Neuroprotective Effects of Resveratrol on a Mouse Model of Parkinson’s Disease via the Wnt/Beta-Catenin Signaling Pathway

YIYING DONG, Z. WANG¹, Z. YAN², C. WU², T. YANG²* AND HANCHENG QIU³

Department of Biomedical Engineering, Medical Center of Chinese People’s Liberation Army (PLA) General Hospital, Beijing 100853, ¹Department of Neurosurgery, Shenyang Jishuitan Hospital, Shenyang, Liaoning 110005, ²Beijing Huilin Zegu Biotechnology Co., Ltd., Haidian, Beijing 100039, ³Department of Neurosurgery, Beijing Tiantan Hospital, Capital Medical University, Fengtai, Beijing 100054, China

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We herein aimed to evaluate the neuroprotective effects of resveratrol on a mouse model of Parkinson’s disease and to explore whether the Wnt/beta-catenin signaling pathway was involved. Thirty male C57 black 6 mice were randomly divided into a control group, a model group (1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine) and a treatment group (1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine resveratrol) (n=10). The model and treatment groups were intraperitoneally injected with 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (30 mg/kg in normal saline) to induce Parkinson’s disease. Then the treatment group was administered with resveratrol (20 mg/kg in 0.5 % sodium carboxymethylcellulose solution) on the 1st d of modeling. The motor coordination ability was studied by the pole climbing test. Astrocyte and microglia activations in the substantia nigra pars compacta were observed by immunohistochemical staining. 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine-induced striatum dopamine release was measured by high performance liquid chromatography and dopamine transporter level was detected with Western blot. SH-SY5Y cell viability and dependence on the Wnt/beta-catenin signaling pathway were assessed by 3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide assay and Western blot, respectively. Resveratrol relieved the motor dysfunction, death of dopaminergic neurons, activation of astrocytes and microglia, as well as reduction of striatum dopamine content induced by neurotoxin 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine in vitro. It also alleviated the damage of SH-SY5Y cells induced by 1-methyl-4-phenylpyridinium. Resveratrol increased the levels of proteins from the Wnt/beta-catenin signaling pathway in the mouse model of Parkinson’s disease and 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine-treated SH-Y5Y cells. Resveratrol exerted neuroprotective effects on the Parkinson’s disease model predominantly by regulating the Wnt/beta-catenin signaling pathway.

Key words: Parkinson’s disease, resveratrol, dopamine, Wnt/beta-catenin

Parkinson’s Disease (PD) is a common progressive neurodegenerative dyskinesia with the prevalence rate of about 1 % among the elderly aged over 65 y old. The main clinical manifestations include tremor, stiffness, bradykinesia and gait apraxia, as well as autonomic dysfunction, sleep disorders and Alzheimer’s disease in the later stage[1,2]. PD is pathologically typified by the progressive loss of dopaminergic neurons in the Substantia Nigra pars compacta (SNpc), but its exact pathogenesis remains unclear[3]. Levodopa replacement therapy is still the first-line treatment for PD, but the progression cannot be prevented, accompanied by many serious adverse reactions after long-term use, such as autonomic symptoms (nausea, orthostatic hypotension), daytime sleepiness and psychiatric complications (illusion, delusion)[4]. Therefore, researchers have endeavored to find new targets for developing drugs for PD, aiming to improve the quality of life of patients.

The Wnt protein family consists of 19 glycosylated proteins that regulate cell growth and differentiation[5]. It is also an important mediator of cell-to-cell communication and signal transduction during the development of the central nervous system. Wnt ligands are a class of secreted glycoproteins that work through paracrine or autocrine. Besides, the Wnt/Beta (β)-catenin signaling pathway (Wnt1 or Wnt3a) and Wnt signaling

*Address for correspondence
E-mail: yangtao686@163.com

Accepted 21 July 2022
Revised 11 December 2021
Received 24 June 2021
Indian J Pharm Sci 2022;84(4):959-968
essentially participate in the development of midbrain dopaminergic neurons[6]. Wnt1 is involved in midbrain formation, progenitor cell differentiation in SNpc and survival of dopaminergic neurons.

