Nrf2-regulated miR-380-3p Blocks the Translation of Sp3 Protein and Its Mediation of Paraquat-Induced Toxicity in Mouse Neuroblastoma N2a Cells

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ABSTRACT

Laboratorial and epidemiological research has established a relationship between paraquat (PQ) exposure and a risk for Parkinson’s disease. Previously, we have investigated the effects of nuclear factor erythroid 2 related factor 2 (Nrf2) and microRNAs in PQ-induced neurotoxicity, addressing the function of miR-380-3p, a microRNA dysregulated by PQ, as well as Nrf2 deficiency. Nrf2 is known to mediate the expression of a variety of genes, including noncoding genes. By chromatin immunoprecipitation, we identified the relationship between Nrf2 and miR-380-3p in transcriptional regulation. qRT-PCR, Western blots, and dual-luciferase reporter gene assay showed that miR-380-3p blocked the translation of the transcription factor specificity protein-3 (Sp3) in the absence of degradation of Sp3 mRNA. Results based on cell counting analysis, annexin v-fluorescein isothiocyanate/propidium iodide double-staining assay, and propidium iodide staining showed that overexpression of miR-380-3p inhibited cell proliferation, increased the apoptotic rate, induced cell cycle arrest, and intensified the toxicity of PQ in mouse neuroblastoma (N2a [Neuro2a]) cells. Knockdown of Sp3 inhibited cell proliferation and eclipsed the alterations induced by miR-380-3p in cell proliferation. Two mediators of apoptosis and cell cycle identified in previous studies as Sp3-regulated, namely cyclin-dependent kinase inhibitor 1 (p21) and calmodulin (CaM), were dysregulated by PQ, but not Sp3 deficiency. In conclusion, Nrf2-regulated miR-380-3p inhibited cell proliferation and enhanced the PQ-induced toxicity in N2a cells potentially by blocking the translation Sp3 mRNA. We conclude that CaM and p21 were involved in PQ-induced toxicity.

Key words: paraquat; neurotoxicity; Nrf2; miR-380-3p; Sp3, p21; CaM.
Paraquat (PQ) has been used worldwide for more than half a century as an efficient agrichemical, yet its application remains highly controversial given its documented toxicity. PQ causes a widespread oxidative/nitrosative stress in cells, after 1-electron reduction chemistry. The dopaminergic neurons readily take up PQ via the dopamine transporter and the organic cation transporter-3 (Barlow et al., 2003; Rappold et al., 2011), and are highly sensitive to PQ-induced oxidative stress (Dröge and Schipper, 2007; Fitsanakis et al., 2002), resulting in loss of dopaminergic neurons in the substantia nigra, one of the cardinal clinical features of Parkinson’s disease (PD). Neurodegeneration models established for PQ toxicity both in vivo and in vitro have thus been widely employed to address mechanisms associated with the etiology of PD (Berry et al., 2010).

PD is a common complex syndrome ascribed to complicated pathogenesis, encompassing motor deficits, static tremors, bradykinesia, rigidity, and postural/gait disturbance, as well as other nonmotor neurological deficits to various degrees, which are generally considered to be manifestation of loss in dopaminergic neurons. The etiology of PD involves both genetic and environmental factors. Genes, such as alpha-synuclein and parkin among various species. The expression of miR-380-3p was found restricted to embryos and brain in adults (Seitz et al., 2010; Schipper, 2007; Fitsanakis et al., 2005). Nuclear factor erythroid 2 related factor 2 (Nrf2) regulates the expression of numerous antioxidant proteins, such as superoxide dismutase 1 and 2 (SOD1, SOD2), glutathione peroxidase 1 and 5 (GPX-1, GPX-5), and NAPD(P)H quinine oxidoreductase (NQO1) (De Long et al., 1986; Johnson and Giulivi, 2005; Liddell et al., 2006; Siegel and Ross, 2000; Song et al., 2014). In our previous study, Nrf2 was found to be protective in PQ-induced neurotoxicity (Li et al., 2012), establishing a critical role for oxidative stress in mediating PQ neurotoxicity.

