

Altered glutathione redox state in schizophrenia

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Abstract. Altered antioxidant status has been reported in schizophrenia. The glutathione (GSH) redox system is important for reducing oxidative stress. GSH, a radical scavenger, is converted to oxidized glutathione (GSSG) through glutathione peroxidase (GPx), and converted back to GSH by glutathione reductase (GR). Measurements of GSH, GSSG and its related enzymatic reactions are thus important for evaluating the redox and antioxidant status. In the present study, levels of GSH, GSSG, GPx and GR were assessed in the caudate region of postmortem brains from schizophrenic patients and control subjects (with and without other psychiatric disorders). Significantly lower levels of GSH, GPx, and GR were found in schizophrenic group than in control groups without any psychiatric disorders. Concomitantly, a decreased GSH:GSSG ratio was also found in schizophrenic group. Moreover, both GSSG and GR levels were significantly and inversely correlated to age of schizophrenic patients, but not control subjects. No significant differences were found in any GSH redox measures between control subjects and individuals with other types of psychiatric disorders. There were, however, positive correlations between GSH and GPx, GSH and GR, as well as GPx and GR levels in control subjects without psychiatric disorders. These positive correlations suggest a dynamic state is kept in check during the redox coupling under normal conditions. By contrast, lack of such correlations in schizophrenia point to a disturbance of redox coupling mechanisms in the antioxidant defense system, possibly resulting from a decreased level of GSH as well as age-related decreases of GSSG and GR activities.

Keywords: Glutathione, glutathione disulfide, glutathione peroxidase, glutathione reductase, cigarette smoking, age, postmortem caudate, schizophrenia

1. Introduction

Biological systems have evolved complex protective strategies against free radical toxicity [1]. Under physiological conditions the potential for free radical-mediated damage is kept in check by the antioxidant defense system, comprising a series of enzymatic and

non-enzymatic components (Fig. 1). The key antioxidant enzymes include superoxide dismutase (SOD), catalase (CAT), and glutathione peroxidase (GSH-Px). Superoxide dismutase catalyzes the conversion of superoxide radicals ($O_2^- \bullet$) to hydrogen peroxide (H_2O_2), which in turn can form the highly reactive hydroxyl radicals. Catalase and glutathione peroxidase convert hydrogen peroxide to water. Glutathione (GSH) is utilized by GSH-Px to yield the oxidized form of glutathione (GSSG), which is converted back to GSH by glutathione reductase (GR). In addition, nitric oxide (NO) which is the product of a five-electron oxidation

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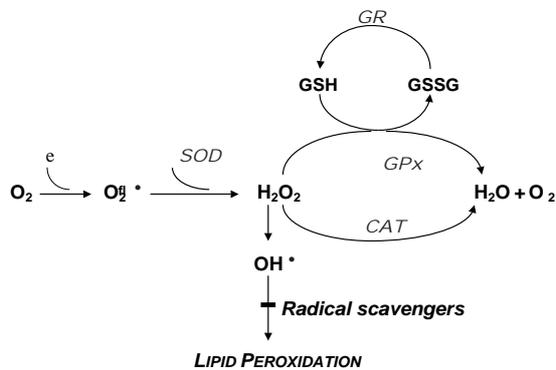


Fig. 1. Production of reactive oxygen species and defense mechanism against damage by reactive oxygen. Abbreviations: $O_2^{\bullet -}$, superoxide radicals; SOD, superoxide dismutase; H_2O_2 , hydrogen peroxide; CAT, catalase; GPx, glutathione peroxidase; GSH, glutathione; GSSG, glutathione disulfide; GR, glutathione reductase, OH^{\bullet} , hydroxyl radicals.

of the amino acid L-arginine, can also produce hydroxyl radicals as well as nitrogen dioxide radical.

There is evidence that free radicals are involved in membrane pathology, and may play a role in schizophrenia [2–4]. Membrane dysfunction can be secondary to free radical-mediated pathology, and may contribute to specific aspects of schizophrenic symptomatology and complications of its treatment. Specifically, free radical-mediated abnormalities may contribute to the development of a number of clinically significant consequences, including prominent negative symptoms, tardive dyskinesia, neurological “soft” signs, poor premorbid function, and CT scan abnormalities. Our previous results showing altered membrane dynamics and antioxidant enzyme activities in schizophrenia, and findings from other investigators are consistent with the notion of free radical-mediated neurotoxicity in schizophrenia [4]. Free radicals are reactive chemical species generated during normal metabolic processes, and in excess, can damage lipids, proteins, and DNA. Regions of high oxygen consumption, lipid content, and transition metals, are at particular risk. Hence, neuronal membranes are uniquely vulnerable to radical-mediated damage.

Studies of the antioxidant defense system are often complicated by cumbersome analytical methods, which require separate and multi-step extraction and chemical reaction procedures. Thus, measurements of multiple parameters are limited in relatively small biological samples. Recently, we [5] have developed a convenient and sensitive tool to measure low molecular weight, redox-active compounds in biological samples using a high-pressure liquid chromatography sys-

tem (HPLC) coupled with a 16-channel Coulometric Multi Electrode Array System (CMEAS). The purpose of the present investigation was to assess the glutathione (GSH) redox system (Fig. 1) by evaluating not only GSH and GSSG levels but also the enzyme activities of GSH-Px and GSH-R simultaneously in the caudate region of postmortem brain samples from patients with schizophrenia as well as controls with and without other psychiatric disorders.