Resveratrol is a natural polyphenolic chemical widely existing in grape skin and other plants, with biological characteristics such as antioxidation, anti-inflammation, anti-aging, antitumor and relief of the endoplasmic reticulum stress. At present, studies regarding the protective effects of resveratrol on dopaminergic neurons focus on resistance to oxidation and inflammation[7]. Nevertheless, the role of the Wnt/β-catenin signaling pathway has seldom been referred. Thereby motivated, we herein aimed to clarify the mechanism by which the Wnt/β-catenin signaling pathway participated in the neuroprotective effects of resveratrol on PD through establishing in vivo and in vitro models using SH-SY5Y cells treated with 1-Methyl-4-Phenylpyridinium (MPP+).

MATERIALS AND METHODS

Animals and cell line:

This study has been approved by the animal ethics committee of Beijing Huilin Zegu Biotechnology Co., Ltd., and great efforts have been made to alleviate the suffering of animals. Thirty Specific Pathogen Free (SPF) male C57 Black 6 (C57BL/6) mice aged 12 w old and weighing 24-26 g were purchased from SPF (Beijing) Biotechnology Co., Ltd. (China, certificate of conformity: SCXK (jing) 2019-0010). They were fed in standard feeding boxes at 20°-26° with a 12 h/12 h light/dark cycle.

Reagents and antibodies:

Resveratrol (purity: ≥98 %), 1-Methyl-4-Phenyl-1,2,3,6-Tetrahydropyridine (MPTP) and 3-(4,5-Dimethylthiazol-2-yl)-2,5-Diphenyltetrazolium Bromide (MTT) were obtained from Sigma (St. Louis, MO, USA). Dulbecco’s Modified Eagle Medium (DMEM), Opti-MEM, Non-Essential Amino Acids (NEAA) and penicillin were purchased from Gibco (Grand Island, NY, USA). Fetal Bovine Serum (FBS) was bought from Scientell (Carlsbad, California (CA), United States of America (USA)). Wnt1 antibody was obtained from Santa Cruz Biotechnology (Santa Cruz, CA, USA). P-Ser9-Glycogen Synthase Kinase 3 Beta (GSK3β) and β-catenin antibodies were provided by Cell Signaling Technology (Beverly, Massachusetts (MA), USA). Dopamine Transporter (DAT) and β-actin antibodies were purchased from Millipore (MA, USA). All other drugs were obtained from Sigma (St. Louis, MO, USA).

Animal grouping and treatment:

Thirty mice were randomly divided into a control group, a model group (MPTP) and a treatment group (MPTP+resveratrol) (n=10). The model and treatment groups were intraperitoneally injected with MPTP (30 mg/kg in normal saline) to induce PD, once every 3.5 d for 5 w (a total of 10 injections). The control group was intraperitoneally injected with a corresponding volume of normal saline. The treatment group was administered with resveratrol (20 mg/kg in 0.5 % sodium carboxymethylcellulose solution) on the 1st d of modeling, once a day for 5 w. The control and model groups were intragastrically treated with corresponding volumes of 0.5 % sodium carboxymethylcellulose solution.

Animal behaviors:

The pole climbing test was used to investigate the motor coordination ability of a mouse[8]. A straight wooden rod with a diameter of 10 mm and a height of 500 mm was used, and a cork ball with a diameter of 25 mm was placed on the top of the rod. The mouse head was placed upside down on the top of the rod and the following three time points were recorded; T1 for the time from the start of movement to the time when the head was turned downward; T2 for the time of climbing down to the middle of pole and T3 for the time of climbing to the bottom of pole from the middle. The above activities were scored as follows, 3 points for shorter than 3 s; 2 points for shorter than 6 s but longer than 3 s and 1 point for over 6 s. Finally, the sum of them (T1+T2+T3) was recorded. The mice were trained for 3 d before the test. Detection was conducted three times during the test, with the interval of 1 min each time and the shortest time was recorded. If the mouse failed to completely flip, slip or drop, the score was 3 points. The pole climbing ability was evaluated 1 w before establishment of the MPTP model and 3 h, 1 d, 3 d and 7 d after it.

