schizophrenia-related molecular perturbations in the brain. Betaine supplementation elicited genetic background-dependent improvement in cognitive performance, and suppressed methamphetamine (MAP)-induced behavioral sensitization. Furthermore, betaine rectified the altered antioxidative and proinflammatory responses induced by MAP in vitro phencyclidine (PCP) treatments. Betaine also showed a prophylactic effect on behavioral abnormality induced by PCP. Notably, betaine levels were decreased in the postmortem brains from schizophrenia, and a coexisting elevated carbonyl stress, a form of oxidative stress, demarcated a subset of schizophrenia with decreased betaine levels.

Interpretation: The present study revealed the role of betaine in psychiatric pathophysiology and underscores the potential benefit of betaine in a subset of schizophrenia.

1. Introduction

There is an urgent need for novel medicines with new actions for psychiatric illness, because almost all of the present therapeutics are...
Almost all of the presently available antipsychotics are designed to act on the monoamine receptors or transporters, and their limited efficacy demands novel drug development paradigms. Metabolomic profiling approach in disease biology has been instrumental in deciphering mechanistic insights into disease pathophysiology, enabling the identification of novel therapeutic targets. Altered brain metabolome has been a consistent observation in schizophrenia, and it is also reflected in the peripheral samples. Betaine (glycine betaine or trimethylglycine) is one such metabolite that was observed to be decreased in the plasma samples of patients with first-episode schizophrenia. This prompted us to test the role of betaine in schizophrenia pathophysiology, and to evaluate its potential as a novel psychotherapeutic, by leveraging metabolomics, behavioral-, transcriptomics and DNA methylation analyses.

**Added value of this study**

We showed that the lack of betaine, stemmed from the genetic loss of Chdh activity in mice, resulted in remnants of psychiatric behaviors and schizophrenia-related molecular perturbations in the brain. Interestingly, the betaine supplementation elicited genetic background-dependent improvement in cognitive performance. Furthermore, betaine supplementation showed psychotropic action in pharmacological animal models of schizophrenia and rectified the altered antioxidative and proinflammatory responses characteristic of the model. In agreement with these observations, betaine levels were seen decreased in the postmortem brains from schizophrenia, and a coexisting elevated carbonyl stress (a form of oxidative stress) demarcated a subset of schizophrenia with “betaine deficit-oxidative stress pathology”. We further showed that the decreased betaine levels were consistent with the elevated carbonyl stress, in glyoxylase 1-deficient hiPSCs and betaine supplementation was efficacious in alleviating elevated carbonyl stress. We also identified a cis-expression quantitative trait locus (QTL) for Chdh expression in postmortem brains from schizophrenia, enabling genotype-based stratification of schizophrenia patients for betaine efficacy.

**Implications of all the available evidence**

The present study demonstrated that deficits of betaine levels in the brain contribute to the psychiatric pathophysiology and also revealed the potential benefit of betaine as a psychotherapeutic in a subset of schizophrenia cases.

designed to act on the monoamine receptors or transporters [1], and there is a substantial population where the presently available drugs are unsatisfactory [2]. Altered brain metabolome and its reflection in peripheral samples is a growing observation in schizophrenia, which could be beneficial for deciphering molecular pathophysiology of schizophrenia and thereby identifying novel druggable targets [3–5]. Betaine (glycine betaine or trimethylglycine) is a metabolite that is reported to be decreased in the plasma sample of patients with first-episode schizophrenia (FESZ) [3]. In their report, they examined the plasma metabolites from FESZs and healthy controls in order to identify biomarkers for schizophrenia. The reduction of the betaine level was replicated in the second sample set, suggesting future use of betaine as a clinical biomarker for schizophrenia.

**Research in context**

**Evidence before this study**

In vertebrates, betaine is ingested from the diet [5] and is also endogenously synthesized in mitochondria from its precursor choline using choline dehydrogenase (CHDH) [Fig. 1] [4]. To date, several biological functions of betaine have been proposed, which include (i) osmotic regulator (compatible solute) [6,7]; (ii) antioxidant/anti-inflammatory activity [8]; (iii) supplier of methyl donor (s-adenosylhomocysteine (Hcy); homocysteine; methyl-THF, 5-methyltetrahydrofolate; MS, methionine synthase; CBS, cystathionine beta-synthase; CGL, cystathionine beta-lyase; MSRA, methionine sulfoxide reductase A; MSRB1/2, methionine sulfoxide reductase B1/2. In vertebrates, betaine is ingested from the diet [5] and is also endogenously synthesized in mitochondria from its precursor choline using choline dehydrogenase (CHDH) [Fig. 1] [4]. To date, several biological functions of betaine have been proposed, which include (i) osmotic regulator (compatible solute) [6,7]; (ii) antioxidant/anti-inflammatory activity [8]; (iii) supplier of methyl donor (s-adenosylhomocysteine (Hcy); homocysteine; methyl-THF, 5-methyltetrahydrofolate; MS, methionine synthase; CBS, cystathionine beta-synthase; CGL, cystathionine beta-lyase; MSRA, methionine sulfoxide reductase A; MSRB1/2, methionine sulfoxide reductase B1/2.

**2. Materials and methods**

See the Supplementary information for the details of the techniques outlined below.

**2.1. Study approval**

All the animal experiments were performed in compliance with relevant laws, and guidelines were approved by the Animal Ethics
Committee at RIKEN (permission numbers: H29-2-204 [3] and 2016-058 [4]) and the Chiba University Institutional Animal Care and Use Committee (permission number: 31–342). Human induced pluripotent stem cell (hiPSC) study was approved by the Human Ethics Committee at RIKEN for iPSC study (Wako-daisan 25–14). The subject for hiPSC study gave informed, written consent to participate in the study after being provided with, and receiving an explanation of study protocols and objectives. Experiments in postmortem brain samples were approved by Fukushima Medical University, Japan (1685 and 2381) and Niigata University School of Medicine, Japan (G2015–0827). All procedures of postmortem brain study were carried out with the informed written consent of the next of kin.

2.2. Animals

The inbred C57BL/6NCrl (B6N) and C3H/HeNCrl (C3HN) mouse strains were obtained from Japan’s Charles River Laboratories (Yokohama, Japan). The animals were housed in groups of four in standard cages, in a temperature and humidity-controlled room with a 12 h light/dark cycle (lights on at 08:00). The animals had free access to standard lab chow and tap water. All animal experiments were done using male animals (six to 15 animals/group, depending on experiments) between 9:30 am and 5:00 pm. In the study using the Chdh KO mice, the wild-type littermates by intercross between heterozygotes were used as control. All the analyses were conducted between nine and 14 weeks of age, except for the novel object recognition test (NORT) using the ICR strain (purchased from Japan SLC, Hamamatsu, Japan), which was started at 7 weeks of age.