2. Methods

2.1. Postmortem brain tissues

The Schizophrenia Research Center Brain Bank at the VA Medical Center in Denver, Colorado, provided postmortem brain tissue. Human brain was collected at autopsy from local services following family consent.

Psychiatric illness status was determined independently by two board-certified psychiatrists from medical records and structured family interviews. Diagnostic and Statistical Manual IV criteria for schizophrenia [6] (American Psychiatric Association, 1994), were met for the schizophrenic subjects used in the study. Control subjects were also evaluated and determined to have had no evidence of psychiatric illness.

At autopsy, the brain was weighed and examined for gross pathology. It was then divided sagittally and one hemisphere, selected randomly, was preserved in formalin for neuropathological analysis at the macroscopic and microscopic level. Microscopic evaluations included standard Bielschowsky silver stain on multiple cerebral areas to rule out abnormal neuropathology, such plaques and tangles, associated with Alzheimer’s and other conditions. Patients with positive neuritic findings or ambiguous neuropathology reports were excluded from the current study. The hemisphere that was not subjected to neuropathological analysis was sliced coronally into 1 cm slices, from which multiple regions were dissected in 1-gram blocks, frozen in dry ice snow, and packaged for storage at $-80^{\circ}C$. Hemispheres are collected randomly.

An extensive review of hospital, autopsy, and neuropathology reports were conducted to determine agonal and postmortem conditions for each subject. Based upon this information, an agonal state score was assigned following the four-point rating scale described by Hardy et al. [7]. These four categories included 1) violent and fast death, almost always of unnatural causes such as shootings (accidental, homicidal, or sui-

cide) or blunt force trauma; 2) fast death of natural causes, such as the sudden, unexpected deaths of individuals who had appeared reasonably healthy. The most common cause of death in this group was myocardial infarction; 3) intermediate death, usually unexpected, even though the patient was ill prior to death; and 4) slow death, occurring after a long illness. The majority of these patients died of cancer or chronic pulmonary disease.

Additional postmortem parameters, such as age, sex, ethnicity, medication history, psychiatric illness status, cigarette smoking history and alcohol use, were evaluated through hospital, autopsy, and neuropathology reports. Family members and physicians were also interviewed to detail the smoking and alcohol history of the subject, including packs of cigarettes smoked per day, and the quantity and type of alcohol consumed. All of the subjects who had used alcohol prior to death had consumed no more than two drinks per day on average. Smokers had used cigarettes until the time of death.

All patients with schizophrenia and other psychiatric disorders were on antipsychotic medications at the time of death. At the time of autopsy, the brain was removed and hemisectioned in the sagittal plane. One half of the cerebrum, cerebellum, and brainstem was immersed in 10% formalin and fixed for 1–2 weeks. The other hemisphere was sliced coronally and transported on aluminum plates, previously chilled on ice, to the laboratory for dissection. To avoid hemispheric bias, alternate (left/right) hemispheres were collected. Immediately after sample collection, tissue was dissected from approximately 50 areas, in one gram pieces, and frozen on dry ice snow. The remainder of the hemisphere was also frozen on slabs of dry ice, and stored with the dissected parts at -80°C . In the present study, the caudate nucleus from schizophrenic and control groups was used for GSH analyses.

2.2. Laboratory assays

2.2.1. Tissue preparation

Approximately 0.2 g of brain sample was first weighed and lyophilized before the dry weight was taken. Samples were then homogenized by a Polytron tissue homogenizer in one ml of 10 mM Sodium Acetate, pH 6.5. Following centrifugation for 10 minutes at 3,000 rpm, the cell-free homogenate was further passed through a disposable membrane (0.22 μm pore size) micropartition system (Millipore Ultrafree-MC) under centrifugation to remove any compounds above 10,000 nominal molecular weight limit. The resulting protein-free supernatant was subjected to HPLC for GSH and GSSG assays.

2.2.2. Measurements of GSH and GSSG by HPLC-CMEAS

The deproteinized sample was analyzed by an ESA CoulArray Model 5600 HPLC system consisting of two Model 582 pumps, one dynamic gradient mixer, two PEEKTM pulse dampers, a Model 542 refrigerated autosampler injector, a CoulArray organizer module, and a serial array of 16 coulometric electrodes. The system is controlled and chromatograms are analyzed by a Dell personal computer Model Dimension XPS R450 using the ESA CoulArray for Windows-32 software program (version 1.04).