MicroRNAs (miRNAs) are small noncoding regulatory RNAs of 18–25 nucleotides in length that regulate gene expression at posttranscriptional level by sequence-specifically binding to the 3′-untranslated (3′UTR) region of the target mRNAs to inhibit the translation or induce the degradation of these mRNAs. miRNAs are involved in survival and differentiation of dopaminergic neurons and are relevant to the pathogenesis and progression of PD (Anderegg et al., 2013; de Mena et al., 2010; Hébert and De Strooper, 2007; Goodall et al., 2013). Apart from the direct association with PD etiology, miRNAs are also closely related to Nrf2: various interactions between the transcription factor Nrf2 and miRNAs have been revealed. For example, miR-200a and miR-141 target the mRNA of Kelch-like ECH-associated protein 1 (Keap1), a master regulator of Nrf2 by deleteriously binding to it, reducing the cytoplasmic abundance of Keap1 protein, thus releasing Nrf2 proteins from sequestration and further degradation (van Jaarsveld et al., 2013; Yang et al., 2014). Furthermore, some miRNAs, such as miR-146a, miR-132, and miR-28, can directly bind to Nrf2 mRNA and repress its translation (Smith et al., 2015; Stachurska et al., 2013; Yang et al., 2011).

Previous studies have revealed 3 mechanisms for the regulation of the transcription of miRs. (1) Methylation level of the upstream Cpg rich region of miRs was found to be inversely related to miR expression (Fiaschetti et al., 2014). (2) Nucleotide polymorphism occurs in the promoter region of miRs and induces a variety in their expression level (Zhang et al., 2014). (3) The activity of the transcription factor bound to the promoter region of miRs likely plays another determinant role in miR expression (Liang et al., 2009). The transcription factor Nrf2 has been shown to regulate the expression of many miRNAs, such as miR-29 and miR-1 (Kurinna et al., 2014; Shah et al., 2014; Singh et al., 2013). Accordingly, it is reasonable to suggest that Nrf2 might regulate the expression of certain miRNAs, concomitant with its well-documented antioxidant effect. In the present study, Nrf2 was shown to promote the expression of miR-380-3p whereas PQ inhibiting the same miR. miR-380-3p (5′-uauagugauuggaccauc-3′) is encoded within chromosome 12 (105711803–109711863, precursor) in the mouse, and is relatively conserved among various species. The expression of miR-380-3p was found restricted to embryos and brain in adults (Seitz et al., 2004), implying a special role that miR-380-3p may have in neuronal function, differentiation and survival.

Previously, a set of miRNAs in mouse substantia nigra was found dysregulated by PQ (Wang et al., 2017). Because the potency of miRNAs has been widely recognized, altered miRNA expression profile is likely to exert significant health effects. In the present study, we addressed one of these dysregulated miRNAs, miR-380-3p, and its function in PQ-induced nerve cell injury. To decipher the mechanism underlying the function miR-380-3p, apart from its direct target, namely specificity protein-3 (Sp3), some genes potentially regulated by Sp3 (Gartel et al., 2000; Pan et al., 2000; Sowa et al., 1999) were also studied. Specifically, we examined the expression of 2 ubiquitously active molecules, namely cyclin-dependent kinase inhibitor 1 (p21) and calmodulin (CaM). Furthermore, we explored its interaction with Nrf2 and the regulatory role Nrf2 exerts over miR-380-3p. This research was predominantly carried out in Neuro2a (N2a) cells, a mouse neuroblastoma cell line that has a long history of serving in studies on dopamine-associated neurotoxicity (LePage et al., 2005).

MATERIALS AND METHODS

Quantitative RT-PCR for validation of miRNA and mRNA expression. Total RNA samples were extracted from N2a cells and reversely transcribed to cDNA using Prime Script RT reagent kit RR037a for the miRNAs, and kit RR047a for the long RNAs (TAKARA, Japan), following the manufacturer’s instructions. The reverse transcription of miRNAs entailed specific stem-loop primers obtained from Ribobio (Guangzhou, China). The subsequent real-time PCR mixture was established with SYBRPremix Ex Taq RR820a (TAKARA), following the manufacturer’s instructions, and performed in 7500 real-time PCR system (ABI) or Light cycler 480 (Roche, Switzerland). The reaction tubes were incubated at 95°C for 30 s, followed by 40 cycles at an interval of 5 s at 96°C and an interval of 34 s at 60°C. For melting curve analysis, the amplifications were incubated at 95°C for 15 s, 60°C for 1 min and 95°C for 15 s. Data were analyzed by 2ΔΔCT. The expression level of miRNA was normalized to U6, and long RNAs were normalized to GAPDH.