Immunohistochemical staining:

The mice were anesthetized through intraperitoneal injection with 0.004 ml/g of 10 % chloral hydrate. After anesthesia, the heart was exposed. A scalp needle was inserted into the left ventricle and the auricula dextra was cut open, into which normal saline and 4 % paraformaldehyde were perfused. The mouse brain was taken out immediately and fixed in 4 % paraformaldehyde overnight. Afterwards, the brain was dehydrated in 20 % saccharose solution for 3 d and in 30 % saccharose solution for another 3 d and stored at -70°. Frozen tissue
was sectioned into 30 μm thick. The brain sections were collected in Phosphate-Buffered Saline (PBS) and stored at 20° prior to use. After being washed with PBS at room temperature, the sections were incubated with 3 % Hydrogen peroxide (H₂O₂) for 30 min to eliminate endogenous horseradish peroxidase and then washed three times with PBS on a shaker, 5 min each time. At room temperature, the sections were blocked by using 5 % Bovine Serum Albumin (BSA) (prepared with PBS/0.1 % Triton X-100) for 1 h, incubated with mouse anti-Tyrosine Hydroxylase (TH) (1:3000), anti-Glial Fibrillary Acidic Protein (GFAP) (1:800) or anti-Macrophage-1 (MAC-1) (1:100) primary antibody overnight at 4° and washed with 0.01 M PBS 3 times, 5 min each time. Subsequently, they were incubated with corresponding horseradish peroxidase-conjugated secondary antibodies for 1 h at room temperature and color-developed with diaminobenzidine. Stereoscopic counting was performed for TH-positive cells in SNpc, GFAP-positive astrocytes and MAC-1-positive microglia.

Measurement of dopamine levels in the striatum:

Dopamine level was measured using High Performance Liquid Chromatography (HPLC) with LC-20A pump (Shimadzu, Kyoto, Japan) and RF-20A fluorescence detector (Shimadzu, Kyoto, Japan). Chromatographic separation of dopamine was conducted using Thermo Hypersil GOLD™ C₁₈ column (4.6×150 mm, i.d., 5 μm; Thermo Fisher Scientific Inc., MA, USA). Mobile phase A consisted of 30 mM citric acid, 40 mM sodium acetate anhydrous, 0.2 mM Ethylenediaminetetraacetic Acid Disodium (EDTA-2Na) salt and 0.4 mM sodium octane sulfonate (pH=3.9) and phase B was methanol. The ratio of phase A to phase B was 86:14 and the flow rate was 1.0 ml/min. Dopamine level was calculated based on the peak area of standard curve using Lab Solutions LC-solution Version 1.2 Workstation (Shimadzu, Kyoto, Japan).

Cell culture and grouping:

SH-SY5Y cells were cultured in DMEM containing 10 % FBS, 1 % NEAA, 100 U/ml penicillin and 0.1 U/ml streptomycin at 37° with 5 % Carbon dioxide (CO₂). The cells were cultured in sterile plastic culture flasks at the density of 1×10⁵/cm² and trypsinized and passaged every 3 d. Then the cells were seeded into 96 or 6-well plates grown cells were digested, diluted into a suspension by 0.25 % trypsin-EDTA and passaged every 24 h. Drug treatment was started after the culture medium was replaced by serum-free one.

MTT assay:

The cell viability was tested by the MTT assay. Well-grown cells were digested, diluted into a suspension by DMEM containing 10 % FBS, seeded into 96-well plates (200 μl each well) and incubated at 37° with 5 % CO₂ for 24 h. After the culture medium was replaced by serum-free medium, the cells were treated for 24 h as mentioned above, from which the medium was discarded. After addition of 200 μl of 0.5 mg/ml MTT solution prepared by PBS, the cells were further cultured at 37° with 5 % CO₂ for 4 h and MTT solution was removed. Subsequently, 200 μl of Dimethyl Sulfoxide (DMSO) was added and the plates were shaken at room temperature for 15 min. The Optical Density (OD) of each well was measured at 450 nm by a micro plate reader. Cell viability was determined by using the percentage of OD of drug treatment group to that of the control group.

Western blotting:

1 w after MPTP administration, brain tissues were rapidly removed onto ice, added in a proportion of 1:10 (v/v) to buffer containing protease inhibitor cocktail (Life Technologies Corp., Carlsbad, CA, USA), ultrasonicated and incubated on ice for 40 min. Cell lysates were centrifuged at 12 000 ×g for 15 min, and the supernatants were collected as whole-cell protein samples. A 1 μl sample of lysate was used to measure the total protein concentration. The samples were mixed with 5× loading buffer and denatured in boiling water for 5 min before storage at -20° for future use.

