Catalpol Exerts a Neuroprotective Effect in the MPTP Mouse Model of Parkinson’s Disease

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The degeneration of dopaminergic (DA) neurons in Parkinson’s disease (PD) is related to inflammation and oxidative stress. Anti-inflammatory agents could reduce the risk or slow the progression of PD. Catalpol, an iridoid glycoside extracted from the roots of Rehmannia radix, has been reported to reduce the release of inflammatory factors and exert neuroprotective effects. 1-methyl-4-phenyl-1,2,3, 6-tetrahydropyridine (MPTP)-treated mice were used as the PD model and the roles of catalpol on DA neurons and its potential mechanism were investigated in this study. We found that catalpol administration mitigated the loss of DA neurons induced by MPTP and increased exploratory behavior along with tyrosine hydroxylase (TH) expression, which was accompanied by astrocyte and microglia activation. Importantly, catalpol administration significantly inhibited MPTP-triggered oxidative stress, restored growth-associated protein 43 (GAP43) and vascular endothelial growth factor (VEGF) levels. Further, we found that catalpol suppressed the activation of MKK4/JNK/c-Jun signaling, and reduced the pro-inflammatory factors and inflammasome in the mouse model of PD. Our results suggest that catalpol relieves MPTP-triggered oxidative stress, which may benefit to avoid the occurrence of chronic inflammatory reaction. Catalpol alleviates MPTP-triggered oxidative stress and thereby prevents neurodegenerative diseases-related inflammatory reaction, highlighting its therapeutic potential for the management of PD symptoms.

Keywords: Parkinson’s disease, catalpol, neuroprotection, apoptosis, inflammation

INTRODUCTION

Parkinson’s disease (PD) is a common and complex age-related neurological disorder of unclear etiology. The clinical symptoms of PD include motor dysfunction, resting tremor, postural and gait disorders, and neurological impairment (Kalia and Lang, 2015). The pathological hallmarks of PD are the loss of dopamine (DA) neurons and the presence of Lewy bodies, with a resultant reduction in dopamine levels in the substantia nigra (SN) pars compacta and striatum (Bendor et al., 2013; Kordower et al., 2013). Although the underlying molecular mechanisms are not well understood, apoptosis, neuroinflammation, and oxidative stress are thought to be involved in these effects (Imamura et al., 2003; Ghavami et al., 2014; Mashima et al., 2018). For example, activated microglia was shown to contribute to the degeneration of DA neurons in animal models of PD.
Rehmannia glutinosa, has many pharmacological activities on reducing dopaminergic neuron damage (Liu et al., 2018; da Silva et al., 2018). Hence, there is an urgent need to seek new drugs with multiple potentials in alleviating the death of DA neurons in PD.

Catalpol, a water-soluble active compound isolated from Rehmannia glutinosa, has many pharmacological activities including antiapoptotic, anti-inflammatory and antioxidative functions as well as neuroprotective effects (Hu et al., 2010; Wang and Zhan-Sheng, 2018; Yan et al., 2018). Catalpol can cross the blood-brain barrier and serve as a protective agent in the treatment of neurodegenerative diseases (Wang et al., 2012; Dinda et al., 2019). Catalpol exposure reduced tumor necrosis factor-α (TNF-α) and nitric oxide synthase expression in cultured primary microglia that was upregulated by the inflammatory primer, lipopolysaccharide (LPS; Tian et al., 2006), and also protected astrocytes from oxidative damage caused by hydrogen peroxide (Bi et al., 2008a). In other studies, catalpol prevented 1-methyl-4-phenyl-1,2,3,4-tetrahydropyridine (MPTP)/1-methyl-4-phenylpyridinium-induced neurotoxicity in mesencephalic neurons not only by enhancing antioxidant activity, but also by inhibiting monoamine oxidase type B–mediated–neurotoxicity in cells (Mao et al., 2007; Tian et al., 2007; Bi et al., 2008b, 2009; Chen et al., 2008), it also restored locomotor function by increasing the striatal DA concentration and enhanced the level of glial cell-derived neurotrophic factor, thereby preventing the loss of tyrosine hydroxylase (TH)-positive neurons and downregulation of striatal DA transporter (DAT) in an MPTP-induced PD model (Xu et al., 2010). Moreover, catalpol protected against rotenone-induced cell damage by increasing the activities of complex I, glutathione (GSH), GSH peroxidase (GPx) and superoxide dismutase (SOD), which reduced lipid peroxidation and preserved mitochondrial membrane potential (Mao et al., 2007). However, the molecular basis of these effects remains unclear.