Dual-luciferase assay. Luciferase assays were carried out to confirm the interaction between miR-380-3p and the 3′UTR of Sp3 mRNA. Two hundred and ninety-three T cells, which are frequently used in dual-luciferase reporter assay and which do not
Table 1. The Sequences Containing Wild-type or Mutant Binding Site (Red) of miR-380-3p, Obtained From the 3’-Untranslated (3’UTR) of Specificity Protein-3 (Sp3) mRNA

| ID | Sequences (5’-3’) |
|----|------------------|
| mmu-Sp3 mRNA 3’UTR (WT) | ATTATGCATAAATCGCAAAATCAGTTTCCAAGGAAATTTGCAATGCTAGGCTTTACTGTGAACAGAGCTAATTATTTTAATTAGGTTAACTGTACAATATTTTAAGCATTACTCTCTTGTAAGATTTGAGAAAAACCTGAGCTTAAAGCATGTTATCAGTTTTGAT%
| mmu-Sp3 mRNA 3’UTR (mutant) | ATTATGCATAAATCGCAAAATCAGTTTCCAAGGAAATTTGCAATGCTAGGCTTTACTGTGAACAGAGCTAATTATTTTAATTAGGTTAACTGTACAATATTTTAAGCATTACTCTCTTGTAAGATTTGAGAAAAACCTGAGCTTAAAGCATGTTATCAGTTTTGAT%

Chromatin immunoprecipitation assay. N2a cells were cultured as described above. Prior to cross-linking, a group of cells were treated with 100 μM PQ for 48 h. Cross-linking was performed with formaldehyde at a final concentration of 1% (wt/wt) and terminated after 5 min by addition of glycine at a final concentration of 0.125 M. Cells were harvested with lysis buffers in chromatin immunoprecipitation (ChIP) kit (BersinBio, Guangzhou, China) according to the manufacturer’s protocol. Chromatin was sonicated with UC3000 (Diagenode, Belgium) to generate DNA fragments with an average size of 500 bp. Then 4 h incubation with monoclonal Nrf2 (CST) or IgG (Millipore) antibody-beads was performed with an average size of 500 bp. Then 4 h incubation with monoclonal Nrf2 (CST) or IgG (Millipore) antibody-beads was performed in vertical mixer at 4 °C. Next, washing and elution were performed (washing and elution buffer, respectively) with a ChIP kit, according to the manufacturer’s instructions. After reversion cross-linking, the DNA was collected by centrifugations and dissolved in ddH2O. Immunoprecipitated DNA was analyzed by qPCR and the enrichment was expressed as percentage of the input for each condition. The sequences of primers used in the following PCR were listed in Table 2.

Western blotting. Protein level of Sp3 in N2a cells was assessed by Western blot and normalized to GAPDH. Total protein samples were extracted from N2a cells with lysis buffer. Protein concentrations were determined by BCA protein assay kit, standardized by bovine serum albumin. Equal amounts of protein were loaded in each lane for sodium dodecyl sulfate polyacrylamide gel electrophoresis (5% polyacrylamide for compression and 10% for separation, wt/wt). Next, the separated proteins were transferred to nitrocellulose membranes. The membranes were blocked with 5% (wt/wt) skimmed milk for 1 h before the overnight incubation with primary antibodies in 5% (wt/wt) skimmed milk (anti-SP3, 1:200, Santa Cruz; anti-BCL2, 1:1000, Abcam; anti-BAX, 1:2000, Abcam; anti-GAPDH, 1:10000, Abcam) at 4 °C. Membranes were rinsed with TBST for 6 times with 10 min after the cycles. To ensure amplification efficiency, BSP products were analyzed by 3% (wt/wt) agarose gel electrophoresis. Ten colonies were collected for each group and then the sequencing of bacteria solutions was entrusted to Sangon Biotech (Shanghai).

Table 2. The Sequences of Primers Used in the qPCR Following Chromatin Immunoprecipitation

| Gene | No. | F/R | Primer Sequence | Site | Product (bp) |
|------|-----|-----|----------------|------|--------------|
| mir379 | 1 | F | CAACCCCTGTCTCTCTGCTC | 323 | 238 |
| | | R | GAACAGAGGGAAGACAGAG | | |
| mir379 | 2 | F | GCCAGCCAGTGTGTGAATT | 1271 | 227 |
| | | R | GAGGTTCCTGGGTATGAGCA | | |
| Beta-actin | 3 | F | AACAGCGGAGCAGTCACACT | 213 | |
| | | R | GGCTCAGCACCATGAGTCAG | | |

Bisulfite sequencing PCR. The primer sets of bisulfite sequencing PCR (BSP) were used to target the CpG: 29 located between Dlk1 and Glt2 as shown in Table 3. Extracted DNA samples from N2a cells were converted using a Bisulfite DNA Modification Kit (BersinBio), following the manufacturer’s instructions. PCR amplification with BSP primers was then predenaturated at 95 °C for 5 min, followed by 30 cycles at an interval of 10 s at 95 °C, an interval of 15 s at 58 °C and an interval of 15 s at 72 °C, and 72 °C for 10 min after the cycles. To ensure amplification efficiency, BSP products were analyzed by 3% (wt/wt) agarose gel electrophoresis.