According to the Bicinchoninic Acid (BCA) quantification results, 20 μg protein was loaded on each lane of 10 % polyacrylamide gel and separated by Sodium Dodecyl Sulfate-Polyacrylamide Gel Electrophoresis (SDS-PAGE) (80 V). Separated proteins were electronically transferred onto polyvinylidene fluoride membranes at 300 mA constant current for 90 min. Blotted membranes were incubated in Tris Buffered Saline+Tween 20 (TBST) (pH 7.4, 10 mM Tris-HCl, 150 mM Sodium chloride (NaCl), 0.1 % Tween-20) containing 5 % non-fat milk for 1 h and then in 5 % BSA-TBST containing one of the following primary antibodies overnight at 4° anti-Wnt1 (1:200), anti-p-Ser9-GSK3β (1:1000), anti-β-catenin (1:1000) or anti-β-actin (1:1000). Immunolabeled membranes were washed three times in TBST and incubated with secondary antibodies for 1 h, followed by three washes with TBST. Finally, immunoblots were scanned by Gel Max imager (Ultra-Violet Products Ltd., Upland, CA, USA). Band
density was measured by Gel Pro Analyzer software
(Media Cybernetics, Inc., Bethesda, MD, USA).

**Statistical analysis:**

All data are represented as mean±Standard Error of the
Mean (SEM). Treatment group means were compared by
one-way or two-way analysis of variance. When analysis
of variance showed significant differences among
groups, pairwise comparisons were carried out by the
Newman-Keuls post hoc test. For all analyses, p<0.05
was considered statistically significant. All statistical
analyses were performed with IBM Statistical Package
for the Social Sciences (SPSS) software (version 21.0,
USA).

**RESULTS AND DISCUSSION**

A mouse model of MPTP was established to evaluate
the protective effects of resveratrol on neurological and
motor deficits. The mice showed obvious acute reaction
after a few min of intraperitoneal injection of MPTP,
such as tail stiffness, vertical hair, decreased activity, etc.,
the symptoms were alleviated after 12 h, but the activity
of the mice was still significantly reduced. In contrast,
the control group did not undergo motor dysfunction. At
3 h as well as on the 1st, 3rd and 7th d after intraperitoneal
injection of MPTP, the behavior of climbing rods was
observed. The rats showed obvious motor dysfunction
3 h and 1 d after intraperitoneal injection of MPTP. The
time required for climbing was significantly increased.
Resveratrol pretreatment significantly increased the
motor function scores of mice on 1 d and 3 d after
intraperitoneal injection of MPTP compared with MPTP
as shown in fig. 1.

In the pole climbing test, resveratrol significantly
alleviated the motor dysfunction of PD mice induced by
MPTP. Consistent with behavioral testing, resveratrol
attenuated the neurotoxicity of MPTP and protected
dopaminergic neurons. Immunohistochemical staining
was used to observe the dopaminergic neurons in SNpc.
Intraperitoneal injection of MPTP reduced the number
of TH positive cells in SNpc. Compared with the MPTP
group; pretreatment with resveratrol significantly
increased the number of TH positive cells in SNpc as
shown in fig. 2.

To further investigate the effects of resveratrol on

![Fig. 1: Pole test scores for each animal were calculated following acute MPTP treatment after 3 h, 1 d, 3 d and 7 d
Note: Data are represented as mean±SEM of 10 individual experiments, (       ) Control; (       ) MPTP and (       ) Res+MPTP group]
MPTP-induced activation of mouse astrocytes. GFAP staining showed the number of GFAP-positive astrocytes in SNpc. Only a weak GFAP-positive astrocyte immune response was observed in SNpc of the control mice. Intraperitoneal injection of MPTP increased the number of GFAP-positive astrocytes in SNpc 3.12-fold compared with that of the control group. Resveratrol pretreatment inhibited MPTP-induced activation of astrocytes (p<0.001, compared with MPTP group) as shown in fig. 3.

In this study, MAC-1 staining was used to label activated microglia to study the effect of resveratrol on MPTP-induced mouse microglia activation. MAC-1 staining showed that only weak MAC-1 positive microglial immune response was observed in SNpc of the control group. The intraperitoneal injection of MPTP increased the number of MAC-1 positive microglia in SNpc 2.85-fold compared with that of the control group (p<0.001). Resveratrol pretreatment inhibited MPTP-induced activation of microglia (p<0.001, compared to MPTP group) as shown in fig. 4.

