Crocin reduced acrylamide-induced neurotoxicity in Wistar rat through inhibition of oxidative stress

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Abstract

Objectives: Acrylamide (ACR) has many applications in different industries. ACR damages the central and the peripheral nervous system in human and animals. Importance of ACR-induced neurotoxicity encouraged researchers to find both different mechanisms involved in ACR neurotoxicity and potent neuroprotective agents. Therefore, this study was designed to investigate the protective effect of crocin, an active constituent of Crocus sativus L. (saffron) on ACR-induced neurotoxicity in Wistar rats.

Materials and Methods: Animals were treated with ACR (50 mg/kg, IP) 11 days for induction neurotoxicity. Crocin (12.5, 25 and 50 mg/kg, IP) were used during treatment with ACR. At the end of treatment, gait score examination was performed. Then, rats were sacrificed and the severity of damage in brain tissue was determined using pathological tests. The level of malondialdehyde (MDA) and glutathione (GSH) content were evaluated in cerebral cortex and cerebellum to determine the role of oxidative stress in this model.

Results: Exposure to ACR induced severe gait abnormalities and pathological changes, but administration of crocin markedly improved behavioral index and histopathological damages. The elevation of lipid peroxidation parallel with reduction of GSH level was observed in cerebral cortex and cerebellum following exposure to ACR. Treatment with crocin markedly decreased MDA level, while elevated GSH content in nervous system as compared to ACR-treated animals.

Conclusion: The administration of crocin markedly improved behavioral and histopathological damages in Wistar rats exposed to ACR. Reduction of oxidative stress can be considered as an important mechanism of neuroprotective effects of crocin against ACR-induced toxicity.

Introduction

Polyacrylamides, high- molecular- weight polymers, are formed by the polymerization of acrylamide (ACR). These polymers are used in different industries such as water and wastewater management, soil coagulation, dye synthesis and in laboratories for gel electrophoresis. In addition to industrial and laboratory uses, ACR forms when carbohydrate- rich foods are cooked in high temperature. Therefore, diet is another main source of environmental ACR exposure in human (1-3). The toxicity of ACR has been studied extensively in both experimental animals and human. It has been shown that exposure to ACR causes genotoxicity, and cellular damage in nervous and reproductive systems in animals. However, only neurotoxicity has been identified by epidemiological studies on human population (4). ACR as a potent neurotoxic agent damages the central and the peripheral nervous system. ACR induces ataxia, skeletal muscle weakness and weight loss in both occupationally exposed human and experimental animal models (3, 4).

An imbalance between the production and removal of free radicals and reactive oxygen species (ROS) increases oxidative stress that is implicated in pathogenesis of some neurodegenerative diseases (5). Furthermore, it has been reported that ACR-induced neurotoxicity in Wistar rats induced lipid peroxidation and reduced antioxidant capacity in nervous system (6, 7). Therefore, oxidative stress plays important role in ACR-induced toxicity (8, 9).
Also, in previous study we showed that ACR could induce ROS production and apoptosis in PC12 cells as an in vitro model of neurotoxicity (10).

Crocin is an active and important compound of Crocus sativus L. (saffron) (11). Crocin is shown to have high antioxidant activity in several in vitro and in vivo models. In both traditional use (12) and modern pharmacological studies, saffron extracts or its active constituents crocin and safranal are shown to have hepatoprotective effects (13), radical scavenger properties (14), antidepressant (15-17), memory-improved (17, 18), anticonvulsant (19, 20), antinociceptive (21), antineuropathic (22, 23), hypotensive (24), anxiolytic and hypnotic (25, 26) and cardioprotective effects (27).

Crocin can reduce oxidative damage caused by ischemia-reperfusion injury in kidney (28) and skeletal muscle (29). This carotenoid increases cell viability in serum deprivation PC12 cells by inducing glutathione (GSH) synthesis, increasing the glutathione reductase (GR) and c-glutamylcysteinyl synthase (c-GCS) activities, and decreasing ceramide formation (30, 31). Further study indicated that crocin is more potent than α-tocopherol, against oxidative stress-induced death of neurons (30, 31). Crocin also reduced the haematological toxicity of diazinon and the augmentation of direct 8-iso-prostaglandin F(2 beta), haematological toxicity of diazinon and the death of neurons (30, 31). Crocin also reduced the activities, and decreasing ceramide formation (30, 31).

