Abstract. The expression levels of micro rna (mir)-340-5p are reportedly decreased in the peripheral blood during acute ischemic stroke; however, the direct effect and mechanism of action of mir-340-5p in ischemic stroke remains largely unknown. The present study aimed to investigate the effects of mir-340-5p, and its mechanism of action, on PC12 cells following oxygen-glucose deprivation/reperfusion (OGD/R) induction. OGD/R-induced PC12 cells served as the cellular model and subsequently, mRNA expression levels of mir-340-5p and neuronal differentiation 4 (Neurod4) were analyzed using reverse transcription-quantitative PCR. Tumor necrosis factor-α, interleukin (IL)-1β and IL-6 expression levels were detected using ELISA kits, and flow cytometry was used to determine the rate of cellular apoptosis. In addition, a nitric oxide (NO) synthase activity assay kit was used to detect NO levels and a NADPH assay kit was used to measure NADPH levels. Western blotting was also performed to analyze protein expression levels of Bax, Bcl-2, cleaved caspase 3 and phosphorylated endothelial NO synthase (eNOS), and the target gene of miR-340-5p was predicted using TargetScan software and verified using a dual-luciferase reporter assay. The expression levels of mir-340-5p were decreased in PC12 cells following OGD/R induction and Neurod4 was identified as a target gene of miR-340-5p. In addition, miR-340-5p overexpression reduced inflammation, apoptotic rate, NO production and NADPH levels, in addition to increasing eNOS expression in PC12 cells following OGD/R induction. Notably, the overexpression of Neurod4 reversed the aforementioned effects of miR-340-5p on PC12 cells following OGD/R induction. In conclusion, the findings of the present study suggested that mir-340-5p may protect PC12 cells against OGD/R through targeting Neurod4, which could provide important implications for the treatment of ischemia-reperfusion injury based on miR-340-5p expression levels in vivo.

Introduction

Ischemic stroke is one of the most frequently occurring conditions among older populations, accounting for a large proportion of the morbidity and mortality rates worldwide (1,2). Smoking habits, hypertension and diabetes have all been listed as risk factors of ischemic stroke (3,4). Ischemic stroke occurs following a blockage in an artery leading to the brain; thus, an insufficient supply of oxygen and glucose reaches the brain that is required for cellular energy, which culminates in irreparable damage (5,6). Currently, there are no effective approved treatments for ischemic stroke (7). Patients are normally treated with recombinant tissue plasminogen activator, undergo surgical excision of the obstruction in the blood vessel or are prescribed protective treatments following stroke, such as fire-needle acupuncture (8). During treatment, the limited reperfusion time window and rapid blood reperfusion cause secondary injuries, namely reperfusion injury, including hemorrhagic transformation and reactive oxygen species (ROS)-induced injuries (9); thus, current treatment regimens are not ideal. Despite significant research being conducted on ischemia-reperfusion injury, little progress has been made. Therefore, it remains an urgent requirement to investigate effective treatment targets and determine the mechanism of action of ischemia-reperfusion injury.

MicroRNAs (miRNAs/miRs) are single-stranded, non-coding RNAs 22 nucleotides in length, which can silence gene expression through transiently promoting translational arrest or inducing the degradation of mRNA (10). Previous studies have reported that miRNAs are involved in the pathophysiology of ischemic stroke through suppressing post-stroke angiogenesis, inhibiting oxidative stress, reducing neuronal loss, suppressing inflammation and preventing excitotoxic injury (11-14).

Notably, miR-340-5p has been observed to relieve chronic constriction injury-induced neuropathic pain and decrease the inflammatory response (15). miR-340-5p has also been demonstrated to suppress hypoxia/reoxygenation-induced apoptosis and oxidative stress in cardiomyocytes (16). Oxidative stress and inflammation are the main causes...
of cerebral ischemia-reperfusion injury, which has been established through numerous previous studies (17-21). For example, in one previous study, miR-340-5p was found to inhibit inflammation and oxidative stress in cardiomyocytes and serve a role in cerebral ischemia-reperfusion injury (16). Furthermore, it was reported that miR-340-5p expression levels were decreased in the peripheral blood following acute ischemic stroke (22), which suggested that miR-340-5p may serve a vital role in ischemic stroke treatment. Based on these previous studies, the present study aimed to investigate the effects of miR-340-5p on a commonly used cell model of ischemia-reperfusion injury, oxygen-glucose deprivation/reperfusion (OGD/R)-induced PC12 cells. The target gene of miR-340-5p was also investigated.