In this study, we investigated the neuroprotective effects of catalpol on DA neurons and analyzed its underlying mechanisms using a mouse model of PD. Catalpol treatment prevented DA neurons against apoptosis by downregulating the mitogen-activated protein kinase kinase (MKK) 4/c-Jun N-terminal kinase (JNK)/c-Jun signaling pathway in MPTP-treated mice. Furthermore, catalpol not only reduced the levels of pro-inflammatory factors and inflammasome activation, but it also inhibited oxidative stress in the SN.

MATERIALS AND METHODS

Animals and Drug Treatments

Male C57BL/6 mice (10 weeks old) were obtained from the Department of Laboratory Animal Science of China Medical University. The mice \( n = 11 \) pre-group were intraperitoneally injected with catalpol dissolved in saline (15 mg/kg/day; Chengdu Manster Biotechnology Co., Chengdu, China; A0215) or vehicle (saline) for 3 days, followed by MPTP (30 mg/kg/day; Sigma-Aldrich, St. Louis, MO, USA; M0896), MPTP + catalpol, or vehicle starting on day 4 for 5 days. Mice in the MPTP + catalpol treatment group \( n = 9 \) were continually administered catalpol for 6 days, and those primed with MPTP received the vehicle for 6 days. Mice in the control group were given the vehicle saline \( n = 10 \). The experimental procedure is outlined in Figure 1A. The general health and body weight of animals were monitored daily, and animals were handled according to the Guide for the Care and Use of Medical Laboratory Animals (Ministry of Health, Beijing, China). The experimental protocol was approved by the Laboratory Ethics Committee of China Medical University.

Open Field Test (OFT)

The open field test (OFT) is used to assess spontaneous motor function and exploratory behavior. Mice were allowed to adapt to the new environment before beginning the test, as previously described (Zhang Y. H. et al., 2018). A mouse was placed in the center of the arena and the whole experiment lasted 5 min. The exploratory trace, total distance, distance in the zone center and the times that the mouse enters into the center of the arena were recorded using SMART v.3.0 software (Harvard Apparatus, British).

Pole-Climbing Test

A 50-cm pole was split in half, with each section measuring 25 cm. The middle of the pole was marked. A small ball was placed at the top of the pole and covered with gauze to prevent it from slipping. The mouse was placed on the bar and the time taken until it reached the marked line was recorded. The scores were determined as follows: >6 s, 1 point; 3–6 s, 2 points; and <3 s, 3 points.

Rotarod Test

Locomotor ability was tested using a five-lane IITC 755 Series 8 rotarod (IITC Life Science, Woodland Hills, CA, USA) that was rotated starting at a speed of 4 rpm, which was increased to 40 rpm over a period of 5 min in forwarding mode with a 5-min rest interval. Each mouse was tested three times at each time point.

Tissue Preparation

After behavioral testing, mice were anesthetized by intraperitoneal injection of sodium pentobarbital at a dose of 40 mg/kg. They were then perfused with saline and sacrificed by decapitation. Half of the brain was fixed with 4% paraformaldehyde for morphological analysis. The SN and striatum of the other half of the brain were flash-frozen in dry ice and stored at −80°C until biochemical analyses.