2.3. Postmortem brain samples

Postmortem brain tissues (BA17; Brodmann Area 17) from schizophrenia and age-matched control samples were obtained from the Postmortem Brain Bank of Fukushima for Psychiatric Research and Brain Research Institute, Niigata University, Japan (in total n = 24 for schizophrenia and n = 31 for control) [15–17]. Each individual with schizophrenia fulfilled the diagnostic criteria established by the American Psychiatric Association (Diagnostic and Statistical Manual of Mental Disorders: DSM-IV) and had no history of any other neurological disorder or substance abuse. The age matched control samples were with no history of neuropsychiatric disorders. Also, neuropathological evaluation in the control brain samples ruled out any pathological abnormalities characteristic of neurological disorders, though some brains showed indication of mild senility.

2.4. Generation of Chdh-deficient mice

The Chdh-deficient mice were generated by the genome editing methodology using the CRISPR/Cas9 nickase.

2.5. Estimation of betaine and other metabolites

Choline, methionine, betaine, cystathionine, cystine, S-adenosylmethionine (SAM), and S-adenosylhomocysteine (SAH) levels were estimated by liquid chromatography-mass spectrometry (LC/MS). Cysteine, homocysteine and glutathione (GSH) were measured as their reduced forms by high performance liquid chromatography (HPLC). The plasma levels of blood urea nitrogen (BUN), creatinine (CRE), aspartate transaminase (AST), alanine transaminase (ALT), lactate dehydrogenase (LDH) and cholesterol (CHE) were measured using dry-chemistry analysis (Fuji Dri-Chem 3500 V: FUJIFILM Medical Co., Ltd., Tokyo, Japan).

2.6. Behavioral analyses

All behavioral tests relevant to psychiatric illnesses, except for the methamphetamine (MAP)-induced behavioral sensitization test, were performed according to the previously published methods [18]. For the MAP-induced behavioral sensitization test, see the Supplementary Methods. When the effect of betaine supplementation on impairment of the NORT performance by chronic PCP treatment was examined, male ICR mice at 7 weeks of age were given betaine (2.5% as monohydrate in drinking water) or water until the day 29. Either PCP (10 mg/kg/day) or vehicle (saline) was subcutaneously administered once a day to the ICR mice from the day 15 to the day 26. The NORT experiment was started at the day 26 as described elsewhere [19]. The mice were habituated in the experimental environment for 3 days. The training and test sessions were conducted at the days 29 and 30, respectively.

2.7. DNA methylation analysis

DNA methylation in frontal cortex of Chdh knockout (KO) mice (n = 6) and wild-type (WT) control (n = 6) was performed by targeted methylation sequencing of 109Mb of mouse genomic regions, which included CpG islands, known tissue-specific differentially methylated regions (DMR), open regulatory annotations and Ensembl regulatory features. The inbred C57BL/6NCrl (B6N) and C3H/HeNCrl (C3HN) mouse strains were obtained from Japan’s Charles River Laboratories (Yokohama, Japan). The animals were housed in groups of four in standard cages, in a temperature and humidity-controlled room with a 12 h light/dark cycle (lights on at 08:00). The animals had free access to standard lab chow and tap water. All animal experiments were done using male animals (six to 15 animals/group, depending on experiments) between 9:30 am and 5:00 pm. In the study using the Chdh KO mice, the wild-type littermates by intercross between heterozygotes were used as control. All the analyses were conducted between nine and 14 weeks of age, except for the novel object recognition test (NORT) using the ICR strain (purchased from Japan SLC, Hamamatsu, Japan), which was started at 7 weeks of age.

2.8. RNA-seq analysis

Transcriptome analysis in frontal cortical brain region was performed by RNA-seq in (a) Chdh KO mice (n = 6) versus WT controls (n = six), and (b) B6N and C3HN mouse strains administered betaine in comparison to respective controls administered water (n = 6 in each group).

2.9. Real-time quantitative reverse transcription (RT)-PCR

Targeted gene expression was measured by real-time quantitative RT-PCR using TaqMan assays [20].

2.10. Rat primary cortical neuron culture

Rat primary cortical neuron was isolated from the cortices of Sprague-Dawley rats (obtained from Japan’s Charles River) at embryonic day 18.5 (E18.5) as described previously [21]. For the details, see the Supplementary Methods.

2.11. Establishment of GLO1-deficient hiPSCs

Human induced pluripotent cells (hiPSCs) were established from peripheral blood mononuclear cells using Sendai virus vector [22]. Isogenic GLO1-deficient hiPSCs were generated using CRISPR/Cas9-mediated genome editing.

2.12. Carboxyl stress assay

Carboxyl Stress was analyzed by Western blotting using anti-AGE antibody. See the Supplementary Methods.

2.13. cis-eQTL analysis

For eQTL analysis, we selected BHMT1, CHDH and GLO1 and downloaded eQTL data for the brain tissues from GTEx portal (https://gtexportal.org/home/) (Release V7; dbGaP Accession phs000424.v7. p2). The significant eQTL variants with high effect size were selected and filtered for minor allele frequency (MAF) > 30% in Japanese population with reference to the Tohoku megabank genome variation data.
from 3552 whole genome sequences (https://jmorp.megabank.tohoku.ac.jp/201902/downloads) (3.5KJPNv2). These variants were further pruned for linkage disequilibrium (LD) status ($r^2 < 0.4$), yielding variants from independent LD blocks. The shortlisted variants (4 in BHMT, 3 in CHDH and 3 in GLO1) were genotyped in schizophrenia postmortem brain tissues (BA17) (n = 50) by TaqMan SNP genotyping Assays [23]. Gene expression was measured by real-time quantitative RT-PCR using TaqMan assays. Outliers (more or less than mean ± 2SD) were excluded. Association of the variants with the gene expression and metabolites were tested by one-way ANOVA and Student’s t-test.

2.14. Statistical analysis
All values in the figures represent the mean ± SEM. Statistical analysis and graphical representation were performed using GraphPad Prism 6 (GraphPad Software). The total sample size (n) was described in the respective figure legends. Statistical significance was determined using a two-tailed Student’s t-test. When multiple comparisons were needed, one-way or two-way ANOVA with Fisher’s least significant difference (LSD) test, Tukey’s multiple comparison test, Bonferroni’s correction, or Dunnett’s multiple comparison test was used as indicated in the figure legends. A P value of <0.05 was considered as statistically significant.