Therefore, in the current study we decided to find the protective effect of crocin against ACR-induced neurotoxicity in Wistar rats with focus on oxidative stress evaluation in cerebral cortex and cerebellum.

Materials and Methods

Materials

ACR was purchased from Merck. Malondialdehyde tetraethylammonium, reduced glutathione (GSH) and DTNB [5, 5'-dithiobis-(2-nitrobenzoic acid)] were obtained from Sigma. Vitamin E (Ampule 100 IU/ml, Osveh) was used for this study.

Crocin preparation

Stigmas of Crocus sativus L. were purchased from Novin Saffron (collected from Ghaen, Khorasan province, Northeast of Iran) and analyzed in accordance with the ISO/TS 3632-2. Crocin was extracted and purified according to the previously described method (34). The purity of crocin was 98%.

Experimental animals

Male Wistar rats, 230 to 250 g were housed in colony rooms with 12/12 hr light/dark cycle at 21±2 °C and had free access to food and water. All animal experiments were carried out in accordance with Mashhad University of Medical Sciences, Ethical committee Acts.

Experimental design

To induce neurotoxicity in Wistar rats, animals were exposed to ACR (50 mg/kg/day IP) for 11 days (3, 35, 36). Daily dose-rate and corresponding route of administration have been well characterized with respect to pathological expression and neurological deficits. For our study, rats were randomly divided into 7 groups (n=6 in each group) and the treatment was given as follow:

1) Control, Normal saline
2) ACR, 50 mg/kg
3) ACR, 50 mg/kg + crocin 12.5 mg/kg
4) ACR, 50 mg/kg + crocin 25 mg/kg
5) ACR, 50 mg/kg + crocin 50 mg/kg
6) ACR, 50 mg/kg + vitamin E 200 mg/kg
7) crocin 50 mg/kg

ACR was administrated IP once a day for 11 days. Saline was given in the same way to control rats. Crocin and vitamin E administrated IP for 11 days once a day and three times a week, respectively.

The behavioral index (gait scores) examination

At the end of animal treatment, the gait scores were examined according to the methods described by LoPachin (3). Rats were placed in a clear plexiglass box and were observed for 3 min, and a gait score, from 1 to 4, was assigned; where 1 = a normal, unaffected gait; 2 = a slightly affected gait (foot splay, slight hindlimb weakness and spread); 3 = a moderately affected gait (foot splay, moderate hindlimb weakness, moderate limb spread during ambulation); and 4 = a severely affected gait (foot splay, severe hindlimb weakness, dragging hindlimbs, inability to rear).

Histopathology assay

Tissue preparation

After gait score test, animals were sacrificed and the brain of each animal was removed completely and immediately fixed in 10% buffered formalin solution overnight. Then each brain was cut coronally and sliced. Each slice measuring about 2 to 3 mm processed by an automatic tissue processor (Shandon, England) in about 18 hr through the routine histopathological method, and further fixed in formalin, dehydrated in alcohol, cleared in xylene and finally paraffinized. Then paraffin blocks were prepared and sectioned at 3 to 5 µm thickness and the microscopic slides were stained by routine Hematoxyline and Eosin method.

Histopathologic examination

Slides were evaluated by the pathologist using the light microscope (Olympus BX-40, Japan). Necrotic neurons were considered in all areas of
the brain, and the amount of necrosis was estimated in a semiquantitative manner in the whole brain for each animal.

Tissue sampling
At the end of 11 days exposure, after behavioral index examination, rats were sacrificed and the brain tissues (cerebral cortex and cerebellum) were dissected, and then samples were snap-frozen in liquid nitrogen and stored at −80°C until use.

Lipid peroxidation assay
Malondialdehyde (MDA) levels, as marker of lipid peroxidation, were measured in brain tissues. MDA reacts with thiobarbituric acid (TBA) as a thiobarbituric acid reactive substance (TBARS) to produce a pink colored complex, which has maximum absorbance at 532 nm.

Briefly, 3 ml phosphoric acid (1%) and 1 ml TBA (0.6%) were added to 0.5 ml of brain tissue homogenate 10% in KCl and the mixture was heated for 45 min in a boiling water bath. After cooling, 4 ml of n-butanol was added to the mixture and vortex-mixed for 1 min followed by centrifugation at 3000 g for 10 min. The organic layers were transferred to a fresh tube and absorbance was recorded at 532 nm (38). A calibration curve was plotted using malondialdehyde tetrabutylammonium. MDA levels were expressed as nmol/mg protein. The total protein contents of the supernatants were determined by the Bradford assay using bovine serum albumin as standard.