Materials and methods

Cell culture and OGD/R induction. PC12 cells (CRL-1721.1) were obtained from the American Type Culture Collection and were seeded into 96-well plates at a density of 1x10^4 cells/well. Cells were cultured in RPMI-1640 medium (Sigma-Aldrich; Merck KGaA), supplemented with 10% FBS (Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin, and maintained in a humidified atmosphere with 5% CO₂ at 37°C. To study the effects of microRNA-340-5p, the cells were divided into the control group, OGD/R group, mir-NC group and miR-340-5p mimic group. For further study on the underlying mechanism, the cells were divided into an OGD/R group, mir-NC group, miR-340-5p mimic group, miR-340-5p mimic + empty plasmid group and a miR-340-5p mimic + Neurod4 overexpression group. OGD/R was performed in all the groups except for control groups.

OGD/R was performed by culturing the cells in glucose-free medium (RPMI-1640; Gibco; Thermo Fisher Scientific, Inc.) in an incubator with CO₂ (5%; v/v) and N₂ (95%; v/v) at 37°C. After 2 h of incubation, the medium was replaced with normal medium and cells were incubated in an atmosphere of 5% CO₂ and 95% air at 37°C for 12 h.

Cell transfection. Cells were cultured in 6-well plates (1x10^4 cells/well) containing RPMI-1640 medium without antibiotics. Upon reaching 70% confluence, the cells were transfected with a miR-340-5p mimic, miR-negative control (NC), neuronal differentiation 4 (Neurod4) pcDNA3.1 plasmid (10 µl/ml) or an empty pcDNA3.1 plasmid (10 µl/ml) using Lipofectamine® 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.) under serum-free conditions for 6 h. After 48 h of transfection, the cells were collected for subsequent experiments. The miR-340-5p mimic (5'-UUUUAAGC CUGAGACUGAUU-3'), miR-NC (5'-UUUCUGGGAACGU GCCAGGUTT-3'), pcDNA3.1 empty plasmid and Neurod4 overexpression plasmid were all purchased from Shanghai GenePharma Co., Ltd.

Reverse transcription-quantitative PCR (RT-qPCR). Total RNA was extracted from cells using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.). Total RNA was reverse transcribed into cDNA using the qScript miRNA cDNA Synthesis kit (Quantabio), according to the manufacturer's protocol. qPCR was subsequently performed using a TaqMan™ Real Time PCR Mix (Thermo Fisher Scientific, Inc.). The following primer pairs were used for qPCR: GAPDH forward, 5'-AATGGATTGGACGATTGGT-3' and reverse, 5'-TTTGCACTGGTACGTTTGAT-3'; U6 forward, 5'-CTCGTTCTGGGACACA-3' and reverse, 5'-AACGTTTCAAGAATTGG-3'; Neurod4 forward, 5'-AGCTGGTACAACACAACT-3' and reverse, 5'-TTTCAATAAGAGGCGGTCTTC-3'; and miR-340-5p forward, 5'-GCGGTAATAGCAATGAG-3' and reverse, 5'-GGTGCGGTCTGAGAGTCGC-3'. The following thermocycling conditions were used for qPCR: Initial denaturation at 95°C for 8 min; followed by 40 cycles at 95°C for 15 sec and 60°C for 30 sec; and a final extension at 70°C for 35 sec. Expression levels were quantified using the 2-ΔΔCq method (26) and normalized to GAPDH or U6.

Detection of tumor necrosis factor (TNF)-α, interleukin (IL)-1β, monocyte chemoattractant protein-1 (MCP-1) and IL-6. TNF-α (cat. no. BMS662), IL-1β (cat. no. ERIL1B), MCP-1 (cat. no. BMS631INST) and IL-6 (cat. no. BMS625) ELISA kits (Thermo Fisher Scientific, Inc.) were used to detect the levels of TNF-α, IL-1β, MCP-1 and IL-6. Briefly, cells in the different groups were lysed in lysis buffer (Beyotime Institute of Biotechnology) and centrifuged (16,000 x g for 10 min at 4°C) to collect the supernatants. The levels of TNF-α, IL-1β, MCP-1 and IL-6 in the supernatants of each group were analyzed using the corresponding kits, according to the manufacturer's protocols.