3. Results

3.1. Generation of Chdh-deficient mice
In vertebrates, betaine is biosynthesized by two biochemical reactions (Fig. 1). The first step, oxidation of choline to betaine aldehyde, is mediated by choline dehydrogenase encoded by the Chdh/CHDH gene, and in turn, betaine aldehyde is converted into betaine by betaine aldehyde dehydrogenase. To create an animal model with lowered betaine level, the first coding exon, exon 2 of the Chdh gene, was targeted by the CRISPR/Cas9n on the genetic background of the inbred B6N mouse (Supplementary Fig. 3a and b). Homozygotes for the gene disruption, as shown by the unaltered plasma levels of AST, ALT, LDH, CHE, BUN or CRE (Supplementary Table 1). Furthermore, the total levels of glutathione (GSH + GSSG) were not altered by the gene disruption (Fig. 2d and e) in the brain or plasma, suggesting that the lack of Chdh activity itself did not invoke a serious oxidative stress condition in mice.

3.2. Betaine levels and effects of betaine supplementation in Chdh-deficient mice
The disruption of the Chdh gene significantly reduced the betaine levels in the frontal cortex from ~20 pmol/mg tissue to levels near detection limit ($p < 0.01$) (Fig. 2a). This suggested that the endogenous biosynthesis rather than intake from diet maintained the level of betaine, whereas homocysteine concentration was increased by the disruption of Chdh ($p < 0.01$), and it showed a trend towards WT level ($p < 0.1$) upon betaine supplementation (Fig. 2f). While in the brain (frontal cortex), homocysteine levels were unchanged among the groups (Fig. 2g). We also examined the two inbred mouse strains, B6N (the same genetic background with the Chdh-deficient mouse) and C3HN, each of which manifested two extremities for prepulse inhibition function (B6N, highest; C3HN, lowest), a schizophrenia and other mental disorder-related endophenotype [26]. Total homocysteine levels in the brain after betaine supplementation was also unchanged in the two inbred strains (Fig. 2h). The results suggest that homocysteine levels in the brain are robustly maintained constant against perturbation of peripheral betaine levels. Interestingly, the cellular methylation capacity deduced from the SAM/SAH ratio [27,28], showed a significant increase ($p < 0.05$) upon betaine supplementation in the B6N WT brain (Fig. 2i), suggesting a possibility that externally administered betaine could have an impact on the methylation capacity to some extent.

3.3. No adverse effects of betaine deficiency in kidney and liver functions
Since the kidney and liver show high expression of Chdh and contain abundant levels of betaine [25], we examined the effects of gene disruption with respect to their functions. The gene disruption elicited a trend of decreased betaine levels ($p < 0.1$) (Fig. 2c). Betaine levels in the kidney of KO mice were replenished to those of WT littermates upon betaine supplementation (Fig. 2c). Betaine in the liver plays a crucial role in one-carbon metabolism [25], but no impairments of liver function were observed by the Chdh disruption, as shown by the unaltered plasma levels of AST, ALT, LDH, CHE, BUN or CRE (Supplementary Table 1). Furthermore, the total levels of glutathione (GSH + GSSG) were not altered by the gene disruption (Fig. 2d and e) in the brain or plasma, suggesting that the lack of Chdh activity itself did not invoke a serious oxidative stress condition in mice.

3.4. Levels of metabolites in methionine–homocysteine cycle in the brain
Among the metabolites in the methionine–homocysteine cycle that were driven by betaine (Fig. 1), levels of homocysteine, a potent oxidant, changed differentially between the plasma and brain. In the plasma, homocysteine concentration was increased by the disruption of Chdh ($p < 0.01$), and it showed a trend towards WT level ($p < 0.1$) upon betaine supplementation (Fig. 2f). While in the brain (frontal cortex), homocysteine levels were unchanged among the groups (Fig. 2g). We also examined the two inbred mouse strains, B6N (the same genetic background with the Chdh-deficient mouse) and C3HN, each of which manifested two extremities for prepulse inhibition function (B6N, highest; C3HN, lowest), a schizophrenia and other mental disorder-related endophenotype [26]. Total homocysteine levels in the brain after betaine supplementation was also unchanged in the two inbred strains (Fig. 2h). The results suggest that homocysteine levels in the brain are robustly maintained constant against perturbation of peripheral betaine levels. Interestingly, the cellular methylation capacity deduced from the SAM/SAH ratio [27,28], showed a significant increase ($p < 0.05$) upon betaine supplementation in the B6N WT brain (Fig. 2i), suggesting a possibility that externally administered betaine could have an impact on the methylation capacity to some extent.

3.5. DNA methylation and transcriptome analyses in the Chdh-deficient mice
To explore the impact of betaine on DNA methylation status, a targeted analysis of methylated genomic regions at single base pair resolution in the frontal cortex from Chdh-deficient mice was performed. The results revealed no statistically significant differences in DNA methylation when compared to the WT mice after multiple corrections (q value >0.05; Fig. 3a, Supplementary Table 2). As an exploratory approach, we selected differentially methylated gene promoters, employing a conservative estimate of $p < 0.05$ to test the enrichment of canonical pathways and gene ontologies (Fig. 3b, Supplementary Table 3). Significant enrichment was observed for “tight junction signaling” and “sertoli cell-sertoli cell junction signaling” pathways for hypermethylated gene promoters, the latter could be related to the infertility phenotype of Chdh-deficient mice [29]. In the case of hypomethylated promoters, cytoskeletal dynamics pathways such as “integrin signaling” and “ephrin receptor signaling” were enriched (Fig. 3b).

Regarding the transcriptomic level changes elicited by the Chdh disruption in the frontal cortex, a total of 851 genes were significantly dysregulated (546 upregulated and 305 downregulated) in the KO compared to the WT littermates ($p < 0.05$) (Fig. 3c, and Supplementary Table 4). The Ingenuity Pathway Analysis showed significant enrichment for molecular pathways, for example, eukaryotic initiation factor 2 (eIF2) signaling, protein ubiquitination pathway, regulation of
eukaryotic initiation factor 4 (eIF4) and p70 S6 kinase signaling, and mammalian target of rapamycin (mTOR) signaling for the downregulated genes \((p < 0.05)\) (Fig. 3d, and Supplementary Table 5). These results were in accordance with the reduced protein synthesis observed in schizophrenia, which involve the eIF2α, eIF4 and mTOR signaling pathways [30]. The upregulated genes were also enriched for molecular pathways potentially related to the schizophrenia pathophysiology: glutamate receptor signaling and synaptic long-term depression (Fig. 3d, bottom). The upregulation results might be compensatory ones against reduced protein synthesis. Collectively, the altered molecular deficits by the Chdh disruption may represent a “molecular precursor signature” [31] underling schizophrenia pathophysiology.

