Disruption of non-enzymatic antioxidant defense systems in the brain of rats with water-immersion restraint stress

Yoshiji Ohta,1,* Koji Yashiro,1 Koji Ohashi2 and Yoichiro Imai3

1Department of Chemistry, Fujita Health University School of Medicine, Toyoake, Aichi 470-1192, Japan
2Department of Clinical Biochemistry, Faculty of Medical Technology and 3Department of Biochemistry, Faculty of Clinical Engineering, Fujita Health University School of Health Sciences, Toyoake, Aichi 470-1192, Japan

(Received 31 October, 2011; Accepted 18 January, 2012; Published online 8 June, 2012)

We examined whether non-enzymatic antioxidant defense systems are disrupted in the brain of rats with water-immersion restraint stress. When rats were exposed to water-immersion restraint stress for 1.5, 3 or 6 h, the brain had decreased ascorbic acid and reduced glutathione contents and increased lipid peroxide and nitric oxide metabolites contents at 3 h and showed further changes in these components with a reduction of vitamin E content at 6 h. Increased serum levels of stress markers were found at 1.5, 3 or 6 h of WIRS. Oral pre-administration of L-ascorbic acid (1.5 mmol/kg) or vitamin E (0.5 mmol/kg) to rats with 6 h of water-immersion restraint stress attenuated the increases in lipid peroxide and nitric oxide metabolites contents and the decrease in vitamin E content in the brain. Pre-administered L-ascorbic acid attenuated the decreases in brain ascorbic acid and reduced glutathione contents at 6 h of water-immersion restraint stress, while pre-administered vitamin E enhanced the decreases in those contents. Pre-administered L-ascorbic acid or vitamin E did not affect the increased serum levels of stress markers in rats with 6 h of water-immersion restraint stress. These results indicate that water-immersion restraint stress causes disruption of non-enzymatic antioxidant defense systems through enhanced lipid peroxidation and nitric oxide generation in the brain of rats with water-immersion restraint stress.

Key Words: water-immersion restraint stress, rat brain, non-enzymatic antioxidant defense systems, lipid peroxide, nitric oxide

The brain is especially vulnerable to oxidative damage because the tissue has high oxygen consumption, abundant lipid content, and a relative paucity of antioxidant defense systems to prevent ongoing oxidative damage compared with other tissues. It is known that lipid peroxide (LPO) level in the brain of rats is the highest among main nine tissues, i.e., brain, adrenal, liver, kidney, heart, lung, skeletal muscle, spleen, and testis, while the levels of reduced glutathione (GSH) and vitamin E determined as reduced glutathione (GSH) and vitamin E determined as α-tocopherol (α-Toc) in the brain are relatively low among the nine tissues, although the level of ascorbic acid (AsA) (vitamin C) is relatively high among the nine tissues. Acute immobilization/restraint stress induces oxidative stress in the brain of rats, which might be involved in stress-induced neurobehavioural alterations. Liu et al. have shown that exposure of rats to 8 h of immobilization stress enhances lipid peroxidation in the cerebral, cortex, cerebellum, hippocampus, and midbrain of the brain, although LPO level is simultaneously increased in the serum and liver. Zaidi and Banu have reported that exposure of rats to 6 h of immobilization causes a decrease in GSH level and an increase in LPO level in the brain and that pre- or post-administration of vitamin A, vitamin E or vitamin C attenuates GSH depletion and enhanced lipid peroxidation in the brain of rats with immobilization stress. Pal et al. have reported that exposure of rats to 1 h of restraint stress increases LPO levels in the serum and brain and that pre-administration of L-AsA or α-Toc used as vitamin E for five consecutive days attenuated increased LPO levels in the serum and brain. Gulati et al. have shown in rats with restraint stress that 1 h of stress causes an increase in LPO level and decreases in the levels of GSH and nitric oxide (NO) metabolites (nitrite/nitrate, NOx) in the brain, while 6 h of stress causes an increase in NOx level with similar levels of increased LPO and decreased GSH in the brain. However, it has been shown that exposure of rats to 6 h of cold (2–4°C)-restraint stress has no effect on LPO level in the serum and brain and on GSH level in the brain. Thus, there is a clear difference in the changes in the levels of GSH, a major member of non-enzymatic antioxidant defense systems, and LPO in the brain between rats with acute immobilization/restraint stress and cold-restraint stress.