Reactive Oxygen Species (ROS) Measurement

To determine whether catalpol treatment decreases oxidative stress, we detected ROS levels in the SN with the Reactive Oxygen Species Assay Kit (Beyotime Institute of Biotechnology,
Catalpol Protects Dopaminergic Neurons

**FIGURE 1 |** Catalpol alleviates impairment of exploratory behavior in 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP)-treated mice. Mice were intraperitoneally injected with catalpol (15 mg/kg/day) or vehicle (saline) for 3 days, and then administered MPTP (30 mg/kg/day; n = 11), MPTP + catalpol, or vehicle starting on day 4 for 5 days. Mice that were previously treated with MPTP + catalpol (n = 9) were continually administered catalpol for 6 days; those primed with MPTP received the vehicle for 6 days. Mice treated with the vehicle saline served as the control group (n = 10). (A) Treatment schedule. (B) Representative images of movement trials in the open field test (OFT). (C) Quantitative analysis of total distance traveled in the OFT. (D) Distance from the zone center. (E) The number of entries into the zone center in the OFT. (F,G) Time and score in the pole-climbing test. (H) The average amount of time mice remained on the rod in the Rotarod test. Data represent mean ± SEM. The P-values were calculated using a one-way analysis of variance (ANOVA). *P < 0.05 vs. vehicle group; #P < 0.05 vs. MPTP-induced group.

Shanghai, China; S0033). Briefly, the SN was weighed and homogenized in ice-cold phosphate-buffered saline, then centrifuged at 500 × g for 10 min at 4°C. The supernatant was used for the ROS assay. ROS production was evaluated using the oxidant-sensitive probe 2,7-dichlorofluorescein diacetate according to the manufacturer’s instructions. Fluorescence intensity was measured at 485 nm excitation and 525 nm emission wavelengths using a microplate reader (Cytation5; Frontiers in Aging Neuroscience | www.frontiersin.org 3 November 2019 | Volume 11 | Article 316
Determination of SOD Activity

The SOD activity was measured by ultraviolet spectroscopy with the SOD Assay Kit (Jiancheng Biology, Nanjing, China; A001-1) according to the manufacturer’s instructions. Briefly, the SN was removed from the mouse brain and weighed and homogenized in ice-cold SOD sample preparation solution. The lysate was centrifuged at 4,000 × g for 10 min at 4°C, and the supernatant was used for the assay.

Western Blot Analysis

Mouse brain tissue was collected and homogenized by sonication in ice-cold RIPA buffer (Beyotime Institute of Biotechnology; P0013B) containing a protease inhibitor cocktail (Sigma-Aldrich, St. Louis, MO, USA; 8340) for 3 h at 4°C. After centrifugation at 12,000 × g at 4°C for 20 min, protein concentration in the supernatant was determined with bicinchoninic acid protein assay reagent (Beyotime Institute of Biotechnology; P0010). The supernatant containing 25 μg of protein was resolved by 12% sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and the proteins were transferred to a polyvinylidene difluoride membrane (Millipore, Billerica, MA, USA; IPVH00010) that was probed with the following antibodies: rabbit anti-TH (1:1,000, Millipore); rabbit anti-DAT (1:500), rabbit anti-SOD1 (1:500), rabbit anti-GPx4 (1:500), mouse anti-glyceraldehyde 3-phosphate dehydrogenase (GAPDH; 1:10,000), rabbit anti-β-actin (1:10,000), rabbit anti-growth-associated protein (GAP) 43 (1:500), and rabbit anti-vascular endothelial growth factor (VEGF; 1:500; all from Proteintech, Rosemont, IL, USA); mouse anti-B cell lymphoma (Bcl) 2 (1:1,000), mouse anti-Bcl2-associated × protein (BAX; 1:1,000), rabbit anti-caspase-3 (1:1,000), mouse anti-caspase-9 (1:1,000), rabbit anti-p-MKK4 (1:1,000), rabbit anti-MKK4 (1:1,000), rabbit anti-p-c-Jun (1:1,000), rabbit anti-c-Jun (1:1,000), rabbit anti-p-JNK (1:1,000), rabbit anti-JNK (1:1,000), rabbit anti-nucleotide-binding domain (NBD) and leucine-rich-repeat-containing (LRR; known as NLR family member) × 1 (NLRX1; 1:1,000), rabbit anti-IL-1β (1:1,000), and rabbit anti-glial fibrillary acidic protein (GFAP; 1:1,000; all from Cell Signaling Technology, Danvers, MA, USA); mouse anti- ionized calcium binding adaptor molecule (Iba)1 (1:1,000, Thermo Fisher Scientific, Waltham, MA, USA); and rabbit anti-α-synuclein (1:1,000), rabbit anti-TNF-α (1:500), and rabbit anti-nucleotide binding oligomerization domain-like receptor, pyrin domain-containing (NLPR) 3 (1:1,000; all from Abcam, Cambridge, MA, USA). The membrane was incubated with horseradish peroxidase-labeled secondary antibodies (1:10,000, Thermo Fisher Scientific, Waltham, MA, USA), and protein band intensity was quantified using Prism v.7.0 software (Graph Pad, La Jolla, CA, USA).