3.6. Chdh-deficient mice displayed remnants of depressive behavior

We examined 15 distinct behavioral phenotypes relevant to psychiatric illnesses (Supplementary Fig. 6). In the forced swim test, immobility time at bin 5 (4-5 min) was significantly prolonged \((p < 0.05)\), and total immobility time showed a trend of increase \((p < 0.1)\) in the Chdh-deficient mice compared to the WT littermates (Figs. 3e and f), suggesting a depressive trait in the gene-deficient mice. In the other behavioral tests, no significant differences were observed between the WT and Chdh-deficient mice (Supplementary Fig. 6). The limited behavioral deficits elicited by the Chdh disruption could be partly due to the fact that betaine was not completely depleted from the whole body, because betaine was available from the diet or through intestinal microflora [32,33].

3.7. Effects of betaine supplementation in inbred mouse strains

To determine the therapeutic potential of betaine in psychiatric disorders, betaine was administered via drinking water (2.17% betaine) for 3 weeks to the two inbred mouse strains; B6N and C3HN. In general, efficacy of drugs, in particular neurotropics, varies with respect to the individual patients or genetic backgrounds. The same 15 distinct behavioral phenotypes as evaluated in the Chdh-deficient mice were measured. Notably, betaine supplementation elicted the differential behavioral effects in the two strains. In the novel object recognition test

![Fig. 2](image-url)

Fig. 2. Effects of Chdh deficiency and betaine supplementation on the metabolite contents. Metabolite contents were measured in the Chdh-deficient (Homo) and wild-type Chdh littermate control (WT) mice with (betaine) or without (water) chronic betaine supplementation. (a) Betaine levels were significantly reduced in the frontal cortex of Chdh-deficient mice, which were restored by betaine supplementation. (b) In plasma, betaine levels were not statistically significant among the tested groups. (c) Betaine levels were drastically reduced in the kidney of Chdh-deficient mice and were restored to WT level. Total glutathione (GSH) contents \((\text{GSH} + \text{GSSG})\), in the (d) brain and (e) plasma did not reveal any statistically significant differences among the tested groups. (f) Total plasma homocysteine increased in the Chdh-deficient mice when compared to WT, and showed a trend towards WT level after betaine supplementation; however, in the brain, the levels of total homocysteine were unchanged in the tested groups (g–h). (i) SAM/SAH ratio, indicating the cellular methylation potential, was significantly increased with betaine supplementation in WT B6N mice. Data represent mean ± SEM. *\(p < 0.05\), **\(p < 0.01\), ***\(p < 0.001\); Tukey’s multiple comparison test among the four groups, \(n = 5\) to seven per group.
Fig. 3. DNA methylome, transcriptome and behavioral analyses of Chdh-deficient mice. (a) Targeted DNA methylation analysis across the genome, from the frontal cortex of Chdh-deficient mice, did not reveal any statistically significant differences (after correcting for multiple tests; q-value < 0.05) when compared to the wild-type control. (b) Pathway analysis of genes with a p < 0.05 (uncorrected) indicated an enrichment of “tight junction signaling” and “sertoli cell-sertoli cell junction signaling” pathways for hypermethylated promoters, implicated in spermatogenesis. Hypomethylated genes were enriched for pathways involving cytoskeletal dynamics; n = six in each group. (c) Volcano plot shows differentially expressed genes between Chdh KO mice and wild type controls from frontal cortex; n = six in each group. Green and blue dashed lines indicate p value thresholds of 0.05 and 0.01, respectively. Top hits among the differentially expressed genes (p < 0.01 and absolute fold-change > 2) are highlighted in the plot. (d) Differentially expressed genes (p < 0.05) were analyzed for the enriched canonical pathways, which revealed enrichment for signaling pathways involved in the regulation of translational control and protein synthesis/degradation, for downregulated genes. (e) Results of forced swim test are presented. 1 bin corresponds to 1 min. Note that the Chdh-deficient mice showed increased immobility in bin five (p = 0.01). Data represent mean ± SEM. *p < 0.05, **p < 0.01; Fisher’s Least Significant Difference (LSD) test. n = 21 per group. (f) Cumulative values of immobility time for five minutes (bins two to six) were calculated. Note that KO homozygotes revealed a trend of increase in immobility time (p = 0.09). Data represent mean ± SEM. n = 21 per group.
NORT, B6N mice showed significantly longer stay time at a novel object in the retention session in the betaine-supplemented group compared to the water-drinking control, suggesting improved cognitive memory performance in the B6N strain (Fig. 4a). In contrast, C3H mice showed no significant improvement in the performance by betaine supplementation (Fig. 4a). Although we do not know the reason for the difference between the two strains, it may be related to the much lower preference to a novel object at the basal level when compared to familiar one in C3H mice. Both B6N and C3HN mouse strains showed better performance with betaine supplementation in the Y-maze test.
which assessed the spatial working memory ability, compared to the control (Fig. 4b). However, no significant changes were observed in the NORT or Y-maze test, resulting from the genetic loss of Chdh activity (Supplementary Fig. 6), potentially due to the supplementation of betaine from the chow and/or intestinal bacterial flora. There were no significant differences between the betaine-administered and control groups in B6N or C3HN animals in the other behavioral tests, and importantly no behavioral measures were impaired by betaine treatments in both strains (data not shown). No gross abnormalities in blood chemistry were also observed (data not shown).

Transcriptome analysis of the frontal cortex revealed that the patterns of gene expression caused by betaine were distinct between the two strains, presumably stemming from the underlying genetic differences and aligning with their behavioral differences (Figs. 4c, d, and Supplementary Table 6). Interestingly, the principal component analysis (PCA) of transcriptome data revealed clear segregation of mouse strains in relation to betaine supplementation (based on principal components 1 and 2) (Fig. 4e). The principal components 3 and 4 could clearly discriminate the effect of betaine supplementation irrespective of strain differences (Fig. 4f). Gene ontology enrichment analysis of genes whose expression was positively correlated with PCA loading factor in these principal components revealed biological processes relevant for the cognition, memory, and neurodevelopment related terms (Supplementary Table 7). Meanwhile, certain common molecular phenotypes were also evident (Supplementary Fig. 7, Supplementary Table 8, and Supplementary Table 9). These results highlight an interaction of genetic background × drug (betaine). Gene ontology analysis of upregulated genes by betaine supplementation in both strains identified mitogen-activated protein kinase (MAPK) signaling cascade (Supplementary Table 8), where relevant genetic impairments have been documented in schizophrenia [34].