Cell culture. The murine neuroblastoma cell line N2a was purchased from Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China), and maintained in Dulbecco’s Modified Eagle’s Medium (DMEM, HyClone, Utah) containing 10% (wt/wt) fetal bovine serum (HyClone) and 1% (wt/wt) penicillin and streptomycin (Gibco, Massachusetts) at 37 °C in a humidified atmosphere containing 5% (vol/vol) CO₂. For resuspension, the cells were trypsinized by Trypstin-EDTA (Gibco), and the cells were passaged when attaining 80% confluence.

Cell treatments. For PQ treatment, the cells were maintained in complete medium containing 50, 100, and 200 μM PQ, alone or following a 6 h pretreatment with 40 μM BHQ. Cells without any treatment were used as control. The N2a cells were transfected with miRIDIAN mir-380-3p mimics, hairpin inhibitors (Dharmacon, Colorado) or siRNAs (GenePharma, Shanghai, China) using DharmaFECT reagent (Dharmacon), and the corresponding scrambled sequences were also transfected in cells as
negative controls. The transfections were carried out in Reduced Serum Media (Opti-MEM, Gibco) and limited to 6 h to reduce cytotoxicity. The transfection of siRNA and corresponding scrambled sequences was after the same protocol. The mimics and siRNAs were applied at 40 nM, and inhibitors at 100 nM, under conditions analogous to the corresponding negative controls, respectively.

Cell proliferation assay. Cell proliferation was measured using Cell Counting Kit-8 (CCK-8) reagent (Solarbio, Beijing, China) according to the manufacturer’s protocol. The N2a cells were seeded into 96-well plates (5 × 10^4 cells per well) and incubated. Upon reaching 40% confluency, the cells were transfected with miRIDIAN miR-380-3p mimic or hairpin inhibitor. The corresponding scrambled sequences were adopted in negative control groups. The transfected cells were then exposed to 0, 50, 100, or 200 μM of PQ for 48 h, followed by cell proliferation assessment. A mixture of 90 μl medium and 10 μl CCK-8 solution was used to replace the PQ-stained medium in each well, incubated for 2 h and the absorbance values were measured at 450 nm using a microplate reader.

Cell cycle analysis. Cells were trypsinized, harvested, washed twice with cold PBS, fixed with 75% (wt/wt) ice-cold ethanol at −20°C overnight, washed twice with cold PBS, and then incubated with PI/RNase Staining Buffer (BD Biosciences, New Jersey) for 15 min at room temperature. The tubes were stored at 4°C protected from light prior to analyzing and the PI signal was examined by a flow cytometry (FACSVerses, BD Biosciences) within 1 h.

Apoptosis assay. Apoptosis was quantified using FITC Annexin V Apoptosis Detection Kit 1 (BD Biosciences) following the manufacturer’s protocol, and with flow cytometry (FACSVerses, BD Biosciences). Cells were trypsinized and collected 48 h after transfection, washed twice with cold PBS, and resuspended at 1 × 10^6 cells/ml. Hundred microliters of the cell solution was transferred to a new tube and mixed with 5 μl of FITC Annexin V and 5 μl of PI, gently vortexed. After incubation for15 min in the dark, 400 μl of 1 × Binding Buffer was added to each tube, and the cells were analyzed by flow cytometry within 1 h.

Statistical analysis. The results were analyzed by SPSS 22 and expressed as mean ± SD. One-way analysis of variance (ANOVA) and Student-Newman-Keuls test for pairwise comparisons were used to detect significant differences among the groups under varying experimental treatments. The differences between 2 groups were determined by t test or Welch’s t test when the variances were unequal. The interactions between factors were determined by two-way ANOVA. p < .05 was the criterion for statistical significance.

