7,8-Dihydroxyflavone Protects PC12 Cells against MPP+-Induced Cytotoxicity by Heme Oxygenase-1

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Abstract: 7, 8-dihydroxyflavone (7, 8-DHF), a newly-selective tyrosine kinase receptor B (TrkB) agonist, has been well studied for its neurotrophic functions. Our previous study has shown that 7,8-DHF could exert potent neuroprotective effect against 6-hydroxydopamine (6-OHDA)-induced cytotoxicity in PC12 cells without TrkB receptor expression. However, this phenomenon induced by 1-methyl-4-phenylpyridinium(MPP+) has not been elucidated. This study is designed to investigate whether heme oxygenase-1(HO-1)signaling pathway mediate the protection by 7,8-DHF. The results were as follows: MTT assay showed that treatment with MPP+ significantly suppressed the cell viability. 7,8-DHF pretreatment suppressed MPP+-induced cytotoxicity and providing strong protection in PC12 cells. Incubation of 7,8-DHF with the PC12 cells up to 6 h showed it up-regulated the HO-1 expression. Moreover, the PC12 cells pretreatment with ZnPP were found to partially block the protective effect of 7,8-DHF. The present results provided the evidence that 7,8-DHF could protect the PC12 cells against MPP+-induced cytotoxicity through HO-1 pathway.

Keywords: 7,8-Dihydroxyflavone, 1-Methyl-4-Phenylpyridinium, PC12 Cells, Heme Oxygenase-1, ZnPP

Introduction
Parkinson’s disease (PD), a progressive neurodegenerative disease, is characterized by the selective loss of dopaminergic neurons in the substantia nigra[1, 2]. PD is characterized by tremor, bradykinesia, rigidity and postural instability [3, 4]. Oxidative stress is considered the most important causal factors of neuronal death in PD [5, 6]. Although its etiology remains unknown, nuclear factor erythroid 2-related factor 2 (Nrf2)/HO-1 pathway has been implicated in the cellular protection against oxidative stress [7–9]. Under basal conditions, Nrf2 is sequestered in the cytoplasm by the inhibitory Kelch-like ECH-associated protein-1 (Keap1). Upon stimulation, Nrf2 dissociates from Keap1 and translocates to the nucleus, where it binds to the antioxidant-response element (ARE), leading to upregulation of several antioxidant enzymes such as HO-1 and various antioxidants.

Reports have found that an impaired antioxidant system has been observed in PD, including decreased heme oxygenase-1 (HO-1) expression [10, 11]. HO-1 is an inducible enzyme that has been shown to participate as an essential defensive mechanism for neurons exposed to oxidant challenges, being related to antioxidant defenses in certain neuropathological conditions [12].

7,8-Dihydroxyflavone (7,8-DHF), a member of flavonoid family, has been identified as a selective tyrosine kinase receptor B (TrkB) agonist and exerts potent neuroprotective and neurotrophic effects in Parkinson’s disease (PD), and other neurological disorders [13, 14]. Except activating TrkB receptors, antioxidant property of 7,8-DHF also contributes to its protection against oxidative stress-induced damage [15–17].

Our previous studies have demonstrated that 7,8-DHF protects PC12 cells against 6-OHDA-induced cytotoxicity through its antioxidant action and activating PI3K/Akt [18]. Recently, several studies reported that cytoprotective effects of 7,8-DHF against oxidative stress are associated with the Nrf2-induced expression of HO-1 in lung fibroblasts, keratinocytes and C2C12 myoblasts [19, 20]. In the present study, we further elucidated whether HO-1 pathway mediate the protective effect of 7,8-DHF against MPP+-induced PC12 cell injury.

2. Materials and methods

2.1 Materials 7,8-DHF was purchased from TCI America (USA). Thiazolyl blue tetrazolium bromide (MTT), MPP+ and Zinc protoporphyrin IX (ZnPP, a specific inhibitor of HO-1) were Sigma products (St.
Louis, MO, USA). Dulbecco’s modified Eagle’s medium (DMEM) and fetal bovine serum (FBS) were bought from Hyclone (Logan, USA). Rabbit polyclonal anti-HO-1 was from Santa Cruz. Rabbit anti-β-actin was from Bioss Beijing. BCA protein assay kit was from Thermo Fisher Scientific (USA).

### 2.2 Cell culture

The PC12 cells, a Rat pheochromocytoma cell, were supplied by the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The cells were maintained in DMEM supplemented with 10% FBS, 100 IU/ml of penicillin and 100 μg/ml of streptomycin at 37°C in a humidified atmosphere of 5% CO_{2} and 95% air. It was necessary to change the medium every two or three days. The cells were maintained in 96-well plates (MTT assay) or 6-well plates (Western blotting assay) for indicated times.