3.8. Effects of betaine supplementation on MAP-induced behavioral sensitization and gene expression

We next examined whether betaine could perform psychotropic action in an animal model of schizophrenia by leveraging behavioral sensitization paradigm, where repeated administration of psychostimulants in rodents can enhance the stimulating effect on locomotor activity. This behavioral paradigm is assumed to model positive symptoms of schizophrenia and the relapse process [35]. It is of note that peripheral betaine concentration in schizophrenia was inversely correlated with positive symptom scores of the PANSS (Positive and Negative Syndrome Scale) test [3]. Water or betaine (2.17%) was administered 3 weeks before methamphetamine (MAP) injection throughout to the day of MAP challenge (Fig. 5a). In the B6N mice, the challenge injection of MAP elicited enhanced locomotor activities compared to those of the first MAP administration in both water and betaine groups (Fig. 5b). Remarkably, betaine suppressed the degree of elevation in the locomotor activities, which was counted as the differences between the challenge MAP-induced activities and the first MAP-induced activities (p < 0.01) (Fig. 5c). In the C3HN mice, repeated MAP injections did not evoke a marked locomotor sensitization (Fig. 5d). Accordingly, the effect of betaine on the suppression of behavioral sensitization was not clear in the C3HN mice (Fig. 5e), demonstrating the different pharmacogenetic profiles between the two strains.

While the gene expression analyses revealed that betaine deficits and supplementation influenced multiple schizophrenia-relevant transcriptome signatures, we specifically focused on the gene expressions relevant to oxidative stress and neuroinflammation in the MAP model, because [1] such mechanism has been presumed as a pathophysiological component of behavioral sensitization [36] and schizophrenia [37], and [2] betaine’s potency of antioxidant/anti-inflammatory activity is proposed (Supplementary Fig. 1) [8]. It is suggested that betaine could promote nonenzymatic antioxidant activity through the accelerated turnover of methionine in the methionine–homocysteine cycle. Methionine can act as a scavenger of reactive oxygen species (ROS) [38], and oxidized methionine, methionine sulfoxide, can be reduced back to methionine by methionine sulfoxide reductase A (MSRA), B1 (MSRB1) and B2 (MSRB2) (Fig. 1) [39]. Repeated MAP pretreatments coordinate dampened the expressions of antioxidant genes (Gpx1, Gpx4, Sod1, Nos2, Msrb1 and Msrb2 and Mpr1) (p < 0.05) induced by the MAP challenge, and in contrast upregulated the proinflammatory gene Nos2 (p < 0.05), from the comprehensive panel of genes tested (Fig. 5f, Supplementary Table 10 and Supplementary Fig. 8). Betaine cotreatment antagonized the effects of repeated MAP pretreatments on the expression of multiple antioxidant genes (p < 0.05 for Gpx4, Sod1, Msrb1 and Msrb2; p = 0.05 for Mpr1) (Fig. 5f). Betaine cotreatment also revealed a trend towards lowered response of proinflammatory gene expression (Nos2, p = 0.06) after the MAP challenge (Fig. 5f). These results suggest that betaine’s action against MAP-induced sensitization involves, at least in part, antioxidant/proinflammatory response system.

3.9. Altered oxidative stress and proinflammatory conditions, and working memory deficit were rescued by betaine administration in vitro and in vivo phencyclidine models

We further pursued betaine’s action against oxidative stress conditions in vitro phencyclidine (PCP) model. Chronic administration of PCP, a noncompetitive N-methyl-D-aspartate (NMDA) receptor blocker, produces schizophrenia-like behaviors in humans [40]. This psychotomimetic effect is known to be partly associated with oxidative stress given by PCP [19]. Because the deteriorating effect of PCP on neuronal system is also observed in the primary neuron culture [41], we examined the expression changes of the same genes as in the MAP treatment experiments, plus Sod3 (this gene is expressed in rat brain but not in mouse brain), in this paradigm (Fig. 6a). Among those genes, the expressions of Nfe2L2, Rela, Cat, Gpx1, Sod2, Sod3, Cth, Tnf, Nfk1b1 and Nos2 showed coordinate upregulation (p < 0.05) under the PCP (1 μM) exposure (Fig. 6b, also see Supplementary Fig. 9). And notably, the addition of betaine (500 μM) led to significant expression suppression (Fig. 6b). These results demonstrate again a role of betaine in the antioxidant and proinflammatory system.

As a next step, we evaluated the effect of betaine on behavior in an in vivo PCP model (Fig. 6c). ICR mice chronically treated with PCP showed significantly reduced novel object recognition preference in the NORT paradigm as previously reported [19], most likely representing a working memory deficit (Fig. 6d). Intriguingly, the pretreatment of mice with betaine ameliorated the damaging effect of PCP (p < 0.001) (Fig. 6d).

3.10. Analyses of betaine levels and carbonyl stress in postmortem brain samples

Next, we asked whether betaine-related pathology can be seen in schizophrenia. Interestingly, reduced levels of betaine were observed in the postmortem brain tissue from patients with schizophrenia when compared to the controls (Fig. 7a), while the other metabolites
Fig. 5. Betaine suppressed methamphetamine-induced behavioral sensitization and restored altered oxidative stress/neuroinflammatory conditions. (a) Methamphetamine (MAP) sensitization tests were conducted with (+) or without (−) chronic betaine supplementation in two inbred strains, B6N and C3HN. The activities by first-day injection and challenge injection, and the differences in the locomotor activities “Challenge – Day 1” [(locomotor activity induced by challenge injection) − (locomotor activity induced by first-day injection)] are presented (b−e). Betaine supplementation suppressed MAP-induced sensitization in B6N mouse (b, c); whereas, C3HN was not susceptible to the MAP-induced sensitization when compared to B6N (d, e). Data represent mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001; Bonferroni’s multiple comparison test after repeated ANOVA in (b, d), and Student t-test in (c, e); n = 12 per group. (f) After B6N mice received repeated MAP injections followed by challenge injection, expressions of antioxidant and proinflammatory genes in the frontal cortex were examined. Data represent mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001; Bonferroni multiple comparison test between two preset pairs: Cont vs. MAP, and MAP vs. MAP+B; n = 5−6, per group. Cont, control (H2O + repeated saline + MAP challenge); MAP, methamphetamine (H2O + repeated MAP + MAP challenge); MAP+B, methamphetamine + betaine (betaine + repeated MAP + MAP challenge).
remained unchanged (Figs. 7b-g) [15–17] (Supplementary Table 11). SAM contents showed a decreased trend \( (p < 0.1) \), whereas SAH levels were unaltered in the schizophrenia group, resulting in reduced SAM/SAH ratio in the schizophrenic brain \( (p < 0.01) \) (Figs. 7h-j), which can be linked to the decreased betaine levels (Fig. 1). The betaine levels or the SAM/SAH levels were not affected by the amounts of antipsychotics taken (Supplementary Table 12).