Immunohistochemistry and Immunofluorescence

Brain tissue sections were incubated overnight at 4°C with rabbit anti-TH antibody (1:100), then treated with the appropriate biotinylated secondary antibody for 2 h at room temperature followed by a third antibody for 30 min (MXB, Fuzhou, China). The sections were developed in diaminobenzidine (Sigma-Aldrich, St. Louis, MO, USA) for 3–5 min, then immersed in distilled water to halt the reaction. The sections were dehydrated and sealed with neutral gum and images were acquired on a light microscope (ECHO, San Diego, CA, USA). For immunofluorescence labeling, mouse brain sections were blocked with sheep serum (Beyotime Institute of Biotechnology) at room temperature for 1 h, then incubated overnight at 4°C with a mixture of primary antibodies consisting of mouse anti-GFAP (1:100) and rabbit anti-TH (1:100), or mouse anti-TH (1:100; ImmunoStar, Hudson, WI, USA) and rabbit anti-Iba1 (1:400; Wako Pure Chemical Industries). After rinsing, the sections were incubated with goat anti-rabbit IgG (H + L) and goat anti-mouse IgG (H + L) cross-adsorbed secondary antibodies (both 1:500, Thermo Fisher Scientific, Waltham, MA, USA) for 2 h at room temperature. Images were acquired with a confocal laser scanning microscope (A1; Nikon, Tokyo, Japan).

Statistical Analysis

Data are represented as mean ± standard error of the mean (SEM), and each set of experiments was repeated three times in parallel. Differences among means were compared using one-way ANOVA. Data were analyzed using Prism v.7.0 software. The results were considered significant at $P < 0.05$.

RESULTS

Catalpol Enhances Exploratory Behavior in MPTP-Treated Mice

To investigate whether catalpol alters the motor function and exploratory behavior in MPTP-treated mice, we used the OFT, pole climbing test and Rotarod test to assess motor ability. The distance traveled and the number of entries in the zone center were greater in the MPTP + catalpol group than in the MPTP group, whereas total distance traveled was similar across groups (Figures 1B–E). We then assessed the effect of catalpol on locomotion ability with the pole climbing and Rotarod tests but found no significant differences in scores between the three groups in either test (Figures 1F–H). These results indicate that catalpol reverses the reduction in exploratory behavior caused by MPTP treatment. MPTP or catalpol treatment did not markedly affect the body weight of the mice (Supplementary Figure S1).
FIGURE 2 | Catalpol rescues dopaminergic (DA) neurons in the SN and striatum in an MPTP-induced mouse model of Parkinson’s disease (PD). (A) Immunohistochemical detection of tyrosine hydroxylase (TH) in DA neurons of the substantia nigra (SN) and striatum of mice treated with vehicle, MPTP, or MPTP + catalpol. Scale bar = 100 µm. (B-E) Western blot analysis of TH and α-synuclein (α-syn) levels in the SN. (F-I) Protein levels of TH and DA transporter (DAT) as detected by western blotting. Data represent mean ± SEM; n = 7 mice per group. The P-values were calculated using one-way ANOVA. *P < 0.05 vs. vehicle group; #P < 0.05, ##P < 0.01 vs. MPTP-induced group.
of JNK activation, we evaluated MKK4 phosphorylation status.

Catalpol Modulates MKK4/JNK/c-Jun Signaling to Inhibit the Apoptosis of DA Neurons

As JNK/c-Jun signaling is important for apoptosis, we next examined whether catalpol inhibits apoptosis by regulating this pathway. A western blot analysis revealed that MPTP increased JNK and c-Jun phosphorylation, which was reversed by catalpol (Figures 4C–F). Given that MKK4 signaling is a key regulator of JNK activation, we evaluated MKK4 phosphorylation status in the SN and determined that it was altered by catalpol, although total MKK4 protein level was unchanged compared to the MPTP-treated group (Figures 4A,B). These findings indicate that catalpol blocks apoptosis via modulation of MKK4/JNK/c-Jun signaling.