We determined the release of striatal dopamine by HPLC. The striatum dopamine level was significantly decreased after MPTP administration. Compared with the MPTP group, the MPTP+ resveratrol group had significantly higher level of dopamine in the striatum (p<0.001) as shown in fig. 5.

Western blotting was used to determine the level of DAT in the striatum. Intraperitoneal injection of MPTP significantly decreased the expression of DAT (p<0.01, compared with control group). Resveratrol was able to reverse the decrease in DAT level (p<0.01, compared with MPTP group) as shown in fig. 6.

Intraperitoneal injection of MPTP significantly reduced the levels of Wnt1, p-Ser9-GSK-3β and β-catenin, but the resveratrol+MPTP group had significantly elevated levels (p<0.01, compared with MPTP group) as shown in fig. 7.

SH-SY5Y cells were pretreated with resveratrol (1 μM, 10 μM or 100 μM) for 24 h before administration of 500 μM MPP+. Cell viability was measured by MTT assay. Compared with the MPP+ group, the MPP+resveratrol group showed significant resveratrol dose dependence. Specifically, 10 μM and 100 μM resveratrol pretreatment significantly augmented cell viability (p<0.01) as shown in fig. 8.

Western blotting exhibited that MPP+ treatment significantly reduced the expression levels of Wnt1, p-Ser9-GSK-3β and β-catenin (p<0.01, compared with control group). In addition, 10 μM resveratrol pretreatment for 30 min significantly elevated their levels (p<0.01, compared with MPP+ group) as shown in fig. 9.
Fig. 3: (A) Astrocytes in mouse SNpc and (B) Stereological cell counts of astrocytes in the mouse SNpc  
Note: Data are represented as mean±SEM of six individual experiments. *Compared with control group, p<0.05 and # compared with MRTP group, p<0.05

Fig. 4: (A) Microglia in mouse SNpc and (B) Stereological cell counts of microglia in the mouse SNpc  
Note: Data are represented as mean±SEM of six individual experiments. *Compared with control group, p<0.05 and # compared with MRTP group, p<0.05
Fig. 5: HPLC of dopamine release in the mouse striatum
Note: Data are represented as mean±SEM of 10 individual experiments. *Compared with control group, p<0.05 and †compared with MRTP group, p<0.05

Fig. 6: Western blot analysis of DAT expression in the striatum of the acute MPTP mouse model
Note: Data are represented as mean±SEM of 10 individual experiments. *Compared with control group, p<0.05 and †compared with MRTP group, p<0.05
Fig. 7: Western blot analysis of Wnt1, (A) p-Ser9-GSK3β; (B) β-catenin and (C) expression levels in mouse SNpc
Note: Data are represented as mean±SEM of five independent experiments. *Compared with control group, p<0.05 and #compared with MRTP group, p<0.05

Fig. 8: Resveratrol dose-dependently suppressed the decrease in SH-SY5Y cell viability induced by MPP⁺ (500 μM)
Note: Data are expressed as mean±SEM of five independent experiments. *Compared with control group, p<0.05 and #compared with MRP⁺ group, p<0.05

Fig. 9: SH-SY5Y cells were pretreated with 10 μM resveratrol in the absence for 30 min and then challenged with MPP⁺ for 24 h. After treatment, cells were harvested and levels of Wnt1, p-Ser9-GSK3 and β-catenin analyzed. Upper panels show densitometric analysis, while the lower panels show representative blots
Note: Data are expressed as mean±SEM of five independent experiments. *Compared with control group, p<0.05 and #compared with MRP⁺ group, p<0.05
Only a weak GFAP-positive astrocyte immune response number of GFAP-positive astrocytes in SNpc of mice. MPTP was further explored. GFAP staining showed the inflammation response and are involved in the regulation of immune system. Astrocytes are involved in the innate immune brain and play a vital role in the central nervous system. Astrocytes are the most abundant cell types in the brain and play a vital role in the central nervous system. Astrocytes are involved in the innate immune response and are involved in the regulation of immune inflammation. Since neuroinflammatory response plays a key role in the pathogenesis of PD, the effect of resveratrol on mouse astrocyte activation induced by MPTP was further explored. GFAP staining showed the number of GFAP-positive astrocytes in SNpc of mice. Only a weak GFAP-positive astrocyte immune response was observed in SNpc of the control mice. Intraperitoneal injection of MPTP increased the number of GFAP-positive astrocytes in SNpc 3.12-fold compared with the control group. Resveratrol pretreatment inhibited MPTP-induced activation of astrocytes (p<0.001, compared with MPTP group). In the brain tissue, microglia can regulate immune inflammation and MPTP can promote the occurrence of neuroinflammation. Herein, the number of MAC-1 positive microglia in SNpc of mice was observed by MAC-1 staining. Intraperitoneal injection of MPTP increased the number of MAC-1 positive microglia in the SNpc area 2.85-fold compared with the control group (p<0.001). Resveratrol pretreatment inhibited MPTP-induced activation of microglia (p<0.001, compared with MPTP group), suggesting that resveratrol significantly suppressed MPTP-induced astrocyte and microglia activation.