Each sample containing 50 μL protein-free extract was run on a *single* column (ESA Meta-250, 5 μm ODS, 250 \times 4.6 mm ID) with a 150-minute gradient elution that ranged from 0% [Mobile Phase A consisting of 1.1% (w/v) of 1-pentane-sulfonic acid (pH, 3.0)] to 20% Mobile Phase B (MPB, consisting of 0.1 M lithium acetate in a solvent mixture of methanol, acetonitrile and isopropanol, 80/10/10 (v/v/v) with a fixed flow rate of 0.5 ml/min. The temperature of both cells and column was maintained at 25 $^{\circ}\text{C}$. Similar to the previously published method [9,10], the current gradient elution also included a 2 minutes of a high-potential cell-cleaning step to restore the column to 100% MPA at the end of each run. The CMEAS is set to have increments from 0 to 960 mV in 60 mV steps. The properties of various antioxidant compounds or monoamine metabolites are often closely related to the same structural characteristics. The use of several coulometric sensors in series that are maintained at different potentials provides resolution and detection of co-eluting compounds with different electrochemical features.

The ESA CoulArray for Windows 32 package was used for quantitative analysis of peak identity and the peaks relative to absolute or reference standards. This software is able to automatically subtract backgrounds resulting from the gradient drift. The coulometric array is used to generate databases of all redox-active molecules with redox potentials from 0 to 960 mV. Peak identity is verified by the retention time, dominant channel, and the ratio of reactivity on the dominant channel to reactivity on the subdominant channels [9, 10]. The concentration of each peak was calculated according to the dominant channel which has > 70% ratio conformity to contribute to the quantitation [11]. In a typical assay, the standard curve for GSH and GSSG were established between 0.2 to 200 ng that provides three orders of magnitude. Within the range of the standard curve, the coefficients of variation (CV) for within-run assays were in the range of 0.9–6.0% and for between-run assays were in the range of 1.0–9.1%.

2.2.3. Glutathione peroxidase (GPx) assay

Approximately 200 mg of brain tissue was homogenized in 1.5 ml of 50 mM Tris-HCl Buffer containing 1 mM EDTA, pH 7.4. Following centrifugation of the homogenate for 10 minutes at 8500 rpm, aliquots of supernatant between 0.1 to 1.0 mg proteins were used for enzyme assays.

The Cellular Glutathione Peroxidase Assay Kit (Cat. No. 354104) from Calbiochem was used to measure GPx activity indirectly. In principal, GSSG that is produced upon reduction of organic peroxide by cytoplasmic GPx is recycled to its reduced state by glutathione reductase (GR). The oxidation of NADPH to NADP⁺ is accompanied by a decrease in absorbance at 340 nm, which provides a spectrophotometric means of monitoring GPx activity.

2.2.4. Glutathione reductase (GR) assay

The Glutathione Reductase Assay Kit from Calbiochem (Cat. No. 359962) was used to measure GR activity. Briefly, the GSSG is reduced by a multi-step reaction in which GR is initially reduced by NADPH producing a semiquinone of FAD, a sulfur radical and a thiol. The reduced GR reacts with a molecule of GSSG resulting in a disulfide interchange, which produces a molecule of GSH and the GSSG-R_{red}-SG complex. An electron rearrangement in GSSG-R_{red}-SG results in a second disulfide interchange, splitting off the second molecule of GSH and restoring the GR to the oxidized form. The GR activity was thus measured by the rate of NADPH oxidation, which is accompanied by a decrease in absorbance at 340 nm. Since GR is present at rate-limiting concentration, the rate of decrease in the A₃₄₀ is directly proportional to the GR activity in the sample.

2.3. Statistical analyses

Multiple regression analyses were conducted to determine whether the brain collection and storage variables (age, postmortem interval, brain weight, and storage time) were significantly associated with the biochemical measures of interest. Separate regression analyses were conducted for each of the biochemical measures. The brain collection and storage measures were entered into each of the regression analyses as predictor variables. The distributions for all variables were examined using normality plots and the Kolmogorov-Smirnov test, which quantifies the discrepancy between data distribution and an ideal Gaussian distribution. The Dallal and Wilkinson approximation to Lilliefors'

method was used to compute the P values. The data pass the normality test if $p > 0.10$. For normally distributed data, means and variances were compared by the use of *t*-tests and *F*-tests.

One-way analyses of variance (ANOVA) were conducted to evaluate group differences for brain collection and storage parameters between schizophrenics, controls with and without other psychiatric disorders. Analyses of covariance (ANCOVA) were conducted to control statistically for the potential effects of the brain collection and storage variables on the group differences for the biochemical measures of interest. Group means for the GSH redox coupling measures were compared using ANOVAs, with pairwise comparisons (unpaired *t*-tests) following significant findings.

3. Results

3.1. Brain collection and storage parameters

One-way analyses of variance (ANOVA) were conducted to evaluate group differences for the brain collection and storage parameters (Table 1). There are no significant differences of age, postmortem interval (PMI), and storage time among three test groups with exception that brain weights was found significantly higher in schizophrenic than in control samples without psychiatric disorders. The normality plots and the Kolmogorov-Smirnov test indicated that normality transformations of the data were not necessary. On the other hand, as expected, the frequency of smoking was higher in patients with either schizophrenia or bipolar and/or depression than in control subjects without psychiatric disorders. In addition, the ratio of male to female subjects was also higher in the patient samples than in control samples without psychiatric disorders.