### 2.3 Determination of cell viability

The PC12 cells were cultured in 96-well plates at a density of 10^4/well. Following 24 h of incubation, the PC12 cells were treated with MPP^+ (100, 200, 300, 400, 500 μmol/L) for 24 h. And the cells were pretreated with the 7, 8-DHF (1, 5, 25 μmol/L) for 6 h, then exposed to MPP^+ (400 μmol/L) for subsequent 24 h. In some experiments, ZnPP (5 μmol/L) was added in 7,8-DHF treatment, then send to MPP^+ (400 μmol/L). Cell viability was measured by the MTT assay. Briefly, 20 μl of MTT solution (5 mg/ml in PBS) was added into each well. Then, the plates were incubated for 4 h at 37°C. The dark blue formazan product, the reduction of MTT, was dissolved in 150 μl of DMSO. Signal detection was implemented by microplate reader (Molecular Devices, Sunnyvale, CA, USA) at the wavelength of 570 nm. The cell viability was defined as the percentage of the untreated control cells.

### 2.4 Western blot analysis

Total protein was extracted from the PC12 cells. After BCA protein assay, each sample was separated by 10% sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) and then transferred to a polyvinylidene difluoride (PVDF) membrane. The membranes were incubated with 10% nonfat milk to block the non-specific binding sites for 1 h. The membranes were probed with different primary antibodies overnight at 4°C, followed by incubation with goat anti-rabbit HRP-conjugated secondary antibody for 1 h at room temperature. Finally, the protein bands were detected by the ECL System (Bio-Rad, USA) and quantified by densitometric analysis using Image J software.

### 2.5 Statistical analysis

The results were obtained from four to six independent experiments, and presented as means ± SEM. One-way analysis of variance (ANOVA) followed by Turkey’s multiple comparison test was performed to compare the differences between means. Differences were considered statistically significant when P values were less than 0.05.

### 3. Results

#### 3.1 Effect of different dosage of MPP^+ on PC12 cells viability

We first found the neurotoxic effect of different dosage of MPP^+ on neuronal differentiated PC12 cells. Our results showed that a decrease in cell viability was induced by MPP^+ in a concentration-dependent manner. Cell viability was 60% of control cultures after exposure to 400 μmol/L MPP^+ for 24 h (Fig.1). Based on these results, 400 μmol/L MPP^+ was used in the following experiments to assess the neuroprotective effect of 7,8-DHF.

![Fig.1 The PC12 cells were exposed to different dosage of MPP^+ (100, 200, 300, 400, and 500 μmol/L) for 24 h. Cell viability was determined by the MTT method. Data are expressed as percentage of values in untreated control cultures, and are the means±SD of six separate experiments. “*”P < 0.01 compared with the untreated control, “**”P < 0.001 compared with the untreated control.](http://www.iijSciences.com)
dramatically improved the viability up to 87.72% in a dose-dependent manner (Fig. 2). This results suggest that 7,8-DHF pretreatment for 6 h might activate intracellular defense system, leading to more powerful protection against MPP⁺-induced cytotoxicity and the most effective concentrations was 25μmol/L.

Fig. 2 The PC12 cells were pretreated with various concentration of 7,8-DHF(1,5,25μmol/L) for 6 h, and then incubated with or without 400μmol/L MPP⁺ for subsequent 24 h. Cell viability was determined using the MTT assay. The results were obtained from five independent experiments. ***P < 0.01, ****P < 0.0001 compared with the untreated control, ""P < 0.05 compared with MPP⁺ group.

3.3 Neuroprotective properties of different dosage of 7,8-DHF against MPP⁺-induced expression of HO-1

To clarify the involvement of HO-1 signaling pathway in the protective effect of 7,8-DHF, expression of HO-1 was analyzed by Western blot. The results revealed that pretreatment with 7,8-DHF for 6 h enhanced the HO-1 level induced by MPP⁺. We also found that the upregulation of HO-1 expression by 7,8-DHF was also dose-dependent (Fig 3). This results suggest that 7,8-DHF pretreatment for 6 h might activate HO-1 signaling pathway, leading to more powerful protection against MPP⁺-induced cytotoxicity.

Fig. 3 The PC12 cells were treated with 7,8-DHF(1,5,25μmol/L) for 6 h and then exposed in 400μmol/L MPP⁺ for subsequent 24 h. The HO-1 protein levels were estimated by Western blot. The results were expressed by the ratio of target proteins to β-actin and obtained from six independent experiments.. *P< 0.05 compared with the control.