Catalpol Exerts Antioxidant Effects in MPTP-Treated Mice

Oxidative stress is a major cause of apoptosis in DA neurons (Haddad, 2002), we, therefore, evaluated ROS levels in the SN and found that MPTP enhanced ROS levels (Figure 5A). ROS levels were decreased in the presence of catalpol, which also reversed the MPTP-induced suppression of SOD1 activity and protein expression in the SN (Figures 5B–D). Moreover, western blot analysis showed that catalpol treatment restored the expression of the antioxidant enzyme GPx4, which was downregulated in the presence of MPTP (Figures 5C,E). Given that NLRX1 enhances ROS production, we examined whether catalpol-mediated inhibition of ROS is involved in the suppression of NLRX1 and found that expression of the protein was reduced by treatment with catalpol relative to the level in mice treated with MPTP only (Figures 5B,F). These results indicate that catalpol alleviates the apoptosis of DA neurons through its antioxidant capacity.

Catalpol Attenuates Inflammation in MPTP-Injured Mice

PD development and progression are accompanied by an inflammatory response. To determine whether catalpol reduces inflammation in PD, we evaluated the activation astrocyte and microglia based on immunohistochemical detection of GFAP and Iba1 expression, respectively. The immunopositivity of GFAP and Iba1 was increased in mice treated with MPTP compared to the vehicle. However, catalpol had the opposite effect (Figures 6A,B). Consistent with these findings, the western blot analysis showed that GFAP and Iba1 protein levels were elevated in MPTP-treated mice, but these were blocked by catalpol (Figures 7A–C). We also found that catalpol inhibited MPTP-induced inflammation as evidenced by the downregulation of TNF-α, interleukin (IL)-1β and NLRP3 protein expression compared to mice treated with MPTP only (Figures 7A,D–F).

GAP43 promotes nerve regeneration and the establishment of synaptic connections (Allegra Mascaro et al., 2013; Kettenmann et al., 2013). Moreover, the inhibition of GAP43 may be due to the increase of apoptosis (Wang Z. et al., 2018). To assess MPTP-induced injury of DA neurons and to investigate whether catalpol is beneficial for regeneration, we determined by western blotting that GAP43 protein expression was lower in mice treated with MPTP as compared to vehicle (Figures 7G,H). However, catalpol administration restored GAP43 protein level. In addition to its proangiogenic role, VEGF also promotes neuronal survival (Greenberg and Jin, 2005; Okabe et al., 2014); here we found that MPTP reduced VEGF protein expression, which was restored by catalpol (Figures 7A,I). These data indicate that catalpol protects DA neurons against MPTP-induced injury by inhibiting inflammation and stimulating neural regeneration.
DISCUSSION