RESULTS

Nrf2 Transcriptionally Regulates miR-380-3p Expression in N2a Murine Neuroblastoma Cell Line

Nrf2 is sequestered by Keap1, anchored to the cytoplasm and degraded by Cullin 3, and seldom been activated in the absence of particular triggers, such as oxidative stress [Itoh et al., 1999]. Herein, tert-Butylhydroquinone (tBHQ), a specific activator of Nrf2 (Li et al., 2005, 2012, 2014), was applied to N2a cells at 40 μM alone or followed by 100 μM PQ. The former releases Nrf2 from the sequestration of Keap1, and thus enhances the transcriptional function of Nrf2. The NAD(P)H dehydrogenase [quinone] 1 (NQO1) mRNA level, which was already verified as transcriptionally regulated by Nrf2 [Dinkova-Kostova and Talalay, 2010], was increased by treatment with tBHQ (Figure 1A, p < .0001 or p < .001, respectively). The miR-380-3p level increased along with the tBHQ-induced activation of Nrf2 (Figure 1B, p < .001 or p < .0001, respectively).

To validate the hypothesis that Nrf2 transcriptionally regulates miR-380-3p, we obtained the binding site motif of mmu-Nrf2 (Figure 1C) from the transcriptional factor encyclopedia, a database established by the center for molecular medicine and therapeutics (CMMT). Many eukaryotic miRs, just like miR-380, are encoded in clusters resembling the operons common in prokaryotes and all miRs within 1 cluster share the upstream promoter region. The 2000 bp upstream of miR-379, the first miR in the cluster according to genome mapping, should contain the potential promoters that regulate the expression of miR-380-3p and thus was searched for possible binding sites. By motif-based analysis, an -ATTTTTCTGAGTTAG- (1780–1794) was predicted as a possible Nrf2 binding site within the 2000 bp sequence.

Next, we adopted ChIP to confirm the prediction. Using ChIP followed by qPCR covering 2 properly selected sites in the potential promoter region, the combination of Nrf2 and the promoter region of miR-380-3p (Figs. 1D and 1E, p < .01 or p < .001, respectively) was identified.

miR-380-3p Suppresses the Translation of Sp3 mRNA

A location pairing the seed sequence of miR-380-3p was found in the 3’UTR region of Sp3 mRNA (+tgtagacctgattaagaa, 3117–3123), indicating a canonical match likely to regulate the translation of Sp3 mRNA (Wang, 2014). Accordingly, Sp3 mRNA and protein levels after alterations in miR-380-3p expression were determined as to test our prediction.

Though miR-induced mRNA degradation mostly occurs in plants, it has also been reported in vertebrates. qRT-PCR was performed to measure the Sp3 mRNA level in N2a cells after transfection of miR-380-3p mimics/inhibitors. The Sp3 mRNA abundance was indistinguishable from control after the transient transfections (Figs. 2A–D, p > .05), whereas its protein level was changed in response to alterations in miR-380-3p expression (Figs. 2E and 2F, p < .05).

However, the changes in Sp3 protein level were insufficient to properly validate the regulation or to locate the exact binding site in the 3’UTR of Sp3 mRNA. Accordingly, we cotransfected the dual-luciferase reporter (psiCHECKTM-2 Vector) carrying the predicted binding site (Table 1), along with miR-380-3p mimics or the corresponding negative control sequence, into 293 T cells. We found that miR-380-3p mimics suppressed the translation of Renilla luciferase (Figure 2G, p < .0001). Furthermore, mutation (Table 1) in the predicted binding site abolished the suppression by the mimics (Figure 2H, p > .05).
Overexpression of miR-380-3p Enhances the Toxicity of PQ in N2a Cells

By transient transfection of miR-380-3p mimics/inhibitors, synthetics resembling or counteracting mature miRNAs, we achieved immediate functional gain/loss of miR-380-3p (Figs. 3A–C, p < .05 or p < .01, respectively), and observed the ensuing effects. Cell proliferation, apoptosis rate, and cell cycle distribution assays were carried out, to establish the role of miR-380-3p in nerve cell survival and proliferation.

The miR-380-3p mimics inhibited cell proliferation (Figure 3D, p < .001, p < .05, respectively) and increased apoptosis rate of N2a cells (Figure 3F, p < .001), altering the cell phase proportion, reducing the mitotic cells and keeping more cells in S phase (Figure 3G, p < .01, respectively). However, the inhibition of miR-380-3p did not cause detectable alterations in any of the above endpoints (Figs. 3E–G, p > .05).

Upon exposure to PQ, miR-380-3p mimics caused lower cell proliferation, higher apoptosis, and cycle arrest in N2a cells (Figs. 3D, 3H, and 3I, p < .0001, p < .01, p < .05, respectively), whereas miR-380-3p inhibition failed to induce any detectable alterations (Figs. 3E, 3H, and 3I, p > .05).