3.4 Blocking effect of HO-1 inhibitor on protection by 7,8-DHF

To confirm that the protective effects of 7,8-DHF are mediated through the activation of HO-1 pathway, the PC12 cells were pretreated with ZnPP, a specific HO-1 inhibitor, and 7,8-DHF before the incubation of MPP⁺, the cell viability was then assessed. 7,8-DHF abrogated the cell injury induced by MPP⁺, and this protective effect were partially blocked by ZnPP (Fig 4). These results suggest that HO-1 might be involved in the protective effect of 7,8-DHF.
with MPP+ (400µM) for 24 h. The cell viability was estimated by the MTT assay. The results are obtained from six independent experiments. ***P < 0.0001 compared with the control, **P < 0.001 compared with MPP+ group, *P < 0.05 compared with MPP group+7,8-DHF.

4. Discussion

There is growing evidence that 7,8-DHF exerts neuroprotection by acting as a potent antioxidant [15-17]. 7,8-DHF is considered as a direct free radical scavenger due to two adjacent hydroxyl groups as electron donors [21, 22]. Our previous study also showed that the protective effect of 7,8-DHF in 6-OHDA-treated PC12 cells are associated with increased HO-1 activation [18]. In conclusion, the protective effect of 7,8-DHF was closely associated with its antioxidant property. Recently, 7,8-DHF has been reported to activate Akt and ERK-dependent Nrf2 signaling cascades in cultured human HaCaT keratinocytes and lung fibroblasts [23], leading to the up-regulation of HO-1 and cytoprotection against oxidative stress. Therefore, it is likely that Nrf2/HO-1 pathway might be involved in the protective effect of 7,8-DHF in MPP+-induced PC12 cells injury.

1-Methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP) induces toxicity through its conversion of astrocytes to MPP+ in the reaction catalyzed by monoamine oxidase type B (MAO-B) [24]. MPP+ is selectively transported into the dopaminergic neurons through a high-affinity dopamine (DA) transporter and accumulates in the neuronal mitochondria increases sensitivity to oxidative attack, and eventually causes apoptotic or necrotic neuronal cell death. In this study, we observed that exposure to MPP+ markedly reduced the PC12 cells viability and increased the HO-1 levels [25, 26]. However, these effects were suppressed by pretreatment with 7,8-DHF.

Elevated HO-1 expression levels are involved in antioxidant defense in Parkinson disease patients [27]. Accumulating evidence has demonstrated that activation of HO-1 pathway plays a central role in cellular antioxidant defense [7, 28, 29]. Under normal conditions, Nrf2 was bonded in the plasma with Keap1, which facilitate the ubiquitination and subsequent proteolysis of Nrf2. Once exposure to various types of stress, the Nrf2/Keap1 complex is dissociated, leading to the phosphorylation of Nrf2, a critical process in the nuclear translocation of Nrf2 and activation of target genes including HO-1. In our study, although 7,8-DHF did not increase the whole cell levels of Nrf2, it facilitated the translocation of Nrf2 to the nuclear, increasing the HO-1 expression. These results show treatment with 7,8-DHF activate Nrf2/HO-1 in PC12 cells.

Using the MTT assay, it was found that pretreatment with ZnPP, a HO-1 inhibitor, markedly blocked the protection by 7,8-DHF after exposure to MPP+. These results suggest that Nrf2-dependent induction of HO-1 may, at least in part, participate the protection by 7,8-DHF in MPP+-injured PC12 cells. Qinyong Ye et al reported that astaxanthin protects against MPP+-induced oxidative stress by the HO-1/NOX2 axis in the PC12 cells [30]. Jiang et al reported that gosdronin protects against MPP+-induced oxidative stress by upregulation of HO-1 through p38 MAPK/Nrf2 pathway in human dopaminergic cells [31]. Lee et al found that sulfuretin that the cytoprotective effect of sulfuretin against tert-butyl hydroperoxide-induced hepatotoxicity through Nrf2/ARE and JNK/ERK MAPK-mediated HO-1 expression [32]. These findings suggest that the protein kinase-mediated activation of Nrf2/HO-1 pathway may be specific to the stimulus and cell type.

As expected, 7,8-DHF alone markedly elevated the HO-1 levels, and pretreatment with ZnPP attenuated 7,8-DHF-induced HO-1 expression. These results suggest that neuroprotective properties was mediated by the HO-1 signaling pathways, and the HO-1-induced upregulation are involved in the protection by 7,8-DHF against MPP+-induced cell injury in PC12 cells.

In conclusion, the present study demonstrated that 7,8-DHF activated the intracellular defense systems through HO-1, and improves the survival of PC12 cells treated with MPP+. 7,8-DHF, acting as a small-molecule TrkB agonist and potent antioxidant, might be promising candidate for the therapy of neurological diseases.

References