Water-immersion restraint stress (WIRS) consists of immobilization/restraint stress and a kind of cold stress in which the exposed temperature is usually around 23°C. WIRS is widely used to evaluate stress-induced acute gastric mucosal lesions in experimental animals. We have reported that exposure of rats to WIRS for 3 to 6 h causes oxidative damage associated with disruption of non-enzymatic antioxidant defense systems and enhanced lipid peroxidation and NO• generation in the liver as well as the gastric mucosa. Furthermore, it has been shown that exposure of rats to 6 h of WIRS causes oxidative damage in the liver, kidney, heart, and skeletal muscle and that vitamin E pre-administered to the stressed rats reduces oxidative damage in the liver, kidney, heart, and skeletal muscle. It has also been shown that vitamin E and L-AsA pre-administered to rats with WIRS reduce oxidative damage by attenuating disrupted non-enzymatic antioxidant defense systems and enhanced lipid peroxidation and NO• generation in the gastric mucosa and liver. However, there is no information on whether WIRS disrupts non-enzymatic antioxidant defense systems through enhanced lipid peroxidation and NO• generation in the brain of rats.

In the present study, therefore, we attempted to clarify whether WIRS disrupts non-enzymatic antioxidant defense systems in the brain of rats by examining the time courses of the changes in AsA, GSH, vitamin E, LPO, and NOx levels in the brain of rats exposed to WIRS for a period of 6 h and the effects of pre-administered L-AsA and vitamin E on these changes in the brain of stressed rats. In addition, we examined WIRS-induced changes in the serum

doi: 10.3164/jcbn.11-14
©2012 JCBN
J. Clin. Biochem. Nutr. | September 2012 | vol. 51 | no. 2 | 136–142

To whom correspondence should be addressed.
E-mail: yohta@fujita-hu.ac.jp
levels of adrenocorticotropic hormone (ACTH), glucocorticoid, and glucose, which are stress markers, in rats pre-administered with and without either L-AsA or vitamin E, because it is known that serum levels of these stress markers are increased in rats with WIRS.\textsuperscript{16,17}

### Materials and Methods

**Materials.** RRR-α-Toc used for vitamin E administration was purchased from Sigma-Aldrich (St. Louis, OM). L-AsA, corticosterone, α,α’-dipyridyl, 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB), ethylenediaminetetraacetic acid (EDTA), GSH, 2-thiobarbituric acid, α-Toc and δ-Toc used for vitamin E determination, and other chemicals were obtained from Wako Pure Chemical Ind. Ltd. (Osaka, Japan). All chemicals used were of reagent grade and were not further purified.

**Animals.** Male Wistar rats aged six weeks were purchased from Nippon SLC Co. (Hamamatsu, Japan). The animals were housed in cages in a ventilated animal room with controlled temperature (23 ± 2°C) and relative humidity (55 ± 5%) with 12 h of light (7:00 to 19:00). The animals were maintained with free access to rat chow, Oriental MF (Oriental Yeast Co., Tokyo, Japan) and tap water for one week. All animals received humane care in compliance with the Guidelines of the Management of Laboratory Animals in Fujita Health University. The animal experiment was approved by Institutional Animal Care and Use Committee.

**Induction of WIRS and AsA and vitamin E administrations.** Seven-week-old rats were starved for 24 h prior to experiments, but were allowed free access to water. Rats were restrained in 23 in wire cages and immersed up to the depth of the xiphoid process.

**Materials.** RRR-α-Toc used for vitamin E administration was purchased from Sigma-Aldrich (St. Louis, OM). L-AsA, corticosterone, α,α’-dipyridyl, 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB), ethylenediaminetetraacetic acid (EDTA), GSH, 2-thiobarbituric acid, α-Toc and δ-Toc used for vitamin E determination, and other chemicals were obtained from Wako Pure Chemical Ind. Ltd. (Osaka, Japan). All chemicals used were of reagent grade and were not further purified.

**Animals.** Male Wistar rats aged six weeks were purchased from Nippon SLC Co. (Hamamatsu, Japan). The animals were housed in cages in a ventilated animal room with controlled temperature (23 ± 2°C) and relative humidity (55 ± 5%) with 12 h of light (7:00 to 19:00). The animals were maintained with free access to rat chow, Oriental MF (Oriental Yeast Co., Tokyo, Japan) and tap water for one week. All animals received humane care in compliance with the Guidelines of the Management of Laboratory Animals in Fujita Health University. The animal experiment was approved by Institutional Animal Care and Use Committee.