PQ Reduces the Expression of miR-380-3p in N2a Cells

PQ reduced the expression of miR-380-3p in N2a cells (Figure 4A, p < .0001 or p < .01, respectively), and longer exposure or higher concentration intensified this effect. Furthermore, the expression of NQO1 was also reduced by PQ when Nrf2 was already activated by BHQ (Figure 4B, p < .001) and the binding between Nrf2 and the miR-380-3p promoter region was also slightly weakened by PQ, though the difference did not attain statistical significance (Figure 4C, p > .05).

The methylation level of an intergenic germ-line derived differentially methylated region (IG-DMR) between Dlk1 and Gtl2 was previously found to be relevant to the regulation of the expression of miR-380-3p and other miRNAs within the cluster (Lin et al., 2003; Seitz et al., 2004). To further determine the mechanism by which PQ influences the miR-380-3p expression, we examined whether PQ changes the methylation level of the IG-DMR. But exposure to PQ did not induce substantial alterations in the methylation of IG-DMR (Figs. 4D and 4E).

PQ Dysregulates Sp3 Expression in N2a Cells

Among all the potential targets of miR-380-3p, Sp3 had been recognized as functional in many systems (Gandhy et al., 2015; Gölner et al., 2001; Krüger et al., 2007; van Loo et al., 2007). Thus, we investigated the involvement of Sp3 in PQ-induced neurotoxicity. First, PQ was found to cause disordered Sp3 expression. At the early stage of exposure or at lower concentration, the Sp3 mRNA and protein level in N2a cells were elevated in response to PQ exposure, but subsequently reduced to levels lower than the untreated group as the treatment proceeded both in time or concentration (Figs. 5A and 5B, p < .001, or p < .0001, respectively).

Next, we determined whether Sp3 had an effect on PQ-induced neurotoxicity. Several siRNAs targeting Sp3 mRNA were designed (Table 4) and the knockdown efficiency was examined (Figs. 5C and 5D, p < .05, p < .01, or p < .001, respectively). After specific knockdown of Sp3, the proliferation of N2a cells, compared to the groups transfected with negative control sequence, was found generally lower (Figure 5E, p < .0001), analogous to the effect of miR-380-3p overexpression.

Because miR-380-3p targets Sp3 mRNA and exerts an effect similar to Sp3 siRNA, it is probable that the effect of miR-380-3p occurs predominantly by regulating Sp3. To assess if Sp3 deficiency would abolish the effects of miR-380-3p and thus to determine whether miR-380-3p was mediated via Sp3, we...
knocked down the Sp3 mRNA prior to the application of miR-380-3p mimics or inhibitors and examined cell proliferation. Knockdown of Sp3 seemed to narrow the gap between cells transfected with mimics and the corresponding negative control groups (Figure 5F, \( p < .001 \), \( p < .0001 \), or \( p > .05 \), respectively). However, cell proliferation was not altered by miR-380-3p inhibition (Figure 5G, \( p > .05 \)).

CaM and p21 Are Dysregulated by PQ

As well as being regulated by Sp3 (Pan et al., 2000; Sowa et al., 1999), p21 is a potent cyclin-dependent kinase inhibitor and a key regulator in cell cycle progression (Gartel and Radhakrishnan, 2005) and apoptosis (Almond and Cohen, 2002), whereas CaM plays a role in many crucial cellular processes, such as metabolism, inflammation, apoptosis, short- and long-term memory, and immune response (Chin and Means, 2000; Lledo et al., 1995; Nishizawa et al., 1988). In this study decreased expression of Sp3, as well as lower concentrations of PQ (50 \( \mu M \)), failed to significantly alter the expression of CaM or p21 (Figure 6A, \( p > .05 \)). However, the p21 mRNA abundance was Sp3-independently reduced by 100 \( \mu M \) PQ (Figure 6B, \( p < .05 \) or \( p < .01 \), respectively), whereas the CaM mRNA level increased...
following exposure to 100 μM PQ (Figure 6C, p < .01). Sp3 knockdown did not alter CaM and p21 expression levels.

To explore other possible pathways through which Sp3 may cast its effects, protein levels of 2 critical apoptosis mediators, Bcl2 and Bax (Cimmino et al., 2005; Czabotar et al., 2009), were examined. The expression of Bax or Bcl2 was not significantly altered by either PQ treatment or Sp3 interference (Figs. 6D and 6E, p > .05).