3.2. Evaluations of glutathione redox state

To evaluate GSH redox state, GSH and GSSG levels were measured by HPLC coupled with a 16-channel Coulometric Electrode Array System. GSH-Px and GSSG-R activities were assayed by a spectrophotometric procedure in the caudate region of postmortem brains from schizophrenic patients and control subjects with and without other psychiatric disorders (Table 2). GSH levels were significantly ($p = 0.017$) lower in patients with schizophrenia than in control subjects without any psychiatric disorders. A similar reduction of GSH level was also found in patients with bipolar

Table 1
Brain collection and storage parameters

Parameters	Schizophrenics	Controls	
		without PD ^a	with other PD
Number of samples	12	16	10
Age, years	60 ± 10 ^b	54 ± 17	50 ± 17
Sex (M/F)	9/3	10/6	9/1
Smoking (yes/no)	11/1	10/6	8/2
Postmortem interval (h)	20 ± 14	15 ± 7	16 ± 7
Brain weight (g)	1404 ± 168 ^c	1282 ± 143	1398 ± 77
Storage time (months)	17 ± 8 ^d	11 ± 8	18 ± 8

^aPD, psychiatric disorders such as bipolar and depression.

^bEach value represents the mean and standard deviation based on the number of examined brain samples.

^cANOVA ($F = 3.4482$; $df = 2.35$; $p = 0.0430$).

^dANOVA ($F = 2.9450$; $df = 2.45$; $p = 0.0658$).

Table 2

Evaluation of the glutathione redox system in the caudate region of postmortem brain tissue between schizophrenic and control groups

Mercaptans	Schizophrenics ($n = 12$)	Controls ($n = 16$)		P (two-tailed, unpaired t tests)		
		without PD ^a ($n = 16$)	with other PD ($n = 10$)	S vs C	S vs C-PD	C vs C-PD
GSH ^b	107 ± 61 ^c	179 ± 89	115 ± 99	0.017	0.838	0.110
GSSG ^b	35 ± 24	33 ± 15	30 ± 25	0.870	0.632	0.670
GPx	4658 ± 1199	5825 ± 1775	6294 ± 1210	0.048	0.005	0.432
GR	685 ± 142	794 ± 103	795 ± 218	0.036	0.194	0.997

^aAbbreviations: PD, psychiatric disorders; S, schizophrenics; C, control; C-PD, control with psychiatric disorder; GSH, glutathione; GSSG, glutathione disulfide; GPx, glutathione peroxidase; GR, glutathione reductase.

^bBoth GSH and GSSG were determined by HPLC-CMEAS procedure.

^cGSH and GSSG were expressed as ng/mg dry wt., whereas GPx and GR were expressed as mU/mg wet wt.

and/or depression (controls with other types of psychiatric disorders) although it was not statistically significant. On the other hand, there were no significant differences of GSSG levels among the three groups of subjects. Consequently, the ratio of GSH to GSSG was significantly ($p = 0.0085$) lower in patients with schizophrenia than in control subjects without any psychiatric disorders (Fig. 2), but not in control subjects with bipolar and/or depression.

Both GPx and GR activities were found significantly ($p < 0.05$) lower in patients with schizophrenia than in control subjects without any psychiatric disorders (Table 2). Moreover, GPx activities in patients with schizophrenia was also significantly ($p = 0.005$) lower than that of patients with bipolar and/or depression. However, no significant differences were found in either GPx or GR between two control groups.

To further assess the GSH redox coupling reactions, levels of substrate (GSH) were correlated with the specific enzyme activities (GPx or GR). In control subjects without any psychiatric disorders, GSH levels were significantly ($p = 0.0088$) correlated to GPx activities (Fig. 3). A similar positive correlation ($p = 0.0052$)

GSH:GSSG ratio

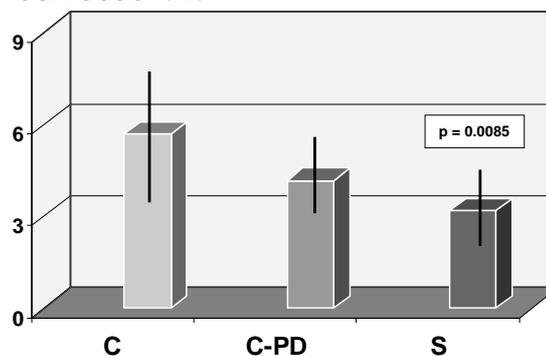


Fig. 2. Comparison of ratio of glutathione (GSH) to glutathione disulfide (GSSG) in postmortem caudates among control subjects without psychiatric disorders (C), control subjects with other psychiatric disorders (C-PD), and patients with schizophrenia (S). $p = 0.0085$ (C vs S, two-tailed unpaired t-test).

was also demonstrated between GSH levels and GR activities (Fig. 4). Furthermore, GPx activities were significantly ($p = 0.0226$) correlated to the GR activities in the same control group (Fig. 5). No significant correlations were found in patients with either schizophrenia