**Induction of WIRS and AsA and vitamin E administrations.** Seven-week-old rats were starved for 24 h prior to experiments, but were allowed free access to water. Rats were restrained in wire cages and immersed up to the depth of the xiphoid process in at 23°C water bath to induce WIRS, as described in our previous reports.\textsuperscript{11–17} In the time course experiment, rats were subjected to WIRS for 1.5, 3 or 6 h. In the experiment of L-AsA and vitamin E administrations, L-AsA and vitamin E (RRR-α-Toc) were dissolved in 5% Tween 80 and the prepared L-AsA or vitamin E solution was orally administered to rats with and without 6 h of WIRS at a volume of 1 ml/100 g body weight with a stomach tube at 0.5 h before the onset of WIRS. The doses of administered L-AsA and vitamin E were 1.5 and 0.5 mmole/kg body weight, respectively. The doses of L-AsA and vitamin E used were determined based on our previous report.\textsuperscript{15} Stressed and unstressed rats not administered with L-AsA or vitamin E received the same volume of 5% Tween 80 in the same manner.

**Determinations of serum and brain components.** All rats used for the determinations of serum and brain components were sacrificed under ether anesthesia at which time blood was collected from the inferior vena cava. Serum was obtained from the collected blood by centrifugation. Immediately after sacrifice, whole brain was removed from each rat. The collected serum and brains were stored at –80°C and the resultant supernatant was filtrated at 4°C under centrifugation (10,000 g for 20 min at 4°C).

**Materials.** RRR-α-Toc used for vitamin E administration was purchased from Sigma-Aldrich (St. Louis, OM). L-AsA, corticosterone, α,α’-dipyridyl, 5,5’-dithiobis(2-nitrobenzoic acid) (DTNB), ethylenediaminetetraacetic acid (EDTA), GSH, 2-thiobarbituric acid, α-Toc and δ-Toc used for vitamin E determination, and other chemicals were obtained from Wako Pure Chemical Ind. Ltd. (Osaka, Japan). All chemicals used were of reagent grade and were not further purified.

**Animals.** Male Wistar rats aged six weeks were purchased from Nippon SLC Co. (Hamamatsu, Japan). The animals were housed in cages in a ventilated animal room with controlled temperature (23 ± 2°C) and relative humidity (55 ± 5%) with 12 h of light (7:00 to 19:00). The animals were maintained with free access to rat chow, Oriental MF (Oriental Yeast Co., Tokyo, Japan) and tap water for one week. All animals received humane care in compliance with the Guidelines of the Management of Laboratory Animals in Fujita Health University. The animal experiment was approved by Institutional Animal Care and Use Committee.

**Induction of WIRS and AsA and vitamin E administrations.** Seven-week-old rats were starved for 24 h prior to experiments, but were allowed free access to water. Rats were restrained in wire cages and immersed up to the depth of the xiphoid process in at 23°C water bath to induce WIRS, as described in our previous reports.\textsuperscript{11–17} In the time course experiment, rats were subjected to WIRS for 1.5, 3 or 6 h. In the experiment of L-AsA and vitamin E administrations, L-AsA and vitamin E (RRR-α-Toc) were dissolved in 5% Tween 80 and the prepared L-AsA or vitamin E solution was orally administered to rats with and without 6 h of WIRS at a volume of 1 ml/100 g body weight with a stomach tube at 0.5 h before the onset of WIRS. The doses of administered L-AsA and vitamin E were 1.5 and 0.5 mmole/kg body weight, respectively. The doses of L-AsA and vitamin E used were determined based on our previous report.\textsuperscript{15} Stressed and unstressed rats not administered with L-AsA or vitamin E received the same volume of 5% Tween 80 in the same manner.

**Determinations of serum and brain components.** All rats used for the determinations of serum and brain components were sacrificed under ether anesthesia at which time blood was collected from the inferior vena cava. Serum was obtained from the collected blood by centrifugation. Immediately after sacrifice, whole brain was removed from each rat. The collected serum and brains were stored at –80°C until use. Serum LPO was assayed by the fluorometric method of Guillemin\textsuperscript{18} using tetramethoxypropane as a standard. The concentration of serum LPO is expressed as that of malondialdehyde (MDA) equivalents.

Serum ACTH, corticosterone, and glucose concentrations in rats exposed to WIRS for 1.5, 3 or 6 h were significantly higher than those in the corresponding rats without stress and there were no differences in the increased serum ACTH, corticosterone, and glucose concentrations in stressed rats between each time point of WIRS (Fig. 1).