**DISCUSSION**

Various hypotheses have been advanced to the mechanism/s of PQ-induced dopaminergic neuron injury. Apart from causing long-lasting dopamine overflow that keeps dopaminergic neuron in constant excitement and leads to excitotoxic injury (Shimizu et al., 2003), PQ also inhibits mitochondrial respiratory complex I and causes mitochondrial dysfunction, thus inducing
neuronal apoptosis or necrosis (Hauser and Hastings, 2013). Furthermore, the synthesis and fibrosis of alpha-synuclein in murine dopaminergic neuron has been shown to be accelerated by PQ, contributing to the formation of Lewy bodies, a cardinal feature of PD (Manning-Bog et al., 2002). In addition, widespread oxidative/nitrosative stress underlies its neurotoxicity, attesting to the critical role of Nrf2 as a key mediator of PQ-induced cell injury.

As critical mediators, potentially regulating the translation of at least 30% of vertebrate proteins (Ellwanger et al., 2011; Krek et al., 2005), miRNAs are highly relevant to neurodegeneration (Bicchi et al., 2013; Nelson et al., 2008; Roth et al., 2016). In our
previous study, the expression of a set of miRNAs in mouse substantia nigra was dysregulated by PQ, and miR-380-3p, which was restricted to brains of adult mice (Seitz et al., 2004), was downregulated as evidenced by ChIP analysis (Wang et al., 2017). Yet, the biological function of miR-380-3p has yet to be determined. Accordingly, herein, we focused on potential target genes of miR-380-3p, such as Sp3. Though the negative effect of miR-380-3p on cell survival and proliferation has been verified in the present study, it has yet to be established that miR-380-3p increases the risk of PD or other neurodegenerative disorders.

Sp3, better known as transcriptional factor Sp3, is a main member of Sp-like transcriptional factor family, and it is ubiquitously maintained in a wide spectrum of species, ranging from nematodes to human beings (Philipsen and Suske, 1999). Sp3 deficiency causes ossification impairment (Göllner et al., 2001), cardiac malfunction (van Loo et al., 2007), erythropoiesis impairment, and placental defects (Krüger et al., 2007). Specific knockdown of Sp3 also reduces cell proliferation in HepG2 cells (Gandhy et al., 2015). Furthermore, Sp3 is an established competitor of Sp1, and moderately reduces the transcriptional function of Sp1 (Yu et al., 2003), the latter promoting the synthesis of microtubule-associated protein tau (Maloney and Lahiri, 2012) and amyloid precursor protein (Docagne et al., 2004), key molecules associated with the progression of PD and Alzheimer’s disease (AD). In the present study, Sp3 deficiency induced by transfected miR-380-3p mimics and small interference RNAs also played a negative role in nerve cell survival and proliferation.

Though all of the above suggest a potential role of Sp3 in neurodegeneration, few studies have addressed the mechanism underlying the role of Sp3 in neurological disorders. In this study, p21 and CaM expression were altered by exposure to PQ, corroborating their role in PQ-induced neurotoxicity. Knockdown of Sp3 failed to alter the regulation of p21 and CaM, implying that Sp3, though verified in previous studies as being transcriptionally regulating p21 and CaM, probably failed to exert a major influence on p21 and CaM expression. As it was previously shown, in addition to its abundance, phosphorylation of Sp3 is positively correlated with its binding to the promoter

**Figure 4.** Paraquat (PQ) reduced the expression of miR-380-3p in Neuro2a (N2a) cells. A, N2a cells were treated with PQ at 100 or 200 μM for 24 or 48 h, and the treatments basically reduced miR-380-3p level in N2a cells, n = 3. **p < .01 or ***p < .001 for the comparisons labeled by the zig-zag lines. B, Both groups were pretreated with 40 μM tBHQ (tert-Butylhydroquinone) for 6 h, and the group exposed to 100 μM PQ for 48 h showed lower NQO1 mRNA abundance, n = 3. ***p < .001 versus the control group. C, Comparison of nuclear factor erythroid 2 related factor 2 (Nrf2)-specific coprecipitation efficiency between the 2 groups in (Figs. 1F and 1G), for 2 sites respectively, n = 3. p > .05 for the group exposed to 100 μM PQ versus the control group at both sites. D and E, Methylation condition of CpG: 29 in untreated N2a cells (D) and those cultured in 100 μM PQ for 48 h (E). The data were analyzed by BIQ analyzer and the probably erroneous or clone sequences were automatically removed before calculation.