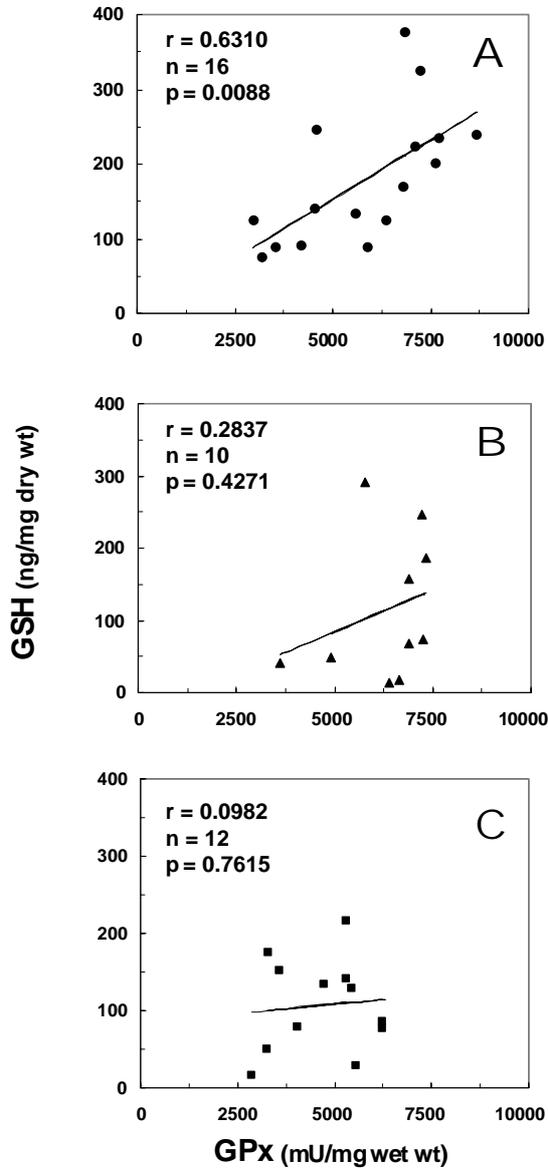


Fig. 3. Correlations of glutathione (GSH) levels to glutathione peroxidase (GPx) activities in postmortem caudate region from control subjects without psychiatric disorders (A), control subjects with bipolar and/or depression (B), and patients with schizophrenia (C).

or bipolar and/or depression (Figs 3–5).

3.3. Effect of gender and smoking status in GSH redox state

Although there are more female subjects in the control samples without psychiatric disorders than in patient sample groups, no significant differences of any GSH redox measures were found in control samples

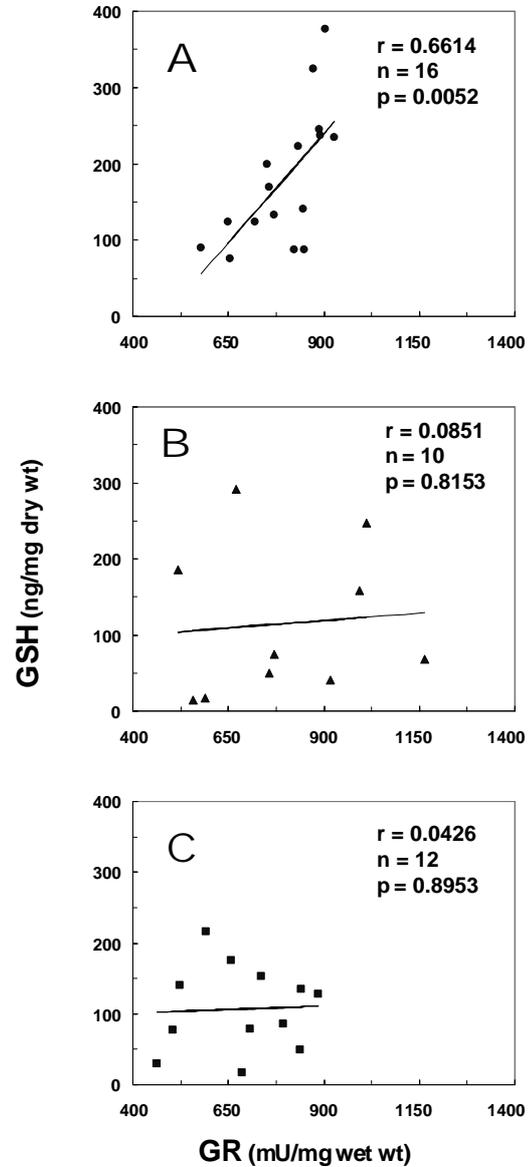


Fig. 4. Correlations of glutathione (GSH) levels to glutathione reductase (GR) activities in postmortem caudate region from control subjects without psychiatric disorders (A), control subjects with bipolar and/or depression (B), and patients with schizophrenia (C).

between female ($n = 6$) and male ($n = 10$) subjects (data not shown).