Serum NOx and LPO concentrations in rats exposed to WIRS for 3 or 6 h were significantly higher than those in the corresponding unstressed rats and the increases in serum NOx and LPO concentrations in stressed rats occurred time-dependently (Fig. 2).

Brain NOx and LPO contents in rats exposed to WIRS for 3 or 6 h were significantly higher than those in the corresponding unstressed rats and the increases in brain NOx and LPO contents in stressed rats occurred time-dependently (Fig. 3).

Brain AsA and GSH contents in rats exposed to WIRS for 3 or 6 h were significantly higher than those in the corresponding unstressed rats, although brain AsA content in rats with 1.5 h of WIRS tended to be lower than that in the corresponding unstressed rats (Fig. 4 A and B). The decreases in brain AsA and GSH contents in stressed rats occurred time-dependently, although the extent of decrease in brain AsA content was larger than that of decrease in brain GSH content at 3 or 6 h of WIRS (Fig. 4 A and B). Brain vitamin E content in stressed rats was significantly lower than that in unstressed rats at 6 h of WIRS (Fig. 4C).

When L-AsA (1.5 mmole/kg) or vitamin E (0.5 mmole/kg) was orally administered to rats with 6 h of WIRS at 0.5 h before the onset of stress, the pre-administered L-AsA and vitamin E had no effect on the increases in serum ACTH, corticosterone, and glucose concentrations at 6 h of WIRS (Fig. 5). Administration of the same dose of L-AsA or vitamin E to unstressed rats in the same manner did not affect the serum ACTH, corticosterone, and glucose concentrations at all (Fig. 5).

The pre-administration of L-AsA or vitamin E significantly attenuated increased NOx and LPO concentrations in the serum of rats with 6 h of WIRS, although the attenuating effect of L-AsA was smaller than that of vitamin E (Fig. 6). Administration of the same dose of L-AsA or vitamin E to unstressed rats in the same manner had no effect on the serum NOx and LPO concentrations (Fig. 6).

The pre-administration of L-AsA or vitamin E significantly...
attenuated increased NOx and LPO contents in the brain of rats with WIRS, although the attenuating effect of L-AsA was smaller than that of vitamin E (Fig. 7). The pre-administration of vitamin E returned the increased brain LPO content to near the level of unstressed rats (Fig. 7B). Administration of the same dose of L-AsA or vitamin E to unstressed rats in the same manner had no effect on the brain NOx and LPO contents (Fig. 7).

The pre-administration of L-AsA to rats with 6 h of WIRS significantly attenuated decreased brain AsA, GSH, and vitamin E contents at 6 h of WIRS (Fig. 8). The pre-administration of vitamin E to rats with 6 h of WIRS significantly caused further reductions of decreased brain AsA and GSH contents at 6 h of WIRS, while the vitamin E pre-administration significantly attenuated decreased brain vitamin E content at 6 h of WIRS and returned the decreased brain vitamin E content to near the level of unstressed rats (Fig. 8). Administration of the same dose of L-AsA to unstressed rats in the same manner increased the brain AsA content significantly, while administration of the same dose of
vitamin E to unstressed rats in the same manner increased the brain vitamin E content significantly (Fig. 8).

Discussion

The present study has clearly shown that WIRS causes disruption of non-enzymatic antioxidant defense systems composed of AsA, GSH, and vitamin E through enhanced lipid peroxidation and NO• generation in the brain of rats. Rats exposed to WIRS for 1.5, 3, or 6 h showed a similar extent of stress response at each time point after the onset of WIRS as described in Materials and Methods. Disruption of brain non-enzymatic antioxidant defense system in rats exposed to WIRS for a period of 6 h occurred 3 and 6 h after the onset of stress, i.e., considerably later than the appearance of stress response. Such a timing of the disruption of non-enzymatic antioxidant defense systems in the brain of rats with WIRS was similar to that in the gastric mucosa and liver. 