Inflammation is implicated in the pathogenesis of PD (Liu et al., 2003; Herrero et al., 2015; Ransohoff, 2016) increases in innate immune components such as interleukin (IL)-1β, IL-6 and TNF-α have been detected in the cerebrospinal fluid and SN of PD patients (Fiszer et al., 1994; Liu et al., 2003). Additionally, in postmortem studies and animal models of PD, ROS and nitrogen species were found to be increased, which was accompanied by activation of microglia and reactive astrocytes (Herrero et al., 2015; Ransohoff, 2016). Serum proinflammatory cytokine concentrations are also increased in PD patients (Dobbs et al., 1999; Scalzo et al., 2010). In rodents, inflammatory stimuli including bacterial LPS (Qin et al., 2007) and viral pathogens (Ogata et al., 1997) contribute to the loss of DA neurons. Bacterial or viral infection may increase the onset of PD in a susceptible population (Jang et al., 2009). In animal models of MPTP-induced PD (Dauer and Przedborski, 2003), TNF-α level in the brain was increased, implying that adaptive immunity was activated in response to MPTP-induced neurodegeneration (Barcia et al., 2011). Peripheral and/or central neuroinflammation is associated with the prodromal phase of PD (Hirsch and Hunot, 2009); thus, inflammation is an important consideration when investigating the mechanisms underlying the loss of DA neurons in PD. Interestingly, LPS can cause prolonged activation of MKK4 (Waetzig et al., 2005), leading to the phosphorylation of JNK, which mediates the proinflammatory function of microglia (Verrecchia et al., 2003; Waetzig et al., 2005). Phosphorylated JNK stimulates c-Jun activity to enhance the transcription of pro-apoptotic cytokines (Whitfield et al., 2001; Dunn et al., 2002). It was reported that anti-inflammatory intervention can reduce the risk of developing PD (Noyce et al., 2012), while pharmacological or genetic inhibition of inflammatory factors...
FIGURE 5 | Protective effects of catalpol against oxidative stress in the SN of MPTP-treated mice. (A) SOD1, NLRX1, and GPX4 protein expression as determined by western blotting. Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) served as the loading control. (B–D) Quantitative analysis of SOD1, NLRX1, and GPX4 protein levels. (E) Changes in SOD1 activity in the SN of mice. (F) Reactive oxygen species (ROS) production in the SN detected with the dichlorofluorescein diacetate probe. Data represent mean ± SEM; n = 6–8 mice per group. The P-values were calculated using one-way ANOVA. *P < 0.05 vs. vehicle group; #P < 0.05, ##P < 0.01 vs. MPTP-induced group.

conferred protection against MPTP-induced neurodegeneration (Du et al., 2001; Nomura et al., 2011; Świątkiewicz et al., 2013). The bioflavonoid compound pycnogenol, which is commonly used as a dietary supplement to suppress inflammation, alleviated MPTP-associated motor impairment and the inflammatory responses of astrocytes and microglia in a mouse model of PD (Khan et al., 2013). Likewise, minocycline which has anti-inflammatory properties blocked the activation of macrophages in the brain to protect against apoptosis in vitro (Duan et al., 2002) and in animal models of PD (Du et al., 2001; Wu et al., 2002). Thus, neuroinflammation is a major mechanism underlying PD, and early intervention with anti-inflammatory agents is a potential strategy for inhibiting PD progression. It is worth noting that Dinda et al. (2019) compared the effects of catalpol, geniposide, and harpagoside. Catalpol exhibits significant advantages of neuroprotective potential in the intervention of PD (Dinda et al., 2019). Catalpol could protect the neurons from MPP⁺-induced toxicity in vitro (Tian et al., 2006). Catalpol can pass the blood-brain barrier. Catalpol treatment markedly protected the neurons of PD mice brain from degeneration through improving antioxidant status (Mao et al., 2007). Catalpol obviously mitigated the loss of dopamine transporter density and TH positive neurons and improved the locomotor ability of PD mice (Xu et al., 2010). We previously reported that catalpol could alleviate β-amyloid toxicity and neuroinflammatory reactions (Wang N. et al., 2018). It is considered that catalpol has the potential property of slowing down the neurodegenerative process in PD.