Reduced glutathione (GSH) assay
Brain GSH content was determined by the method of Moron et al (1979) with minor changes. Briefly, 300 μl of tissue homogenates were mixed with 300 μl of 10% tricolar acetic acid (TCA) and vortexed. After centrifugation at 2500 g for 10 min, supernatants were mixed with reaction mixtures containing 2 ml phosphate buffer (pH: 8) and 500 μl DTNB [5,5′ di thio bis-(2-nitrobenzoic acid)]. Within 10 min, the absorbance was measured at 412 nm using a spectrophotometer (Jenway 6105 uv/vis, UK) (39). GSH contents were determined from a standard curve produced using commercially available GSH. Levels of GSH were expressed as nmol/mg protein.

Statistical analysis
Results are expressed as mean±SD. Statistical analyses were performed with ANOVA followed by Tukey–Kramer test to compare the differences between the means. Differences were considered statistically significant when P<0.05. Statistical analysis for the gait abnormalities were performed with nonparametric test Kruskal-Wallis followed by Dunn’s multiple comparison test.

Results
Effect of ACR on the behavioral index (gait scores) in rats and protective effect of crocin
Exposure to ACR (50 mg/kg, IP) for 11 days led to progressive gait abnormalities in rats as shown in Figure 1. In animals that treated with combination of crocin (25 mg/kg and 50 mg/kg) and ACR gait abnormalities significantly reversed in the dose-dependent manner (P<0.05 and P<0.01 vs ACR treated group).

Histopathological examination of brain tissues
Evaluation of brain tissues in control group exhibited normal cells (Figure 2A), but severe neuronal necrosis was observed in rats that were exposed to ACR. Histopathological evaluation of brain tissues exhibited the dark neurons in cerebral cortex, cerebellum, hippocampus and basal ganglia (Figure 2B, D, E and F). Treatment with crocin diminished the severity of necrosis to the extent that in some treated brains, no considerable necrosis was detectable (Figure 2C, Table 1).

Effect of crocin on lipid peroxidation induced by ACR
Lipid peroxidation in the cerebral cortex and cerebellum was determined by measuring MDA content. Rats that were exposed to ACR showed a significant (P<0.001) increase in the level of MDA from 1.00 ± 0.31 to 4.04 ± 0.60 nmol/mg protein in cerebral cortex and from 1.26 ± 0.32 to 3.00 ± 0.41 nmol/mg protein in cerebellum tissue when compared to control rats (Figure 3A, B). Treatment with crocin at doses of 12.5, 25 and 50 mg/kg significantly decreased the level of MDA in cerebral cortex (P<0.001). In cerebellum tissue, crocin 12.5 mg/kg had no effect on MDA level, but crocin 25 and 50 mg/kg significantly (P<0.001) inhibited lipid peroxidation.
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Figure 2. Histopathological alterations (Hematoxylin and Eosin staining by light microscope) of brain tissues in control and experimental groups of rats. Sections were visualized under light microscope at the magnification of × 200 for A, and × 100 for B, C, D, E and F. A, control rats showing normal histology in cerebral cortex. B, D, E and F, ACR-exposed rats showing the dark neurons in cerebral cortex, cerebellum, basal ganglia and hippocampus, respectively. C, ACR + Crocin 50 mg/kg group showing less numbers of dark neurons in cerebral cortex.

ACR: Acrylamide

Table 1. Effect of crocin on the histopathological changes in brain tissues following exposure to ACR. ***P<0.001 vs. Control, ###P<0.001 vs. ACR treated animals

| Group                                   | % Necrosis |
|-----------------------------------------|------------|
| Control                                 | 0          |
| ACR (50 mg/kg)                          | ###100     |
| ACR + Crocin (12.5 mg/kg)               | ***70      |
| ACR + Crocin (25 mg/kg)                 | ***60      |
| ACR + Crocin (50 mg/kg)                 | ***30      |
| ACR + Vit E (200 IU/kg)                 | ***30      |
| Crocin (50 mg/kg)                       | 0          |

ACR: Acrylamide

Effect of crocin on GSH content in brain tissues following exposure to ACR

Exposure to ACR exhibited a significant decrease in GSH content in cerebral cortex and cerebellum (P<0.001) (Figure 4A, B). Crocin dose-dependently and significantly increased the levels of GSH in brain tissues compared to ACR treated rats. But, 12.5 mg/kg crocin could not recover GSH content in cerebral cortex and cerebellum. Treatment with 50 mg/kg crocin alone had no effect on the GSH content (P>0.05).

