EXPERIMENTAL STUDY

Protection with thymoquinone against formaldehyde-induced neurotoxicity in the rats

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ABSTRACT

INTRODUCTION: This study aimed to investigate the neurotoxic damage of formaldehyde (FA), which is commonly used in medicine and industrial fields, for the hippocampus of rats and the protective role of thymoquinone (TQ) against this neurotoxicity.

METHODS: There were five groups with eight rats in each. Two control groups were formed, in one of them physiological saline was applied and in the other one corn oil was applied. FA was injected in Group 3. Group 4 was exposed to FA and TQ simultaneously. Group 5 received TQ only. At the end of the experiment animals were sacrificed and brain tissues were removed for biochemical and histopathological investigation.

RESULTS: Catalase (CAT), glutathione peroxidase (GSH-px) and superoxide dismutase (SOD), all known as enzymes with antioxidant activities, were increased in FA and TQ simultaneously administered group. FA caused prominent subarachnoidal hemorrhage and vacuolization. Vacuolization was not observed but occasional subarachnoidal hemorrhage was detected in the FA+TQ group.

CONCLUSION: Neurotoxic damage in hippocampus induced by FA was reverted by administration of TQ (Tab. 1, Fig. 1, Ref. 26).

KEY WORDS: formaldehyde, hippocampus, neurotoxic damage, thymoquinone.

Material and methods

Animals and applications

A total of 40 male rats (Wistar Albino) were used for the current study. Rats were divided into five groups, with eight rats in...
each. Temperature was set to 22 ± 2 °C. The light was arranged in the form of 12 hours for each, daytime and nighttime. The rats from the first control group (Group 1) were injected with physiological saline solution intraperitoneally. Rats in group 2 were administered 1 ml of corn oil via gavage. Rats in group 3 were injected FA via intraperitoneal route with a dose of 10 mg/kg diluted in 10 % physiological saline solution. Rats in group 4 were injected FA at the same dose and duration as in the third group and additionally were administered 20 mg/kg TQ in 1 ml corn oil via gavage. Rats in group 5 were administered 20 mg/kg TQ with 1 ml corn oil via gavage.

Fifteen days later the animals were sacrificed and brain tissues were removed for biochemical and histopathological examinations. One hemisphere of each rat was fixated in 10 % formaldehyde for light microscope examinations. Hippocampus region of the other hemisphere was removed for biochemical examination, placed into glass bottles, labeled, and stored frozen for the eventual determination of SOD, GSH-Px and CAT activities.

**Biochemical analyses**

The tissues were weighed and homogenized in ice-cold Tris-HCl buffer (50 mM, pH 7.4). All procedures were performed at +4 °C. Tissue homogenates were centrifuged to remove debris, and the clear supernatant fluids were separated and kept at −80 °C until the enzyme activity measurements were performed (about a week later).

**Determination of SOD activity**

The method to determine SOD activity for current study is based on recording the optic density (OD) of blue formazan produced while Nito blue tetrazolium (N.B.T) and superoxide radicals generated using xantine and xantine oxidase (XOD) react at 560 nm wavelength. SOD inhibits formazan reaction by disposing superoxide radicals. Inhibition rate depends on SOD activity in the environment. SOD activity is expressed as U/g Hb and U/g protein in blood and tissue, respectively. The process is briefly as follows: erythrocyte hemolysate or tissue homogenate is diluted with phosphate buffer. A buffer solution composed of 0.05 mmol/L sodium and 0.025 mmol/L 2-(4-iodophenyl)-3-(4-nitrophenol)-5-phenyltetrasolium chloride (INT) are mixed. 125μl xantine oxidase (80 U/ml) is added to the mixture and incubated for 5 minutes at 37 °C. Reaction is started by adding 8.8 mmol/L hydrogen peroxide and reductions in absorbance are recorded for 3 minutes (18).

**Determination of CAT activity**

The method is described by Aebi. At 25 °C in erythrocyte hemolysate or tissue homogenate, decomposition rate of the substrate H2O2 is tracked spectrophotometrically at 240 nm for 30 seconds. Activity is expressed as U/g Hb and U/g protein in blood and tissue, respectively (19).

**Histological examination**

For histopathological analysis, brain tissues were fixed in 10 % neutral buffered formaldehyde and embedded in paraffin. 5 μm thick sections obtained from paraffin blocks were stained with hematoxylin-eosin (H–E). Samples were examined and scored by a blind observer using a Nikon Eclipse i5 light microscope with a Nikon DS-Fi1c camera and the Nikon NIS Elements version 4.0 image analysis systems (Nikon Instruments Inc., Tokyo, Japan).

**Statistical analysis**

SPSS 20 package programme was used. ANOVA test was applied and p < 0.05 was considered as significant. All numerical results was expressed as mean ± SD.

**Results**

The weight gain in FA group was much less than the weight gain in control group. The weight gain in treatment group was found to be less than that of the control group and higher than that of the FA group.