Flow cytometric analysis of apoptosis. Flow cytometry was used to analyze the effects of miR-340-5p overexpression on cell apoptosis with an Annexin V-FITC Apoptosis Detection kit (bVbioscience; Thermo Fisher Scientific, Inc.). Cells were grouped into a control group, OGD/R group, mir-NC group and miR-340-5p mimic group. To determine the effects of the target binding between Neurod4 and miR-340-5p on the rate of cell apoptosis, the cells were divided into an OGD/R group, mir-NC group, miR-340-5p mimic group, miR-340-5p mimic + empty plasmid group and a miR-340-5p mimic + Neurod4 overexpression group. OGD/R was performed in all the groups except for control group. Following their respective treatments, cells (1x10⁵ cells/well) in the different groups were washed with PBS and resuspended in the binding buffer. Cells were subsequently incubated with 5 µl Annexin V-FITC and 10 µl propidium iodide staining solution for 15 min at room temperature in the dark. Apoptotic cells were analyzed using a BD FACS Calibur™ flow cytometer (BD Biosciences) and BD CellQuest software (version 5.1; BD Biosciences).

Measurement of nitric oxide (NO)/NADPH levels. The levels of NO and NADPH were analyzed using a Nitric Oxide Synthase Activity assay kit (colorimetric; cat. no. ab211083; Abcam) and NADPH assay kit (colorimetric; cat. no. ab186031; Abcam), respectively. Briefly, cells in the different groups were lysed in lysis buffer (Beyotime Institute of Biotechnology) and collected prior to being analyzed for the levels of NO and NADPH using their corresponding kits, according to the manufacturer's protocols.
Western blotting. Protein expression levels were analyzed using western blotting. Total protein was extracted from cells using lysis buffer (Beyotime Institute of Biotechnology) and lysates were centrifuged (16,000 x g for 10 min at 4°C). Total protein was quantified using a bicinchoninic acid assay kit [Yeasen Biotechnology (Shanghai) Co., Ltd.] and proteins (30 µg/lane) were separated via SDS-PAGE on a 10% gel. The separated proteins were transferred to PVDF membranes (EMD Millipore) and blocked in 5% skimmed milk for 2 h at room temperature. The membranes were incubated at 4°C overnight with the following primary antibodies: anti-Bcl-2 (1:1,000; cat. no. ab196495; Abcam), anti-Bax (1:1,000; cat. no. ab215717; Abcam), anti-Bad (1:1,000; cat. no. ab32445; Abcam), anti-cleaved caspase 3 (1:1,000; cat. no. ab49822; Abcam), anti-caspase 3 (1:1,000; cat. no. ab13847; Abcam), anti-phosphorylated (p)-endothelial NOS (eNOS; 1:500; cat. no. ab215717; Abcam), anti-eNOS (1:500; cat. no. ab199956; Abcam) and anti-GADPH (1:500; cat. no. ab9485; Abcam).

Following the primary antibody incubation, the membranes were incubated with a horseradish peroxidase-conjugated secondary antibody (1:5,000; cat. no. ab7090; Abcam) for 1 h at room temperature. Protein bands were visualized using an ImageQuant™ LAS 500 (GE Healthcare) and an ECL western blotting substrate kit (cat. no. ab65623; Abcam). ImageQuant TL software (version 7.0; Cyiva) was used to perform densitometry.

Dual-luciferase reporter assay. Neurod4 was predicted as a target gene of miR-340-5p-5p using TargetScan software (version 7.2; http://www.targetscan.org/vert_72/). Thus, a dual-luciferase reporter assay was performed to verify the target binding of miRNA-340-5p and Neurod4. The wild-type (WT) 3' untranslated region (3'UTR) binding site of Neurod4 was amplified using PCR and cloned into a pmirGLO reporter plasmid (Promega Corporation). The 3'UTR fragment of Neurod4 was also mutated, resulting in a mutant (MUT) 3'UTR, using the Fast MultiSite Mutagenesis System (Beijing Transgen Biotech Co., Ltd.) and cloned into the pmirGLO reporter plasmid. Cells (5x10^4) cultured in 24-well plates were co-transfected with an equal concentration (450 ng/µl) of Neurod4 (WT or MUT) and miR-340-5p mimic or miR-NC using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocols. After transfection for 48 h, the relative luciferase activity was detected using a Dual-Luciferase Reporter assay system (Promega Corporation), according to the manufacturer's protocols. Luciferase activity was normalized to Renilla luciferase activity.