It is known that exposure of rats to 1 h of immobilization/restraint stress enhances lipid peroxidation in the brain. It is also known that brain NO• generation is lower in rats with restraint stress than in unstressed rats at 1 h after the onset of stress, while that generation is higher in stressed rats than in unstressed rats at 6 h. In the present study, time-dependent increases in NOx and LPO contents in the brain of rats with WIRS occurred 3 and 6 h after the onset of stress, i.e., considerably later than the appearance of stress response. These results suggest that WIRS enhances NO• generation and lipid peroxidation in the brain of rats under lasting stress conditions. In addition, time-dependent increases in serum NOx and LPO concentrations occurred in rats exposed to WIRS and were well consistent with those in brain NOx and LPO contents. It has been suggested that excessive glucocorticoids induce oxidative stress in cultured rat hippocampal slices by downregulating the gene expression of glutathione peroxidase, an enzyme to metabolize hydrogen peroxide and lipid hydroperoxides in the presence of GSH, and by upregulating the gene expression of NADPH oxidase, an enzyme to generate superoxide radical, one of reactive oxygen species (ROS). It is known that
lipid peroxidation is caused via ROS in the brain, which is 
involved in the pathogenesis of neurodegenerative diseases. (26) 
Therefore, excessive corticosterone being secreted continuously 
from the adrenal gland through the hypothalamus-pituitary-
adrenal axis may enhance lipid peroxidation in the brain of rats 
with WIRS by stimulating the generation of ROS in the hippo-
campus. It has been reported that exposure of rats to 6 h of 
immobilization stress activates the expression of inducible nitric 
oxide synthase (iNOS) through the mechanism involved in 
excitatory amino acids and subsequent activation of nuclear factor 
κB in the brain cortex, resulting in excessive iNOS-mediated NO• 
generation in the brain. (27) It has been shown that NO • induces 
oxidative stress through the generation of ROS in neuronal 
cells. (28) Therefore, there seems to be a possibility that exposure 
of rats to WIRS causes excessive NO’ generation through activation 
of iNOS in the brain cortex. 

It has been shown that GSH, a major member of non-enzymatic 
defense systems, is depleted in the brain of rats with immobiliza-
tion/restraint stress. (4–9) When compared with unstressed rats, 
brain AsA and GSH contents were decreased in rats with 3 or 6 h 
of WIRS of which decreases were reversely related to the 
increases in brain NOx and LPO contents. The brain AsA content 
tended to decrease in rats with 1.5 h of WIRS. The extent of 
decrease in AsA content was larger than that in GSH content in 
the brain of rats with WIRS. Furthermore, rats with 6 h of WIRS had 
lower brain vitamin E content than unstressed rats. It is known 
that AsA (reduced form) is oxidized to dehydroascorbic acid 
(oxidized form) to regenerate vitamin E from vitamin E radical in 
the liquid/aqueous interface, resulting in support of the chain-
breaking antioxidant action of vitamin E for lipid peroxidation. (29) 

It is also known that GSH recycles AsA from dehydroascorbic 
acid through a non-enzymatic reaction and/or an enzymatic 
reaction. (30) It has been shown that AsA is oxidized first with 
subsequent vitamin E oxidation for protection of brain against 
oxidative damage. (31) These findings suggest that, in the brain of 
rats with WIRS, AsA is consumed first followed by GSH 
consumption with subsequent vitamin E consumption. Thus, brain 
non-enzymatic antioxidant defense system was disrupted in rats 
with WIRS. These findings also suggest that WIRS could disrupt 
non-enzymatic antioxidant defense systems through enhanced 
lipid peroxidation and NO’ generation in the brain of stressed 
rats with WIRS.

It has been reported that pre- or post-administrations of 
vitamins C and E prevents oxidative stress in the brain of rats 
exposed to immobilization stress for 6 h by attenuating decreased 
GSH level and increased LPO level in the tissue, although pre-
or post-administration of vitamins C and E in combination do 
not cause additive prevention of immobilization-stress-induced 
oxidative stress in the brain. (4) We examined the effects of L-AsA 
and vitamin E pre-administered alone on the changes in AsA, 
glutathione, vitamin E, LPO, and NOx contents in the brain of rats 
with 6 h of WIR in order to confirm the disruption of non-enzymatic 
antioxidant defense systems through enhanced lipid peroxidation 
and NO’ generation in the brain of stressed rats with WIRS.