Similarly, there were no significant differences of any GSH redox measures in control samples between smokers ($n = 10$) and non-smokers ($n = 10$) in the present sample size (data not shown). When non-smokers were excluded from the three test groups, levels of GSH and GR were still found significantly lower in patients with schizophrenia than in control subjects without psy-

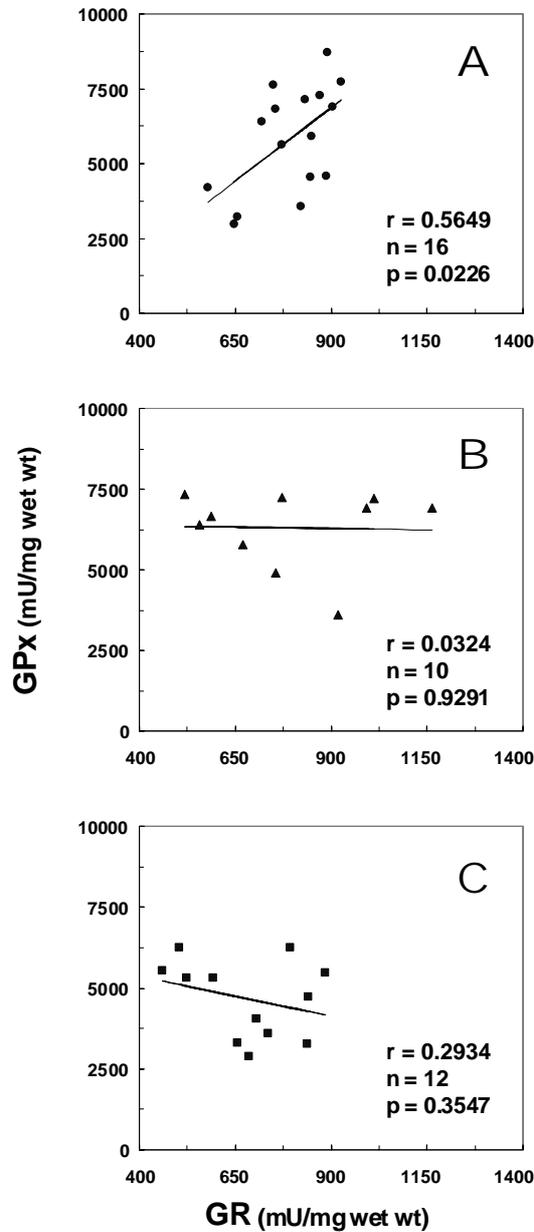


Fig. 5. Correlations of glutathione peroxidase (GPx) to glutathione reductase (GR) activities in postmortem caudate region from control subjects without psychiatric disorders (A), control subjects with bipolar and/or depression (B), and patients with schizophrenia (C).

chiatric disorders (Table 3). Likewise, GPx activities were significantly lower in patients with schizophrenia than in control subjects with bipolar and/or depression. A similar trend was also found between patients with schizophrenia and control subjects without psychiatric disorders, although the difference was not statistically significant. In addition, GSH levels were also signif-

icantly lower in control subjects with bipolar and/or depression than in control subjects without psychiatric disorders (Table 3).

3.4. Effect of age on glutathione reductase reaction

In patients with schizophrenia, both glutathione disulfide (GSSG) and glutathione reductase (GR) were inversely correlated with age ($r = -0.6003$, $p = 0.039$; and $r = -0.6124$, $p = 0.0343$, respectively) in the postmortem caudate region (Figs 6 and 7). Such inverse correlations were not demonstrated in the control groups with and without other psychiatric disorders. On the contrary, GR was positively correlated with age ($r = 0.7140$, $p = 0.0204$) of control subjects with bipolar and/or depressive disorder (Fig. 7).

4. Discussion

4.1. Quantitative determinations of GSH and GSSG

Glutathione plays an important role in metabolism, transport, redox signaling, and cellular protection. Reduced GSH is the major non-protein thiol present in virtually all cells. Its reducing and nucleophilic properties protect cells against destructive effects of reactive oxygen species and free radicals [12]. Therefore, measurements of GSH and its metabolites provide us with useful indices in studying oxidative damage in biological samples. A recent review by Camera and Picardo [13] indicated that GSH levels reported in the literature vary more than 10-fold mainly due to the different methodological approaches such as sample collection, processing and storage, as well as analytical instruments. To minimize GSH autooxidation and rapid enzymatic proteolysis of GSH and GSSG, it is important to store samples at -70°C , and process under refrigeration [13]. Although several chromatographic methods have been developed to measure GSH and GSSG, all have some disadvantages including the need to form derivatives, the inability to determine GSH and GSSG simultaneously, and a lack of detection sensitivity [14]. HPLC with electrochemical detection presents a rapid and sensitive approach to measure GSH and GSSG simultaneously [14,15]. However, detection of disulfides like GSSG by ECD requires application of a higher oxidation potential than GSH. In the present study, we used an HPLC-CMEAS procedure to determine levels of GSH and GSSG. The CMEAS a 16 oxidation potential range from 0 to 1000 mV. The use

Table 3

Comparisons of glutathione redox state in the caudate region of postmortem brain tissue from smokers between schizophrenic and control groups

Mercaptans	Schizophrenics (n = 11)	Controls		P (two-tailed, unpaired t tests)		
		without PD ^a (n = 10)	with other PD (n = 8)	S vs C	S vs C-PD	C vs C-PD
GSH	101 ± 60 ^b	169 ± 95	87 ± 84	0.035	0.350	0.035
GSSG	32 ± 23	35 ± 15	31 ± 29	0.356	0.450	0.339
GPx	4782 ± 1174	5450 ± 1612	6283 ± 1338	0.149	0.012	0.124
GR	688 ± 148	788 ± 104	786 ± 231	0.045	0.159	0.490

^aAbbreviations: PD, psychiatric disorders; S, schizophrenics; C, control; C-PD, control with other types of psychiatric disorder; GSH, glutathione; GSSG, glutathione disulfide; GPx, glutathione peroxidase; GR, glutathione reductase.