Statistical analysis. Statistical analysis was performed using GraphPad Prism 5.0 software (GraphPad Software, Inc.) and data from three independent experiments are presented as the mean ± SD. Statistical differences were determined using a one-way ANOVA, followed by Tukey's multiple comparison test. P<0.05 was considered to indicate a statistically significant difference.

Results

miR-340-5p expression levels are decreased in the OGD/R group and miR-340-5p overexpression reduces the OGD/R-induced inflammatory status. The expression levels of miR-340-5p were significantly reduced in the OGD/R group compared with the control group (Fig. 1A), indicating that miR-340-5p may have a certain role in OGD/R-induced cells. The miR-340-5p mimic was successfully transfected into OGD/R-induced cells; significantly increased expression levels of miR-340-5p were observed in the miR-340-5p mimic group compared with the OGD/R and miR-NC groups (Fig. 1B). Subsequently, the effects of miR-340-5p overexpression on the inflammatory status of cells were investigated. Compared with the control group, the levels of TNF-α, IL-1β, MCP-1 and IL-6 were all significantly increased in the OGD/R group (Fig. 1C), which demonstrated that the OGD/R cell model was successfully induced. The OGD/R-induced inflammatory status, which is indicated by the levels of TNF-α, IL-1β, MCP-1 and IL-6, was decreased in the miR-340-5p mimic group when compared with the OGD/R group (Fig. 1C).

miR-340-5p overexpression reduces the cell apoptotic rate induced by OGD/R in PC12 cells. The rate of cell apoptosis in the OGD/R group was significantly increased compared with the control group (Fig. 2A and B), which further confirmed that the OGD/R cell model was successfully induced. However, OGD/R-induced apoptosis was significantly reduced by the miR-340-5p mimic. To further validate these findings, the expression levels of apoptotic proteins were analyzed. The expression levels of the anti-apoptotic protein Bcl-2 were significantly decreased following OGD/R induction compared with the control group, whereas OGD/R-induced decreases in the Bcl-2 expression levels were significantly increased in the miR-340-5p mimic group (Fig. 3). Furthermore, the expression levels of pro-apoptotic proteins, Bax and cleaved caspase 3/caspase 3, were significantly increased in the OGD/R group compared with the control group (Fig. 2A and B), which further confirmed that the OGD/R cell model was successfully induced. The rate of cell apoptosis in the OGD/R group was significantly increased compared with the control group, whereas OGD/R-induced decreases in the Bcl-2 expression levels were significantly increased in the miR-340-5p mimic group (Fig. 3). The expression levels of pro-apoptotic proteins were analyzed using Pcr and cloned into a pmirGlo reporter plasmid. Cells (5x10^4) cultured in 24-well plates were co-transfected with an equal concentration (450 ng/µl) of Neurod4 (WT or MUT) and miR-340-5p mimic or miR-NC using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocols. After transfection for 48 h, the relative luciferase activity was detected using a Dual-Luciferase Reporter assay system (Promega Corporation), according to the manufacturer's protocols. Luciferase activity was normalized to Renilla luciferase activity.

miR-340-5p overexpression inhibits NO production, reduces NADPH levels and increases the relative expression levels of p-eNOS in OGD/R-induced PC12 cells. The relative expression levels of p-eNOS were determined as the ratio of p-eNOS/eNOS/GADPH. NO, which is induced by OGD/R in PC12 cells, was evaluated herein. The NO levels were significantly increased in the OGD/R group compared with the control group, whereas these OGD/R-induced increases were significantly reduced in the miR-340-5p mimic group (Fig. 4A). Furthermore, p-eNOS/eNOS was significantly decreased in the OGD/R group compared with the control group, and this effect was partly reversed in the miR-340-5p mimic group (Fig. 4B). All these findings suggested that miR-340-5p overexpression may inhibit cell apoptosis through increasing the expression levels of Bcl-2, and decreasing those of Bax, cleaved caspase 3 and caspase 3.