To investigate the relationship between betaine system abnormality and oxidative stress conditions in schizophrenia, we specifically focused on the carbonyl stress in the postmortem brain samples. AGES analysis remained unchanged (Figs. 7b-g) [15–17] (Supplementary Table 11). SAM contents showed a decreased trend \( (p < 0.1) \), whereas SAH levels were unaltered in the schizophrenia group, resulting in reduced SAM/SAH ratio in the schizophrenic brain \( (p < 0.01) \) (Figs. 7h-j), which can be linked to the decreased betaine levels (Fig. 1). The betaine levels or the SAM/SAH levels were not affected by the amounts of antipsychotics taken (Supplementary Table 12).

To investigate the relationship between betaine system abnormality and oxidative stress conditions in schizophrenia, we specifically focused on the carbonyl stress in the postmortem brain samples. AGES analysis
identified the three schizophrenia brain samples (#SZ01, #SZ05 and #SZ12) that showed a strong band (~30 kDa protein(s)) having Nε-(carboxyethyl)lysine (CEL) (Figs. 7k and l), one of AGEs [12]. The three samples showed betaine levels of below average (Fig. 7a), although the correlation of the two measures (betaine levels and CEL modification levels) did not fulfill a statistical significance possibly due to the sample number limitation. This finding suggests a relationship between elevated carbonyl stress and lowered betaine levels. Furthermore, these three patients displayed relatively high symptom scores evaluated by using the Diagnostic Instrument for Brain Studies (DIBS) [42] (Supplementary Table 13). Collectively, the results of postmortem brain study highlighted the existence of a subset of patients with schizophrenia characterized by “betaine deficit-oxidative stress” pathology, manifesting relatively severe psychotic symptoms.

3.11. Betaine’s potency in alleviating carbonyl stress in GLO1-deficient hiPSCs

Because coexistent betaine pathology and elevated carbonyl stress were evident in a subset of schizophrenia cases, the efficacy of betaine supplementation in mitigating the carbonyl stress in isogenic lines of GLO1-deficient hiPSCs was tested. We firstly confirmed that there were no missense variants in the genome of the subject (mentally healthy) for the genes in the carbonyl stress pathway (Supplementary...
Fig. 8. Betaine suppressed accumulation of advanced glycation products (AGEs) in hiPSCs and a genetic variant affected CHDH expression in brain. (a) Genomic structure of the human GLO1 gene is schematically presented. Black and white boxes indicate coding and untranslated regions of exons, respectively. Three hiPSC lines for each genotype (WT-1-3 and KO-1-3) were established. (b) Phase contrast images of WT-1 and KO-1 are presented. WT-2-3 and KO-2-3 showed morphology similar to WT-1 and KO-1 (data not shown). Scale bar: 400 μm. (c) Cell (WT-1 and KO-1) lysates were analyzed in western blotting using an antibody against carboxymethyllysine (CML), a typical type of AGEs. The filter was also probed with anti-GAPDH antibody as a loading control and anti-GLO1 antibody to show a loss of the GLO1 protein in the GLO1KO (−/−) cells. (d) KO-1 was cultured for four days in the absence or presence of indicated concentrations (5, 50, or 500 μM) of betaine. The results of western blot analysis of the cell lysates are presented (top). The position for “band A” is indicated by an arrowhead. The same experiment was repeated using KO-2-3 and WT-1-3 in duplicate. A significant difference (n= six per group; 3 KO or WT lines × duplicate) was observed between the groups treated with 0 μM and 500 μM betaine. The CML signals were normalized to the intensity of GAPDH signals. Data represent mean ± SEM. *p < 0.05; Dunnett multiple comparison test. n= six. (e) Contents of metabolites in the methionine-homocysteine cycle in the GLO1KO hiPSCs. Data represent mean ± SEM. **p < 0.01, ***p < 0.001; two-tailed Student's t-test. WT-1-3 and KO-1-3 were triplicated (n= nine each). (f) CHDH variant, rs35518479, showed a significant association for its expression and SAM/SAH ratio in the brain, though betaine levels were unaltered. Data represent mean ± SEM. *p < 0.05, **p < 0.01, ***p < 0.001; one-way ANOVA and two-tailed Student's t-test.
Fig. 2). The GLO1-deficient hiPSC lines were prepared by harnessing the CRISPR-Cas9 system, and they contained homozygous GLO1 KO alleles (KO-1, KO-2 and KO-3) causing frameshift mutations in exon 1 and premature stop signals (Figs. 8a and b), and showed no protein expression (Fig. 8c). As controls, three lines (WT-1, WT-2 and WT-3) were established that were proven to harbor the normal GLO1 alleles. The GLO1-deficient hiPSC samples displayed a major band of ~55 kDa (called “band A”) with increased modification by carboxymethyllysine (CML), another species of AGEs [12] (representative picture in Fig. 8c). When betaine was added to cell culture, concentration at 500 μM significantly suppressed the CML modification of “band A” (Fig. 8d), indicating betaine’s efficacy against elevated carbonyl stress. The GLO1-disrupted hiPSCs showed comparable morphology (Fig. 8b) with those of the WT, but they manifested reduced differentiation potential into neural lineage, which could not be rescued by the betaine supplementation (data not shown).

Next, we examined the effects of GLO1 ablation on the levels of betaine and metabolites in the methionine-homocysteine cycle. Strikingly, betaine levels were significantly decreased in the GLO1 KO hiPSCs compared to the WT cells (Fig. 8e), suggesting that betaine is consumed to cope with increased carbonyl stress. In parallel, lowered SAM/SAH ratio concomitant with increased SAH levels in the KO cells was seen (Fig. 8e), probably reflecting dampened intracellular homocysteine clearance by BHMT pathway (Fig. 1) in the KO cells.