When L-AsA (1.5 mmole/kg) or vitamin E (0.5 mmole/kg) was orally 
administered to rats with 6 h of WIRS at 0.5 h the onset of stress, both 
pre-administered vitamins did not affect the increases in serum 
ACTH, corticosterone, and glucose concentrations at 6 h of WIRS 
at all. These results indicate that L-AsA and vitamin E pre-
administered to rats with WIRS have no effect on stress response.
Vitamin E functions not only as a chain-breaking antioxidant for lipid peroxidation but also as a scavenger of ROS and NO•. Its known that vitamin E inhibits NO•-induced lipid peroxidation in rat brain homogenates. It is also known that AsA not only supports the chain-breaking antioxidant action of vitamin E for lipid peroxidation but also functions as a scavenger of ROS. The pre-administered L-AsA and vitamin E attenuated the increases in serum LPO and NOx concentrations and brain LPO and NOX contents in rats with 6 h of WIRS. These results support the above-described suggestion that WIRS disrupts non-enzymatic antioxidant defense systems through enhanced lipid peroxidation and NO• generation in the brain of rats with WIRS. The pre-administration of L-AsA attenuated decreased AsA, GSH, and vitamin E contents in the brain of rats with 6 h of WIRS. In addition, the same dose of L-AsA administered to unstressed rats in the same manner increased the brain AsA content. These results indicate that pre-administered L-AsA transported to the brain of rats with 6 h of WIRS restores AsA, GSH, and vitamin E by exerting antioxidant action in a direct manner and/or an indirect manner in the tissue. In contrast, the pre-administration of vitamin E enhanced the decreases in AsA and GSH contents in the brain of rats with 6 h of WIRS, although the vitamin E pre-administration returned the decreased brain vitamin E content to near the level of unstressed rats. The same dose of vitamin E administered to unstressed rats in the same manner increased the brain vitamin E content. As described above, vitamin E pre-administered to rats with 6 h of WIRS attenuated increased LPO and NOx contents in the brain. These results indicate that pre-administered vitamin E transported to the brain of rats with WIRS exerts antioxidant action in the tissue. It is known that not only AsA but also GSH can recycle vitamin E from vitamin E radical. Therefore, the further reductions of decreased brain AsA and GSH contents by vitamin E pre-administration in rats with 6 h of WIRS seems to be explained as follows: reduced AsA and GSH in the brain of rats with 6 h of WIRS are further consumed to maintain pre-administered vitamin E transported to the brain, resulting in prevention of enhanced lipid peroxidation and NO• generation by the transported vitamin E in the tissue. These findings allow us to confirm that WIRS disrupts non-enzymatic antioxidant defense systems through enhanced lipid peroxidation and NO• generation in the brain of rats.

Conflict of Interest

No potential conflicts of interest were disclosed.

Abbreviations

- ACTH: adrenocorticotropic hormone
- AsA: ascorbic acid
- DTNB: 5,5'-dithiobis(2-nitrobenzoic acid)
- EDTA: ethylenediaminetetraacetic acid
- GSH: reduced glutathione
- iNOS: inducible nitric oxide synthase
- LPO: lipid peroxide
- MDA: malondialdehyde
- NO•: nitric oxide
- ROS: reactive oxygen species
- Toc: tocopherol
- WIRS: Water-immersion restraint stress

References

1 Halliwell B. Oxidative stress and neurodegeneration: where are we now? J Neurochem 2006; 97: 1634–1658.
2 Azhar S, Cao L, Reaven E. Alteration of the adrenal antioxidant defense system during aging in rats. J Clin Invest 1995; 96: 1414–1424.
3 Liu J, Wang X, Shigenaga MK, Yeo HC, Mori A, Ames BN. Immobilization stress causes oxidative damage to lipid, protein, and DNA in the brain of rats.
Ohta Y, Chiba S, Tada M, Imai Y, Kitagawa A. Development of oxidative stress in rat brain. *Clin Chim Acta* 2004; 340: 229–233.

Pal R, Gultai K, Chakraborti A, Banerjee B, Ray A. Role of free radicals in stress-induced neurobehavioral changes in rats. *Indian J Exp Biol* 2006; 44: 816–820.

Gulati K, Chakraborti A, Ray A. Differential role of nitric oxide (NO) in acute and chronic stress induced neurobehavioral modulation and oxidative injury in rats. *Pharmacol Biochem Behav* 2009; 92: 272–276.

Chakraborti A, Gulati K, Banerjee BD, Ray A. Possible involvement of free radicals in the differential neurobehavioral responses to stress in male and female rats. *Behav Brain Res* 2007; 179: 321–325.

Gulati K, Chakraborti A, Ray A. Modulation of stress-induced neurobehavioral changes and brain oxidative injury by nitric oxide (NO) mimetics in rats. *Behav Brain Res* 2007; 183: 226–230.