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Neurod4 is a target of miR-340-5p in OGD/R-induced PC12 cells. It was predicted by TargetScan that Neurod4 was a target gene of miR-340-5p (Fig. 5A), thus a dual-luciferase reporter assay was used for further validation. The relative luciferase activity was significantly reduced in the 3'UTR-WT-Neurod4 + miR-340-5p mimic group compared with the 3'UTR-MUT-Neurod4 + miR-340-5p mimic group (Fig. 5B), indicating that miR-340-5p may target Neurod4 in OGD/R-induced PC12 cells. Furthermore, Neurod4 expression levels were significantly increased in the OGD/R group compared with the control group, and this effect was significantly reversed following the addition of the miR-340-5p mimic (Fig. 5C). This observation further validated that Neurod4 may be a target of miR-340-5p in OGD/R-induced PC12 cells.

miR-340-5p overexpression reduces the inflammatory status, whereas Neurod4 overexpression counteracts the effects of miR-340-5p overexpression on OGD/R-induced PC12 cells.

Neurod4 expression levels were significantly increased in the Neurod4 overexpression group compared with the empty plasmid group (Fig. 6A), indicating that Neurod4 overexpression was successful in the PC12 cells. The OGD/R-induced levels of TNF-α, IL-1β, MCP-1 and IL-6 were significantly reduced following miR-340-5p overexpression in the OGD/R + miR-340-5p mimic group (Fig. 6B-E). However, following Neurod4 overexpression, the anti-inflammatory effects of miR-340-5p overexpression on OGD/R-induced PC12 cells were reversed and the inflammatory levels were significantly increased. These findings indicated that the anti-inflammatory effects of miR-340-5p overexpression on OGD/R-induced PC12 cells may be achieved by decreasing the levels of TNF-α, IL-1β, MCP-1 and IL-6 through down-regulating Neurod4 expression.

miR-340-5p overexpression reduces apoptosis and Neurod4 overexpression counteracts the effects of miR-340-5p
The OGD/R-induced increases in cell apoptosis were significantly reduced following mir-340-5p overexpression (Fig. 7a and B); however, the apoptotic rate in OGD/R-induced cells transfected with the miR-340-5p mimic was significantly increased following Neurod4 overexpression. These findings suggested that Neurod4 overexpression may reverse the effects of miR-340-5p overexpression on cell apoptosis in OGD/R-induced PC12 cells. As an
anti-apoptotic protein, Bcl-2 expression levels were observed to be significantly increased in the OGD/R + miR-340-5p mimic group; however, this effect was reversed following Neurod4 overexpression (Fig. 7C). In addition, the expression levels of the pro-apoptotic proteins Bax, Bad, cleaved caspase 3 and caspase 3 demonstrated the opposite trend compared with the Bcl-2 expression levels in each group. Taken together, these findings suggested that miR-340-5p overexpression may protect OGD/R-induced cells from apoptosis through increasing the expression levels of Bcl-2, and decreasing the expression levels of Bax, Bad, cleaved caspase 3 and caspase 3 through decreasing Neurod4 expression.
miR-340-5p overexpression protects OGD/R-induced PC12 cells by reducing NO levels and increasing p-eNOS/eNOS expression levels, whereas this effect is counteracted by Neurod4 overexpression. The OGD/R-induced increases in NO levels were significantly decreased in the OGD/R + miR-340-5p mimic group (Fig. 8A). However, the effect of miR-340-5p overexpression was significantly reversed following Neurod4 overexpression, suggesting that miR-340-5p overexpression may exert a protective effect by decreasing NO levels (Fig. 8A). The expression levels of p-eNOS were significantly increased in the OGD/R + miR-340-5p mimic group compared with the OGD/R group, but this effect was reduced following Neurod4 overexpression (Fig. 8B), indicating that miR-340-5p overexpression may protect OGD/R-induced PC12 cells from injuries by decreasing NO levels and the expression levels of p-eNOS/eNOS through downregulating Neurod4 expression.