In this study, the inflammatory response in the mouse brain was accompanied by MPTP-induced neuronal apoptosis, as evidenced by the upregulation of cleaved caspase-3 and cleaved caspase-9 both effects in the extrinsic apoptosis pathway (Zou et al., 1997) relative to vehicle-treated control animals. Additionally, the decrease in Bcl2/Bax ratio upon MPTP treatment indicated that the intrinsic apoptosis pathway was activated (Falschlehner et al., 2007). The presence of reactive
astrocytes and activated microglia confirmed MPTP-induced inflammation in our mouse model of PD, whereas the downregulation of GAP43 and VEGF proteins reflected MPTP-mediated neurodegeneration. It is generally accepted that GAP43 is an important biomarker for adult neurogenesis. GAP43 decline is related to the increase of apoptotic cells (Wang Z. et al., 2018). Meanwhile, the behavioral deficits observed in MPTP-treated mice were consistent with nigrostriatal DA neuron injury. The schemes of MPTP-caused PD-like pathology include three types: acute, subacute, and chronic (Petroske et al., 2001; Selvakumar et al., 2014; Ren et al., 2015). The subacute model is often chosen to investigate the effects of interventions due to its advantages in the period and the similarities to PD. Whereas, MPTP subacute administration-induced alterations and the proper positive control are still needed to be further confirmed. In the present study, the subacute MPTP paradigm failed to trigger the typical movement deficits despite the dopaminergic neuron damage. Instead of motor defects, the results of the OFT in our study showed that MPTP-managed mice exhibited behavioral defects relative to controls. Nonmotor characteristics of PD with nigrostriatal circuit degeneration perform anxiety-like behaviors (Santiago et al., 2010). So, in the present study, it is important to assess the validity of a candidate based on its effects on the dopaminergic neurons and behavioral examination related to nonmotor characteristics of PD. It is reported that catalpol reduced inflammatory cytokine levels and improved cholinergic function in a mouse model of senescence (Zhang et al., 2013), and protected pre-myelinating oligodendrocytes against ischemia-induced injury in vitro (Cai et al., 2016) while alleviating atherosclerosis (Zhang Y. et al., 2018) and modulating Bcl-2/Bax ratio to suppress endothelial cells apoptosis induced by hydrogen peroxide (Hu et al., 2010). We selected catalpol as our candidate for PD intervention. It has been reported that MPTP treatment could cause olfaction dysfunction, gastrointestinal injury, bladder impairment, and other nonmotor symptoms except for

FIGURE 6 | Catalpol treatment suppresses microglia and astrocyte activity induced by MPTP. (A) Frozen sections of the SN were labeled with anti-TH (red) and anti-GFAP (green) antibodies to detect DA neurons and astrocyte activity, respectively. Scale bar = 100 µm. (B) Brain sections were labeled with anti-TH (green) and anti-Iba1 (red) antibodies to assess microglia activity in the SN. Scale bar = 100 µm; n = 6.
FIGURE 7 | Catalpol suppresses MPTP-induced inflammation and promotes neural regeneration in the SN of mice. (A) Western blot analysis of GFAP, Iba1, IL-1β, TNF-α, and NLRP3 protein expression in the SN of mice. (B–F) Catalpol blocks MPTP-induced astrocyte and microglia activation inhibits IL-1β and TNF-α, and suppresses NLRP3 production. (G–I) Growth-associated protein 43 (GAP43) and vascular endothelial growth factor (VEGF) protein expression detected by western blotting. Data represent mean ± SEM; n = 7 mice per group. The P-values were calculated using one-way ANOVA. *P < 0.05 vs. vehicle group; #P < 0.05, ##P < 0.01 vs. MPTP-induced group.

near-total damage of the striatal dopaminergic input (Taylor et al., 2010; Duty and Jenner, 2011). Several studies have shown that antioxidants pretreatment prior to the induction of PD with MPTP could provide more effective neuroprotection against the cytotoxic effects (Lee et al., 2011; Wang et al., 2015). We then administrated the mice by intraperitoneal injection with catalpol ahead of MPTP treatment. Consistent with previous studies (Bi et al., 2008b; Jiang et al., 2008), we demonstrated that catalpol has neuroprotective effects in PD, as evidenced by the restoration of DA neurons in the SN and striatum of MPTP-treated mice and the corresponding increase in exploratory behavior. Immunohistochemical detection of TH showed that catalpol intervention prevented MPTP-induced loss of TH+ cells, suggesting that it can improve the TH deficiency observed in PD patients (Moore et al., 2005). Catalpol treatment alleviated the MPTP-induced downregulation of GAP43 and VEGF proteins, which suggested that catalpol may induce regeneration. Although no significant differences were observed in the scores during the pole climbing and rotarod tests among the vehicle-, MPTP- and MPTP plus catalpol-treated mice, in the OFT, MPTP-lesioned mice exhibited significant thigmotaxis, traveling a shorter total distance and passing fewer times through the center of the open field relative to the vehicle group, whereas, these behavioral impairments were abolished by catalpol treatment. Our findings demonstrated that catalpol could improve some behavioral defects in PD. Xu et al. (2010) observed significant locomotor impairments in the MPTP-triggered PD mouse model, and catalpol treatment could improve the motor performances of these mice. Those incongruences are in contrast with our results perhaps because of the differences in the PD mouse model type (an MPTP-induced chronic paradigm), the dose of catalpol intervention (high dose at 50 mg/kg) and an extended period of time (8 weeks of catalpol treatment) in Xu’s study.