^bGSH and GSSG were expressed as ng/mg dry wt., whereas GPx and GR were expressed as mU/mg wet wt.

of several coulometric sensors in series that are maintained at different potentials provides resolution and detection of co-eluting compounds such as GSH and GSSG with different electrochemical features. In addition, a filter removed macromolecules. This is considered a valid method to remove proteins because it does not require addition of acids or organic solvents that might influence sample separation, derivatization, and detection [13].

4.2. Evaluations of GSH redox state in schizophrenia

The present study demonstrated a significant decrease of reduced GSH in postmortem caudate from patients with schizophrenia as compared to control subjects without any psychiatric disorders (Table 2). Such a reduction was not associated with brain collection and sample storage parameters. Concomitantly, there was a significant decrease of GSH:GSSG ratio in patients with schizophrenia (Fig. 2). In cerebrospinal fluid of drug-free patients with schizophrenia, Do et al. [16] have previously demonstrated a significant decrease in the levels of GSH as compared to the controls. These same authors using a non-invasive proton magnetic resonance spectroscopy methodology also showed a similar reduction of GSH level in medial prefrontal cortex of schizophrenic patients. Our current findings are consistent with these reports and support a hypothesis of GSH deficiency in schizophrenia. The GSH deficiency induced in the animal model has been related to mitochondrial damage [17,18] and increased free radical insult [19,20]. Moreover, several pathological conditions including Alzheimer's, Huntington's, and Parkinson's diseases have been associated with a deficiency of GSH and imbalanced ratio of GSH to GSSG [21–23].

Furthermore, the present study demonstrated positive correlations between GSH levels and GPx or GR activities in postmortem caudate from control subjects

without any psychiatric disorders (Figs 3 and 4). Concomitantly, GPx was also significantly correlated with GR activities in control subjects (Fig. 5). These positive correlations suggest that a dynamic state regulates the redox coupling under normal conditions. By contrast, lack of such correlations in schizophrenia point to a disturbance of redox coupling mechanisms in the antioxidant defense system, possibly resulting from a decreased level of GSH (Table 2) as well as age-related decreases of GSSG and GR activities (Figs 6 and 7).

4.3. Antipsychotics and the GSH redox state

It is possible that alterations in the GSH redox state is attributable to the long-term treatment with antipsychotic treatment that these patients likely received. In the present study, all subjects except control subjects without any psychiatric disorders were on antipsychotic medications at the time of death. No significant differences, however, were demonstrated in the caudate mercaptans between control subjects with or without non-schizophrenic psychiatric disorders.

On the other hand, GSH plays a pivotal role in protecting cells from oxidative damage such as induced by dopamine (DA) oxidation. Recently, Grima et al. [24] have proposed that GSH deficit in schizophrenia may be associated with DA-induced oxidative stress. Antipsychotic drugs are known to block DA receptors and consequently, may result in increased levels of GSH. Therefore, the decreased GSH levels observed in this study are unlikely due to antipsychotic effects, because GSH levels may be evenly lower in patients with schizophrenia or other psychoses in the absence of antipsychotic treatment.

4.4. Effect of cigarette smoking on GSH redox state

The prevalence of cigarette smoking in schizophrenic patients is between 70%–90%, compared to that of 35–

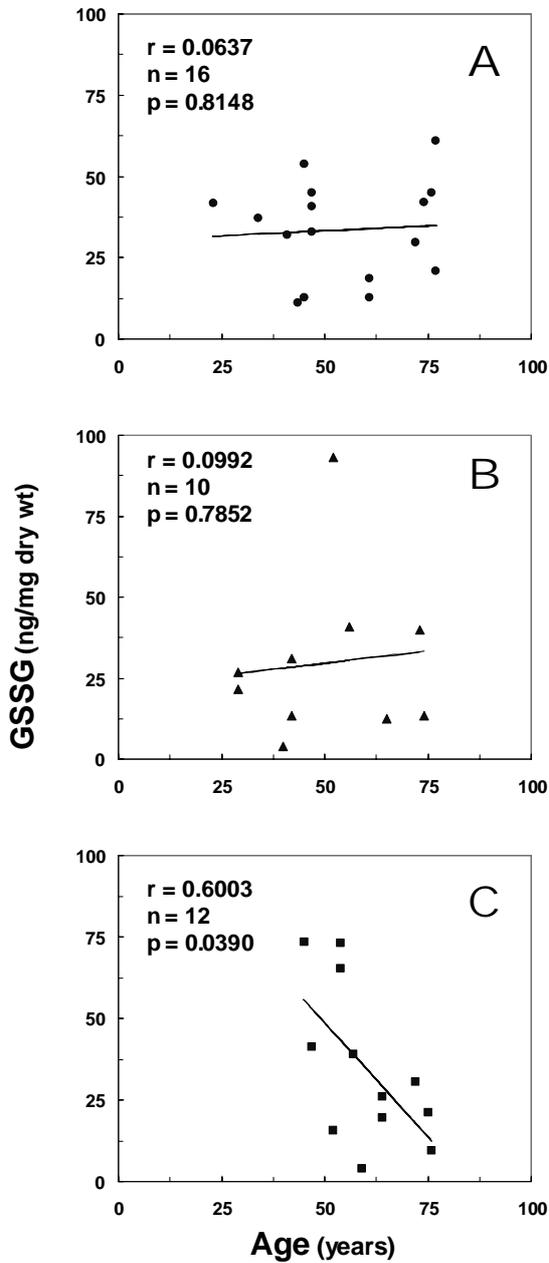


Fig. 6. Correlations of glutathione disulfide (GSSG) levels to age in postmortem caudate region from control subjects without psychiatric disorders (A), control subjects with bipolar and/or depression (B), and patients with schizophrenia (C).