Discussion

In the previous study, we reported ACR induced cytotoxicity in PC12 cells, increased ROS generation and apoptotic cell death, but crocin protected cells from ACR-induced cytotoxicity by interrupting the cell death cascade through inhibition of intracellular ROS production (10). Based on these results from in vitro model, current study was designed to evaluate neuroprotective effect of crocin in Wistar rat.

Results showed that exposure to ACR markedly declined GSH contents while enhanced MDA levels

Figure 3. Effect of crocin on lipid peroxidation in cerebral cortex (A) and cerebellum (B) following treatment with ACR. Animals were treated with different doses of crocin (12.5, 25 and 50 mg/kg) during treatment with ACR 50 mg/kg for 11 days. Data are expressed as the mean±SD, (n=6). ###P<0.001 vs. Control, ***P<0.001 vs. ACR treated animals.

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Figure 4. Effect of crocin on GSH content in cerebral cortex (A) and cerebellum (B) following treatment with ACR. Animals were treated with different doses of crocin (12.5, 25 and 50 mg/kg) during treatment with ACR 50 mg/kg for 11 days. Data are expressed as the mean ± SD, (n = 6). *** P<0.001 vs. Control, ** P<0.01, *** P<0.001 vs. ACR treated animals

ACR: Acrylamide

in cerebral cortex and cerebellum. Crocin administration dose-dependently reduced lipid peroxidation through enhancement of GSH content in both cerebral cortex and cerebellum. Also, crocin recovered histopathological and behavioural damage, which was induced with ACR.

GSH plays an important role in maintaining cell integrity against exogenous and endogenous derived ROS (40). Previous studies indicated that ACR is detoxified by conjugation with GSH (41, 42). Gycidamime, as a metabolite of ACR, could also conjugate with GSH (3, 43). Our study clearly exhibited that exposure to ACR markedly decreased GSH content in cerebral cortex and cerebellum. Reduction of GSH may make the brain tissues more sensitive to the oxidative stress, because following ACR treatment, significant enhancement of MDA level was observed in the cerebral cortex and cerebellum, suggesting the augmentation of the lipid peroxidation. Because of crucial role of oxidative stress in ACR-induced cytotoxicity (6, 8), it is suggested that antioxidants could be considered as an alternative approach in modulation of ACR toxicity (9, 44, 45).

Numerous researches indicated the antioxidant effect of crocin in different conditions. Crocin attenuated lipid peroxidation in kidney (28) and skeletal muscle (29) during ischemia-reperfusion-induced oxidative damage in rats. Moreover, this carotenoid protected the brain against reperfusion-induced oxidative/nitrative injury after global cerebral ischemia in mice (46).

Crocin administration (12.5, 25 and 50 mg/kg) does-dependently increased GSH content in both cerebral cortex and cerebellum, markedly reduced MDA level as a marker of lipid peroxidation, and markedly declined gait abnormalities in experimental groups treated with crocin. Depletion of neural GSH level and enhancement of lipid peroxidation might be one of the primary events in ACR-induced neurotoxicity; therefore, potent antioxidants such as crocin can demonstrate neuroprotective effect in ACR-induced neurotoxicity through interrupting oxidative stress.

Vitamin E has been considered as a neuroprotective agent in many researches, and its protective effect has been widely believed to be due to its antioxidant activity (47). Recently, anti-apoptotic effect of vitamin E has been reported (48). Thus, in current study vitamin E was used as a positive control in protection against ACR-induced neurotoxicity. Results clearly exhibited that there is no difference between vitamin E and crocin (50 mg/kg) effect in inhibition of lipid peroxidation; therefore, crocin may be mentioned as a potent neuroprotective agent.

Conclusion

ACR induced gait abnormalities and histopathological damages via reduction of GSH contents and induction of lipid peroxidation in cerebral cortex and cerebellum. Crocin administration dose-dependently decreased ACR toxicity through suppression of lipid peroxidation and the elevation of GSH content.

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