Méndez-Cuesta LA, Márquez-Valadez B, Pérez-De la Cruz V, et al. Early changes in oxidative stress markers in a rat model of acute stress: effect of L-carnitine on the striatum. *Basic Clin Pharmacol Toxicol* 2011; 109: 123–129.

Capel ID, Dorrell HM, Smallwood AE. The influence of cold-restraint stress on some components of the antioxidant defence system in the tissues of rats of various ages. *J Toxicol Environ Health* 1983; 11: 425–436.

Nishida K, Ohta Y, Kobayashi T, Ishiguro I. Involvement of the xanthine-xanthine oxidase system and neurotriphils in the development of acute gastric mucosal lesions in rats with water immersion restraint stress. *Digestion* 1997; 58: 340–351.

Nishida K, Ohta Y, Ishiguro I. Relation of inducible nitric oxide synthase activity to lipid peroxidation and nonprotein sulphydryl oxidation in the development of stress-induced gastric mucosal lesions in rats. *Nitrir Oxide* 1998; 2: 215–223.

Ohta Y, Kamiya Y, Imai Y, Arisawa T, Nakano H. Role of gastric mucosal ascorbic acid in gastric mucosal lesion development in rats with water immersion restraint stress. *Inflammopharmacology* 2005; 13: 249–259.

Ohta Y, Chiba S, Tada M, Imai Y, Kitagawa A. Development of oxidative stress and cell damage in the liver of rats with water-immersion restraint stress. *Redox Rep* 2007; 12: 139–147.

Ohta Y, Imai Y, Kaida S, Kamiya Y, Kawanishi M, Hirata I. Vitamin E protects against stress-induced gastric mucosal lesions in rats more effectively than vitamin C. *Biofactors* 2010; 36: 60–69.

Kaida S, Ohta Y, Imai Y, Kawanishi M. Protective effect of L-ascorbic acid against oxidative damage in the liver of rats with water-immersion restraint stress. *Redox Rep* 2010; 15: 11–19.

Ohta Y, Kaida S, Chiba S, et al. Involvement of oxidative stress in increases in the serum levels of various enzymes and components in rats with water-immersion restraint stress. *J Clin Biochem Nutr* 2009; 45: 347–354.

Yagi K. A simple fluorometric assay for lipoperoxide in blood plasma. *Biochem Med* 1976; 15: 212–216.

Guillemín R, Clayton GW, Lipscomb HS, Smith JD. Fluorometric measurement of rat plasma and adrenal corticosterone concentration; a note on technical details. *J Lab Clin Med* 1959; 53: 830–832.

Zannoni V, Lynch M, Goldstein S, Sato P. A rapid micromethod for the determination of ascorbic acid in plasma and tissues. *Biochem Med* 1974; 11: 41–48.

Sedlak J, Lindsay RH. Estimation of total, protein-bound, and nonprotein sulphydryl groups in tissue with Ellmann’s reagent. *Anal Biochem* 1968; 25: 192–205.

Kamiya Y, Ohta Y, Imai Y, Arisawa T, Nakano H. A critical role of gastric ascorbic acid in the progression of acute gastric mucosal lesions induced by compound 48/80 in rats. *World J Gastroenterol* 2005; 11: 1324–1332.

Ohkawa H, Ohishi N, Yagi K. Assay for lipid peroxides in animal tissues by thiobutiric acid reaction. *Anal Biochem* 1979; 95: 351–358.

Green LC, Wagner DA, Glogowski J, Skipper PL, Wishnok JS, Tannenbaum SR. Analysis of nitrate, nitrite, and [15N]nitrate in biological fluids. *Anal Biochem* 1982; 126: 131–138.

You JM, Yun SJ, Nam KN, Kang C, Won R, Lee EH. Mechanism of glucocorticoid-induced oxidative stress in rat hippocampal slice cultures. *Can J Physiol Pharmacol* 2009; 87: 440–447.

Adibhatla RM, Hatch JF. Lipid oxidation and peroxidation in CNS health and disease: from molecular mechanisms to therapeutic opportunities. *Antioxid Redox Signal* 2010; 12: 125–169.

Méndez-Cuesta LA, Márquez-Valadez B, Pérez-De la Cruz V, et al. Early changes in oxidative stress markers in a rat model of acute stress: effect of L-carnitine on the striatum. *Basic Clin Pharmacol Toxicol* 2011; 109: 123–129.