54% for all psychiatric patients and 30–35% for general population [25–27]. Cigarette smoke contains, in addition to nicotine, thousands of substances in the gas and tar phases, many of which are free radicals such as peroxy radicals, oxides of nitrogen, hydroquinones and other radical species [28–31]. Several studies have

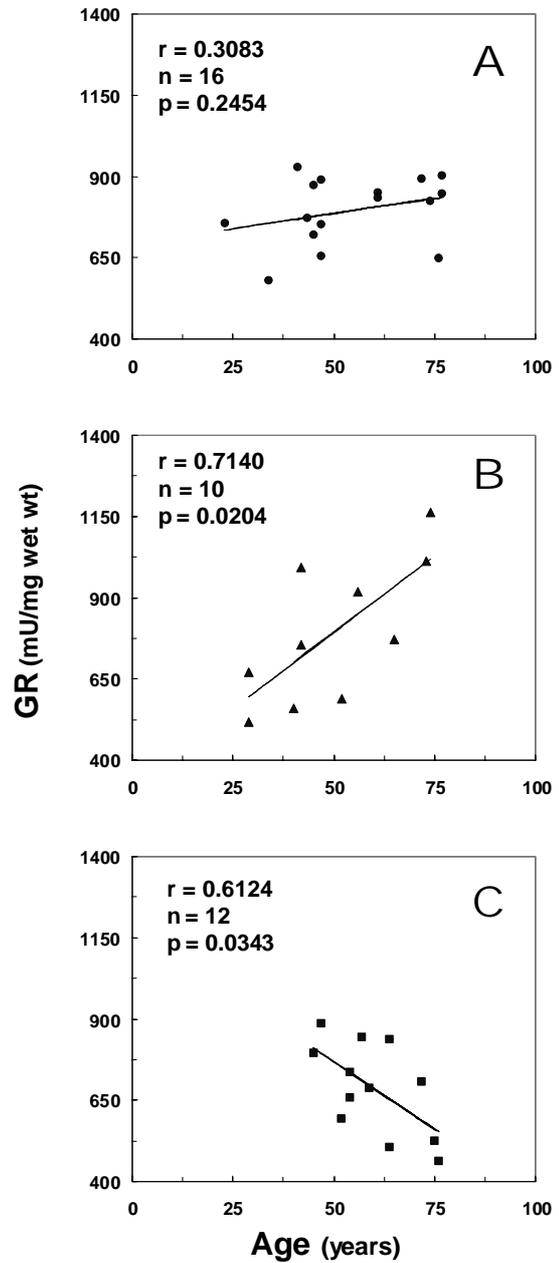


Fig. 7. Correlations of glutathione reductase (GR) activities to age in postmortem caudate region from control subjects without psychiatric disorders (A), control subjects with bipolar and/or depression (B), and patients with schizophrenia (C).

shown that exposure of isolated cells such as epithelial cells and neutrophils or cultured cells to cigarette smoke can lead to rapid depletion of intracellular GSH levels [32–34]. Cigarette smoke induces cell permeability [35,36] and oxidation of lipids and proteins [37, 38] can be prevented by a low-molecular weight an-

tioxidant such as GSH.

As expected, the majority of samples from schizophrenic or bipolar and/or depression groups were obtained from smokers (Table 1). However, the reduced GSH levels found in both patient groups do not appear to be related to cigarette smoking (Table 3).

4.5. GSH redox coupling and reactive nitrogen species

Nitric oxide (NO) is known to have an inhibitory effect on mitochondrial respiration. Following exposure to nitric oxide, GSH will react with NO⁻ to form hydroxylamine and oxidized GSH (GSSG), and with NO⁺ to produce S-nitrosoglutathione [39]. Moreover, peroxynitrite will also react with GSH via either a one or two electron oxidation process, depending on the physiological pH and GSH concentrations [40]. Thus, the intracellular GSH concentration appears to be a critical factor in dictating susceptibility to NO and its derivatives [41]. Under physiologic conditions, NO and its metabolites react with a variety of thiol compounds to form dissociable complexes [42], thereby regulating its inhibitory function [43]. We have recently demonstrated a significantly increased level of NO in the caudate region of postmortem brain from these same individuals with schizophrenia [44], supporting the notion of free radical-mediated neurotoxicity in schizophrenia. Because the reaction of NO with free thiols competes with the same substrate such as glutathione (GSH) for decomposition of hydrogen peroxide by GSH peroxidase, the excessive NO formation may further lead to significant depletion of GSH in schizophrenia.

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