3.12. Pharmacogenetic evaluation of betaine efficacy in schizophrenia

We have revealed the differential efficacy for betaine in the mouse strains with distinct genetic background (Fig. 5). Since a subset of schizophrenia patients were characterized with betaine deficiency and oxidative stress pathology and the betaine treatment was effective in alleviating carbonyl stress, we speculated that the efficacy for betaine might have a genetic predisposition. Cis-eQTL analysis of common genetic variants in BHMT1, CHDH and GLO1 genes in postmortem brain tissues revealed a significant association of a CHDH variant, rs35518479, with its expression levels in the brain (p = 0.03) (Fig. 8f, Supplementary Figs. 10a and b). The A allele carriers showed a significantly higher CHDH expression levels in the brain when compared to the G/G homozygotes. Although there were no significant differences in betaine and SAM levels according to the genotypes (Supplementary Fig. 10c), the SAM/SAH ratio (methylation index) was higher in the A allele carriers (p = 0.02) (Fig. 8f). The results suggest that CHDH expression levels may affect the turnover rate of methionine-homocysteine cycle, and depict a potential utility of CHDH-eQTL in stratifying schizophrenia patients for betaine treatment.

4. Discussion

In the present study, we provided a proof of concept for the potential of betaine in the treatment of schizophrenia from the analyses of both mouse and human samples, along with the identification of “betaine deficit-oxidative stress” pathology in a subset of schizophrenia, which may result in relatively severe symptoms. Our systematic examination of betaine’s role (see Supplementary Fig. 1) showed that the betaine’s action as a psychotropic is imparted partially through antioxidant/pro-inflammatory effects. In addition, a regulation of DNA methylation capacity may be possible, from metabolomics/methylome analyses of mouse and human samples. Notably, analyses of two different inbred mouse strains highlighted an interaction of genetic background × drug (e.g. betaine), and distinct pharmacogenetic profiles (e.g. MAP) according to genetic background. Also, no differences in the brain betaine levels were observed between the two strains, before and after betaine supplementation (data not shown), ruling out strain (genetic) differences in brain betaine turnover.

Although the more detailed mechanism of betaine’s biological effects remains to be clarified in the future study, it is worth noting that methionine, the primary enzymatic product of BHMT (betaine-homocysteine methyltransferase), functions as a powerful antioxidant molecule. Furthermore, it should be pursued whether the expression modulation of genes for antioxidant/proinflammatory cascades by the betaine treatment is mediated via epigenetic changes elicited by the methylation of histone or DNA, where SAM is used as a methyl donor.

Betaine, tri-methylglycine, is very simple in the chemical structure. It receives step-wise breakdown to glycine in mammals: betaine to di-methylglycine, di-methylglycine to mono-methylglycine, and mono-methylglycine to glycine (Fig. 1). It is unlikely that these downstream compounds may play a positive role to exert betaine’s effect, because di- and mono-methylglycine were under the detection limit (data not shown) in mouse and postmortem human brain samples, probably reflecting very rapid degradation of di-methylglycine into glycine by demethylation. In addition, the concentration of glycine seemed relatively stable even by the manipulation of the Chdh gene or the supplementation of betaine in mice, excluding the idea stated above.

In this study, we detected a substantial accumulation in CEL adduct of ~30 kDa protein(s) (Fig. 7k) and CML adduct of ~55 kDa protein(s) (Fig. 8c) in postmortem brain and GLO1-KO iPSCs, respectively. It was unexpected to see AGEs in a relatively protein-specific manner under carbonyl stress, since the Maillard reaction (Supplementary Fig. 2) is deemed to occur non-enzymatically [12]. Molecular identification and characterization of the specifically modified proteins and the mechanism of the specific modifications remain to be studied.

Although we did not address in this study, another known function of betaine is osmotic regulation (Supplementary Fig. 1), and the deficits in brain betaine levels may contribute to cellular osmotic perturbation [6,7,43], which is reported to inhibit methionine uptake, inhibit protein synthesis, and affect the mRNA translation by dysregulation of phosphorylation and mTOR signaling cascades [44–46]. These pathways are shared with those identified in the transcriptomics analysis of Chdh-deficient mice (Fig. 3f). Although taurine, myo-inositol, glycine and glutamine are thought to play major roles as osmolytes in the brain because of their higher cellular contents; betaine is favored in neural cells as an osmolyte [6]. Thus, the osmotic issue should be pursued in future studies.

Since betaine has already been approved as a therapeutic drug for homocystinuria, an autosomal recessive inherited disorder due to a deficiency of cystathionine beta synthase (CBS, Fig. 1) [47], it has the advantage of being repositioned for psychiatric illness. The utility is also substantiated by our observation that betaine can penetrate the blood-brain barrier and is well tolerated with no serious adverse effects.

Interestingly, we identified an eQTL for CHDH expression, which substantiates genotype-based stratification of schizophrenia patients for betaine efficacy. Future studies are warranted to validate the utility of “betaine deficit-oxidative stress pathology” as a biomarker and to identify other genetic underpinnings for betaine’s efficacy. It is also worthwhile to test the availability of betaine in clinical setting for schizophrenia and other neuropsychiatric conditions.

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Declaration of interests

The authors report no biomedical financial interests or potential conflicts of interest.

Author contribution

TO, SB, and TY designed the study. TO, SB, MT, MM, HO, AW, YI, YF, YT, YS, YH, CS-M, YN, YH, KE, NM, and AH-T performed the experiments. AN, JM, MH, YK, AK, and HY collected the postmortem brain samples and clinical information. TO, SB, MT, HO, AW, and YI acquired the data, TO, SB, MT, AW, YI, KH, and TY analyzed the data, and TO, SB, and TY wrote the manuscript.

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Appendix A. Supplementary data