**Biochemical results**

The mean CAT, GSH-Px and SOD enzyme activities were decreased in formaldehyde administered rats compared to control group (p < 0.001) (Tab. 1). Additionally, the mean CAT, GSH-Px

| Group              | CAT (U/g) | SOD (U/g) | GSH-Px (U/g) |
|--------------------|-----------|-----------|-------------|
| I – Control        | 41.6±2.9  | 4.2±0.5   | 43.4±1.3    |
| (Physiological saline solution) |           |           |             |
| II – Control (Corn Oil) | 34.4±5.8  | 3.3±0.2   | 48.4±2.0    |
| III – Formaldehyde | 25.5±0.9  | 2.7±0.2   | 26.7±2.9    |
| IV – Formaldehyde + Thymoquinine | 41.2±4.3  | 4.0±0.4   | 55.1±3.0    |
| V – Thymoquinine   | 40.9±4.4  | 3.4±0.3   | 37.1±1.3    |
| p < (I–III)        | <0.001    | <0.001    | <0.001      |
| p < (III–IV)       | <0.001    | <0.001    | <0.001      |

Values are expressed as mean ± standard error. n – number of subjects; CAT – catalase; SOD – superoxide dismutase; GSH-Px – glutathione peroxidase
and SOD enzyme activities were increased in FA and TQ administered rats compared to FA administered rats (p < 0.001) (Tab. 1).

**Light microscopic results**

Histopathological results are shown in Figure 1. Histological features of the brains of the rats from the control and TQ groups were normal in appearances (Fig. 1 A–C). Intracellular vacuolization, intracerebral hemorrhage, and subarachnoidal hemorrhage were detected in the rats from FA group (Fig. 1D–F). In the FA + TQ group vacuolization or hemorrhage were not detected, while partial subarachnoidal hemorrhage was observed (Fig. 1G–I).

**Discussion**

The results obtained from biochemical and histological examination revealed neurotoxic and destructive effects of FA on hippocampus region. According to the results, TQ was able to prevent the mentioned damage.

SOD, CAT and GSH-px are the enzymatic antioxidant systems of the cells. In a trial aimed to determine the toxic effect of FA on hippocampus, FA was detected to cause oxidative stress, to be neurotoxic and to decrease the levels of antioxidant enzymes SOD and GSH-Px (20). In another study, SOD and GSH-px activities in prefrontal tissue samples of rats which were applied FA injection were found to be significantly lower compared to control group. Therefore, FA was pointed out to cause neuronal toxicity (21). Excess doses of FA application for 30 days significantly reduced hippocampal activity and demonstrated diminishment of NMDA receptor activity, besides, hippocampal FA accumulation was reported in the process of diseases which impair memory, like senile dementia (22). Research concerning accumulation of FA in visceral tissues other then brain was also carried out. Methanol was administered to rats, and in the light of the preliminary information that methanol is metabolised in liver and transformed to FA, antioxidant enzyme systems in liver tissues of rats were assessed. Enzymatic antioxidant systems like SOD, CAT and
GSH-Px and nonenzymatic antioxidant systems were found to be impaired (23). We found out that FA triggered reductions in antioxidant enzyme levels of hippocampal tissues, which was similar with previous studies. In this view, we suggest that FA causes neurotoxicity in hippocampus via oxidative damage.

TQ is a potent antioxidant according to many recent studies. In a study performed in 2017, SOD and CAT enzyme activities were reduced after administration of lipopolysaccharides (LPS), followed by an increase after TQ administration. Moreover, LPS were shown to enhance the production of proinflammatory cytokines like IL-1, IL-6 and TNF by activating neuroinflammatory reactions in hippocampus and potentially contributing to neuronal dysfunction, causing impairment of memory and learning. Subsequently, these harmful effects were reverted by administration of TQ (24). In a different trial of brain ischemia, SOD and CAT activities and GSH levels were increased after giving oral TQ. It was supposed that TQ reverted the damage after brain ischemia, possibly by reducing the oxidative stress (25). Likewise, in our study, TQ was shown to increase SOD, CAT and GSH-Px activity and revert the neurotoxicity caused by FA on hippocampus.

Intraventricular injection of FA was shown to impair memory and learning, besides, caused apoptosis and lipid peroxidation in hippocampal neurons in a recent study (26). In one study basophilic dark stained picnotic nuclei of hippocampal neurons have been detected after experimental FA toxicity (20). In another study, apoptotic cells were observed in prefrontal cortices of animals exposed to FA. FA was then reported to be the agent causing damage in these tissues (21). As mentioned above, FA triggers neuronal toxicity throughout the whole brain tissue by several effects. We also found vacuolisation and subarachnoid and intracerebral hemorrhage in brain tissues of rats exposed to FA. When TQ was applied along with FA, no intracerebral hemorrhage and vacuolisation and less subarachnoid hemorrhage was detected. Thus, we may suggest that TQ has a protective effect on brain tissue.

**Conclusion**

The results of current experimental study indicate that FA causes neurotoxicity damage in rats’ hippocampal tissues and TQ reverses these damages. Moreover, the effects of TQ are accompanied with an increase in antioxidant enzyme activities and a decrease in oxidative damage biomarker levels.

The key eligibility of our study is that to the best of our knowledge, it is the first study to research protective effect of TQ against toxic effects of FA on tissues. Also, we demonstrated tissue damage caused by FA and healing provided by TQ both with histologic examination and changes in antioxidant enzyme activities.

The most clear limitation of our study is that FA was administered via intraperitoneal route. Due to the fact that FA has a marked hepatic first pass metabolism, FA activity may have been diminished in our study. Other remarkable deficiency is that we did not examine memory and learning functions of rats, which are useful for evaluation hippocampal functions.

We suggest that further research focused on antioxidant systems and behavioral systems including testing memory and learning functions may reinforce the favourable effects of TQ against tissue damage.

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