To further validate the inhibitory effect of resveratrol on the function of MPTP-damaging dopaminergic neurons, we measured the release of striatal dopamine by HPLC. The dopamine level in the striatum was significantly reduced after MPTP administration in mice. Compared with the MPTP group, the striatum dopamine level was significantly increased in the MPTP+resveratrol group (p<0.001). DAT is a transmembrane protein that reabsorbs dopamine from the synaptic cleft to presynaptic neurons, controls dopamine transport and maintains a balanced dopamine level. To further investigate whether resveratrol plays an important role in the stable expression of DA, the level of DAT in mouse striatum was determined by Western blotting. The results showed that intraperitoneal injection of MPTP in mice can significantly reduce the expression of DAT (p<0.01, compared with control group), resveratrol could reverse the decrease of DAT level induced by MPTP in mice (p<0.01, compared with MPTP group).

Accumulating evidence has proven that the dysregulation of Wnt signaling plays a crucial role in the pathogenesis of PD and Wnt are crucial regulators of inflammatory response. In the brain, both astrocytes and microglia express Wnt receptors and respond to Wnt proinflammatory or anti-inflammatory activity. However, the relationship between the resveratrol anti-inflammatory pathway involved in the damage and repair of nigrostriatal dopaminergic neurons and the Wnt/β-catenin signaling pathway remains unclear. In this study, intraperitoneal injection of MPTP in mice significantly reduced Wnt1, p-Ser9-GSK-3β and β-catenin, and resveratrol+MPTP group Wnt1, p-Ser9-GSK-3β and β-catenin were significantly elevated (p<0.01, compared...
to the MPTP group). To evaluate the direct effects, we investigated the protective effects of resveratrol on dopaminergic neurons in inhibiting MPP+ neurotoxicity. SH-SY5Y was pretreated with resveratrol (1 μM, 10 μM or 100 μM) for 24 h before administration of 500 pM MPP+. Cell viability was measured by MTT assay. Compared with the MPP+ group, the MPP+ + resveratrol group had significant dose dependence on resveratrol, and 10 μM and 100 μM resveratrol pretreatment significantly increased cell viability (p<0.01). We further evaluated the role of the Wnt/β-catenin signaling pathway in the involvement of resveratrol in neuroprotection. Moreover, Western blotting exhibited that MPP+ treatment significantly reduced the expression levels of Wnt1, p-Ser9-GSK-3β and β-catenin (p<0.01, compared with control group) and 10 μM resveratrol pretreatment for 30 min significantly raised the levels (p<0.01, compared with MPP+ group).

In summary, resveratrol relived the neurotoxicity of MPTP against dopaminergic neurons in SNpc and maintained the function of dopaminergic neurons and the reuptake of striatal dopamine. Meanwhile, resveratrol significantly alleviated the toxicity of SH-SY5Y neurons induced by MPP+. Furthermore, the levels of Wnt1, p-Ser9-GSK-3β and β-catenin in SNpc of PD model mice and SH-SY5Y cells decreased, whereas resveratrol significantly elevated such levels. Resveratrol exerted neuroprotective effects on the PD model predominantly by regulating the Wnt/β-catenin signaling pathway.

Author’s contributions:
Yiying Dong and Zhonghai Wang have contributed equally to this work.

Conflict of interests:
The authors declared no conflict of interest.

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