In obese mice, catalpol was shown to mitigate inflammation by blocking JNK signaling (Santiago et al., 2010). In the present study, MPTP induced an increase in MKK4 and JNK phosphorylation, which was accompanied by dopamine depletion in a mouse model of PD. The increased phosphorylation of c-Jun in the brain of MPTP-treated mice suggested an enhancement of c-Jun transcription through JNK activation (Zhou et al., 2015) as previously reported (Derijard et al., 1994), suggesting that MPTP-induced MKK4/JNK/c-Jun signaling is involved in DA neurons degeneration in PD. Catalpol treatment reversed the MPTP-induced phosphorylation of MKK4 and JNK and c-Jun expression, suggesting that it suppresses MKK4/JNK/c-Jun signaling. Given the critical role of JNK in linking inflammation and apoptosis (Dunn et al., 2002; Waetzig et al., 2005; Perier et al., 2007), catalpol may prevent neurodegeneration by targeting this pathway.
Oxidative stress promotes inflammation and causes cell damage (Haddad, 2002). In general, inflammation is a protective response against cell injury that involves the destruction and removal of detrimental components to restore cellular function. DA neurons are highly susceptible to ROS-mediated injury (Miyazaki and Asanuma, 2008); on the other hand, ROS activate pro-inflammatory signaling in glia, thereby stimulating a sustained inflammatory response. Chronic inflammation ultimately destroys normal tissue. TNF-α, which is released by reactive astrocytes and activated microglia, modulates oxidative stress and inflammation in neurodegenerative diseases (Fischer and Maier, 2015). Elevated TNF-α levels due to systemic inflammation are associated with the decline of DA neurons in PD (Sriram et al., 2006; Allegra Mascaro et al., 2013), and TNF-α level was found to be elevated in the midbrain of a PD mouse model; in fact, the initial oxidative insult in the induction of PD caused a loss of ventral midbrain DA neurons that triggered an inflammatory response (Srivastava et al., 2012). TNF-α promotes neurodegeneration by stimulating ROS production and release. In our study, TNF-α and IL-1β levels were increased in the brain of MPTP-treated mice relative to controls, which is in accordance with previous findings (Chung et al., 2010; Lofrumento et al., 2011). Interestingly, catalpol administration reversed the MPTP-induced decreases in SOD1, NLRX1, and GPX4 protein expression, indicating that the antioxidant defense capacity was enhanced. Moreover, catalpol reduced the levels of the proinflammatory cytokines TNF-α and IL-1β and NLRP3 inflammasome components. These results indicate that catalpol mitigates MPTP-induced oxidative stress to prevent chronic inflammation and neurodegeneration.

In summary, the results of this study demonstrate that catalpol treatment alleviates MPTP-induced degeneration of DA neurons in vivo. The neuroprotective effects of catalpol are at least partly due to the attenuation of inflammation, enhance antioxidant defense and reduce oxidative stress, and inhibition of apoptosis in DA neurons of the SN and striatum through downregulation of MKK4/JNK/c-Jun signaling. Our findings highlight the clinical potential of catalpol for the management of PD symptoms. The pre-treatment with candidate agents for PD intervention is limited by the specific biomarkers that could confirm the asymptomatic stage of PD. Extensive experiments, pre-clinical and clinical evaluation need to be performed.

DATA AVAILABILITY STATEMENT

All datasets generated for this study are included in the article/Supplementary Material.

ETHICS STATEMENT

The animal study was reviewed and approved by The Laboratory Ethics Committee of China Medical University.

AUTHOR CONTRIBUTIONS

L-YW and XY conceived the study and designed the experiments. X-XL and Y-NZ performed the experiments. L-YW and XY interpreted the data and wrote the manuscript. Z-YH critically reviewed the manuscript and obtained funding for the study. All authors have read and approved the final version of the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fnagi.2019.00316/full#supplementary-material.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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