**Discussion**

Ischemic stroke remains one of the major causes of morbidity; however, how to treat the condition remains largely unknown. The current therapeutic options for ischemic stroke are unsatisfactory; therefore, there is an urgent requirement to further study the underlying mechanisms to identify novel treatment options for ischemic stroke (28,29). In the present study, miR-340-5p was observed to exert protective effects over OGD/R-induced PC12 cells through targeting Neurod4 expression.
Figure 7. Neurod4 overexpression counteracts the effect of miR-340-5p overexpression on the apoptotic rate in OGD/R-induced PC12 cells. (A) Apoptotic rate was determined in the different groups. (B) Semi-quantification of (A). (C) Expression levels of Bax, Bcl-2, cleaved caspase and cleaved caspase 3 in different groups. *P<0.05, **P<0.01, ***P<0.001 vs. OGD/R group; ###P<0.001 vs. OGD/R + miR-nc group; ΔP<0.05, ΔΔP<0.01, ΔΔΔP<0.001 vs. miR-340-5p mimic + empty plasmid group. miR, microRNA; NC, negative control; Neurod4, neuronal differentiation 4; OGD/R, oxygen-glucose deprivation/reperfusion.
The insufficient presence of oxygen and glucose for normal metabolism is the main cause of injury in stroke (30) and as the therapeutic window is limited, early recanalization has proved helpful in preventing ischemic neuronal loss (31). Thrombolytic therapies are commonly used for the treatment of ischemic stroke (32); however, the recovery of blood flow has been found to promote secondary injuries (33). Thus, reducing reperfusion injuries is of great significance for improving the therapy available for ischemic stroke.

In the present study, PC12 cells that received oGd/r served as the cellular model, which has been used in numerous previous studies (24,25,34,35). Consistent with the previous studies, the cell apoptotic rate, the inflammatory status and NO levels were increased following oGd/r induction, indicating that the cellular model was successfully established (36,37).

In a previous study, the expression levels of miR-340-5p were rapidly decreased in the peripheral blood of patients who had suffered from an acute ischemic stroke, which indicated a potentially protective role for miR-340-5p in ischemic stroke (22). Consistent with this previous study, miR-340-5p expression levels were decreased in the OGD/R group in the present study.

Cell apoptosis and inflammation are two important factors involved in ischemia-reperfusion injury (38-40). In the current study, following the overexpression of miR-340-5p in PC12 cells, the cell apoptotic rate, inflammatory status and NO levels induced by OGD/R were reduced, providing validation for the protective role of miR-340-5p in cells induced by OGD/R.

Neurod4 is an important factor involved in neuronal differentiation; its expression levels have been reported to be increased under various stimuli, which was negatively correlated with the degree of neuron maturation (41). In the present study, Neurod4 was identified as a target gene of miR‑340‑5p. Neurod4 expression levels have also previously been reported to be increased following maternal hypoxia (42). In the present study, Neurod4 expression levels were increased in the OGD/R group compared with control group, and the OGD/R-induced
increases in Neurod4 expression levels were reduced following miR-340-5p overexpression. To the best of our knowledge, the latter finding was reported for the first time in the present study, which suggested that miR-340-5p may protect PC12 cells from OGD/R injury through targeting Neurod4 expression. In addition, in a previous study, the inhibition of Neurod4 expression reduced the inflammatory levels and suppressed oxidative stress in spinal cord injury (41,43). These injuries caused by inflammation and oxidative stress are suggested to be the two major factors involved in ischemia-reperfusion injury (40,44-46).

In the present study, increased levels of no decreased, which subsequently resulted in ischemia-reperfusion with mir-340-5p may provide a novel strategy for treating ischemia-reperfusion injury. However, future studies are required to determine other targets that could be targeted for the treatment of ischemic stroke.

In a cerebral ischemia-reperfusion injury model, the levels of NO, a physiological messenger, were reported to be upregulated and the expression levels of p-eNOS/eNOS were decreased, which subsequently resulted in ischemia-reperfusion injuries (47). In the present study, increased levels of NO induced by OGD/R were subsequently reduced following miR-340-5p overexpression. The opposite trend was observed to occur to the expression levels of p-eNOS/eNOS following miR-340-5p overexpression. Notably, the effects of miR-340-5p overexpression on NO levels and p-eNOS/eNOS expression levels were reversed by Neurod4 overexpression. These findings further indicated that miR-340-5p may protect against injuries from OGD/R through inhibiting the production of NO and increasing the expression levels of p-eNOS/eNOS through targeting Neurod4.

In conclusion, the findings of the present study suggested that miR-340-5p may exert protective effects over OGD/R in PC12 cells through targeting Neurod4 expression. These results may provide novel strategies for alleviating the injuries obtained from ischemia-reperfusion and pave the way for future research on ischemia-reperfusion injury.

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Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

JW designed the present study, prepared the manuscript and was involved in performing the experiments. GL performed some parts of the experiments and helped to design the study. Both authors read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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