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References

[1] Carpenter Jr WT, Davis JM. Another view of the history of antipsychotic drug discovery and development. Mol Psychiatry 2012;17(12):1168–73.
[2] Molins C, Roldan A, Corripio I, Isohanni M, Miettunen J, Seppala J, et al. Response to polygenic association analysis of 27 genes from the GABAergic system in Japanese individuals affected with schizophrenia. Schizophr Res 2017;185:33–40.
[3] Watanabe A, Toyota T, Owada Y, Hayashi T, Iwayama Y, Matsumata M, et al. Fabp7 maps to a quantitative trait locus for a schizophrenia endophenotype. PLoS Biol 2007;5(11):e229.
[4] Hoffman DR, Cortnerse WR, Duerr JA. Relationship between tissue levels of S-adenosylhomocysteine, S-adenosylmethionine, and transmethylation reactions. Can J Biochem 1979;57(1):56–65.
[5] James SJ, Melnicky S, Pogribnina M, Pogribny I, Caudill MA. Elevation in S-adenosylmethionine in brain, kidney, and liver. Front Physiol 2014;5:159.
[6] Watanabe A, Toyota T, Owada Y, Hayashi T, Iwayama Y, Matsumata M, et al. Betaine uptake, its influence on other Osmolytes and its potential role in neuroprotection from osmotic stress. Neurochem Res 2017;42(12):3490–503.
[7] Wehner DG, Al-Abed I, Early BR, Liotta LA, Fahey JL, et al. Unique metabolomics signature in plasma: evidence that early intervention may impact on disease progression and outcome in schizophrenia. PLoS Med 2006;3(8):e327.
[8] Balan S, Yamada K, Iwayama Y, Hashimoto T, Toyota T, Shimamoto C, et al. Comprehensive analysis of 27 genes from the GABAergic system in Japanese individuals affected with schizophrenia. Schizophr Res 2017;185:33–40.
[9] Nemoto T, Kasa K, Ishii H, Kawauchi H, Hoshi Y, et al. A snapshot of plasma metabolites in first-episode schizophrenia: a capillary electrophoresis time-of-flight mass spectrometry study. Transl Psychiatry 2014;4:e0379.
[10] Balan S, Toyota T, Owada Y, Hayashi T, Iwayama Y, Matsumata M, et al. Betaine synthesis in schizophrenia patient-derived olfactory cells. Transl Psychiatry 2015;5:e663.
[11] Khanmohammadi S, Khanal A, Tsichas K, Ujike H. Methamphetamine-induced behavior sensitization and its implications for relapse of schizophrenia. Schizophr Res 1994;12(3):251–7.
[12] Shi J, Wang Y, Zhang S, Wang Z, Li Y, et al. Enhanced carbonyl stress in a subpopulation of schizophrenia. Arch Gen Psychiatry 2010;67(6):589–97.
[13] Miyashita M, Arai M, Kobori A, Ichikawa T, Toriumi K, Nizato K, et al. Clinical features of schizophrenia with enhanced carbonyl stress. Schizophr Bull 2014;40(5):576–86.
[14] Hino M, Kunii Y, Matsumoto J, Wada A, Nagaoka A, Niwa S, et al. Decreased VEGFR2 expression and increased phosphorylated Akt1 in the prefrontal cortex of individuals with schizophrenia. J Psychiatr Res 2016;82:100–8.
[15] Chirot S, Miura J, Matsumoto J, Hino M, Wada A, Niwa S, et al. Elevated postmortem striatal t-DARPP expression in schizophrenia and associations with DRD2/ANKK1 polymorphism. Prog Neuropsychopharmacol Biol Psychiatry 2014;53:123–33.
[16] Hino M, Kunii Y, Niwa J, Matsumoto J, Hino M, Wada A, Niwa S, et al. Decreased VEGFR2 expression and increased phosphorylated Akt1 in the prefrontal cortex of individuals with schizophrenia. J Psychiatr Res 2016;82:100–8.
[17] Matsumoto J, Nakashima H, Kunii Y, Sugiuara Y, Yuki D, Wada A, et al. Decreased 16:0/20:4-phosphatidylcholine levels in the postmortem prefrontal cortex of elderly patients with schizophrenia. Sci Rep 2017;7:45050.
[18] Shimamoto C, Oshi N, Maekawa M, Watanabe A, Ohba H, Arai R, et al. Functional characterization of FABP3, 5 and 7 gene variants identified in schizophrenia and autism spectrum disorder and mouse behavioral phenotypes. Hum Mol Genet 2014;23(24):e012744.
[19] Saito T, Yamao S, Ota M, Egashira T, Yae K, Kuwamoto D, et al. Generation of induced pluripotent stem cells from human terminally differentiated circulating T cells. Cell Stem Cell 2010;7(1):11–4.
[20] Matsumoto J, Nakanishi H, Kunii Y, Sugiura Y, Yuki D, Wada A, et al. Enhanced postmortem striatal t-DARPP expression in schizophrenia and associations with DRD2/ANKK1 polymorphism. Prog Neuropsychopharmacol Biol Psychiatry 2014;53:123–33.
[21] Shimamoto C, Oshi N, Maekawa M, Watanabe A, Ohba H, Arai R, et al. Functional characterization of FABP3, 5 and 7 gene variants identified in schizophrenia and autism spectrum disorder and mouse behavioral phenotypes. Hum Mol Genet 2014;23(24):e012744.
[22] Seki T, Yamao S, Ota M, Egashira T, Yae K, Kuwamoto D, et al. Generation of induced pluripotent stem cells from human terminally differentiated circulating T cells. Cell Stem Cell 2010;7(1):11–4.
[23] Watanabe A, Toyota T, Owada Y, Hayashi T, Iwayama Y, Matsumata M, et al. Betaine synthesis in schizophrenia patient-derived olfactory cells. Transl Psychiatry 2015;5:e663.
[24] Watanabe A, Toyota T, Owada Y, Hayashi T, Iwayama Y, Matsumata M, et al. Betaine synthesis in schizophrenia patient-derived olfactory cells. Transl Psychiatry 2015;5:e663.
[25] Boch J, Kempf B, Schmid R, Bremer E. Synthesis of the osmoprotectant glycine betaine in Bacillus subtilis: characterization of the gbsAB genes. J Bacteriol 1996;178(17):5312–9.
[26] Lui WY, Muuk D, Lee WM, Cheng CY. Sertoli cell tight junction dynamics: their regulation during spermatogenesis. Biol Reprod 2003;68(4):1087–97.
[27] English JA, Fan Y, Fockey M, Lopez LM, Hryniewiecka M, Wynne K, et al. Reduced betaine synthesis in schizophrenia patient-derived olfactory cells. Transl Psychiatry 2015;5:e663.
[43] Kempson SA, Montrose MH. Osmotic regulation of renal betaine transport: transcription and beyond. Pflugers Arch 2004;449(3):227–34.

[44] Uesono Y, Toh EA. Transient inhibition of translation initiation by osmotic stress. J Biol Chem 2002;277(16):13848–55.

[45] Patel J, McLeod LE, Vries RG, Flynn A, Wang X, Proud CG. Cellular stresses profoundly inhibit protein synthesis and modulate the states of phosphorylation of multiple translation factors. Eur J Biochem 2002;269(12):3076–85.

[46] Aramburu J, Ortells MC, Tejedor S, Buxade M, Lopez-Rodriguez C. Transcriptional regulation of the stress response by mTOR. Sci Signal 2014;7(332):re2.

[47] Wilcken DE, Wilcken B, Dudman NP, Tyrrell PA. Homocystinuria—the effects of betaine in the treatment of patients not responsive to pyridoxine. N Engl J Med 1983;309(8):448–53.