**Table 4.** The Sequences of Designed siRNAs Targeting Specificity Protein-3 (Sp3) mRNA

| ID            | Sense (5’-3’)          | Antisense (5’-3’)          |
|---------------|------------------------|---------------------------|
| mmu-Sp3-659   | GCACCCAGGAUCAGAUUCAUTT | AUGAAUCUGAUCCUGUGGCTT     |
| mmu-Sp3-914   | GCAGAUUGGUGGCAGACUUTT  | AUGAUGGCAACCAUUUGGCTT     |
| mmu-Sp3-1643  | GCUCGCCAACAAAUUACUUTT  | AAGUAUUUGUGGCAGACGTT     |
| Negative control | UUCUCGGACAGGGACGUTT | ACGUGACAGGUUGGAGAAATT |
It has also been reported that SUMOylation (SUMO, small ubiquitin-related modifier) of Sp3 is one of the main regulatory events controlling the activity of this transcription factor, and potentially the molecular switch that determines whether Sp3 functions as a transcriptional activator or a repressor (Stielow et al., 2010). Such regulatory events may also be involved in the toxic cascade induced by PQ, altering the function of Sp3. As a main member of the Sp family, Sp3 has its binding box in the promoter region in a wide range of genes, which in fact provides it access to the control room of all kinds of downstream molecules via which Sp3 conducts its function. But in the present study, p21, CaM, and 2 well-known apoptosis mediators, Bcl2 and Bax, were unlikely to be among these downstream molecules. However, the N2a cells adopted in the present study was found to give responses different from other neuron or neuron-like cells (LePage et al., 2005), so it is too early to exclude these molecules from all situations (Figure 7).

Our novel studies established miR-380-3p bound to the 3'UTR of Sp3 and restrained the translation of Sp3 protein, suppressing N2a cell proliferation and enhancing the toxicity of PQ. In addition, knockdown of Sp3 altered N2a cell proliferation in a fashion analogous to miR-380-3p, and narrowed, or even eliminated, the gap between cells overexpressing miR-380-3p and the negative control groups. Surprisingly, inhibition of miR-380-3p did not induce detectable alterations in cell proliferation, apoptosis rate or cell cycle proportion, which is probably due to the low level of endogenous miR-380-3p in N2a cells; miR-380-3p was only detected in embryos and the adult brain at a fairly low level (Seitz et al., 2004). The low abundance of miR-380-3p likely makes it more difficult for inhibitors to exert their effects,
Figure 6. Calmodulin (CaM) and cyclin-dependent kinase inhibitor 1 (p21) were dysregulated by paraquat (PQ). A, Total RNA samples were extracted from Neuro2a (N2a) cells 48 h after transient transfection. Knockdown of specificity protein-3 (Sp3) did not alter the p21 and CaM mRNA level in N2a cells, n = 3, p > .05. B and C, Total RNA samples were collected 48 h after transient transfection and exposure to PQ. p21 and CaM mRNA level were altered by Sp3 siRNA but PQ, n = 3. *p < .05 or **p < .01 for the contrasts labeled by zig-zag lines. D, Western blots of Bcl2 protein from N2a cells 48 h posttransient transfection and treatment with PQ, n = 3, p > .05. E, Western blots of Bax protein from N2a cells 48 h posttransient transfection and treatment with PQ, n = 3, p > .05.

Figure 7. Proposed scheme of Nrf2/miR380-3p/Sp3 pathway. Paraquat translocates nuclear factor erythroid 2 related factor 2 (Nrf2) to the nucleus (the dashed oval) upon dislocation from Kelch-like ECH-associated protein 1 (Keap1). Next, Nrf2 activates the expression of miR-380-3p. It further attenuates the translation of specificity protein-3 (Sp3) mRNA. Sp3 might regulate the expression of downstream molecules such as Calmodulin (CaM) and cyclin-dependent kinase inhibitor 1 (p21).
mandating higher concentrations of inhibitors to achieve measurable inhibition. Thus, the Sp3 protein level was not elevated to a degree that was sufficiently high to impart obvious alterations in cell proliferation, apoptosis rate, or cell cycle proportion. In addition, N2a cells have been found by previous researchers to be much less sensitive to certain neurotoxins than some other neuron or neuron-like cells, mandating caution to extrapolate in capacity of miR-380-3p deficiency in vivo or in other cell lines from negative results in N2a cells (LePage et al., 2005).

Nrf2 is a transcriptional factor of many antioxidant enzymes. It is recognized as a crucial protective mediator in oxidant-induced neuronopathies. Earlier studies have established that several miRs are transcriptionally regulated by Nrf2 (Kurinna et al., 2014; Shah et al., 2014; Singh et al., 2019). Here, we report, for the first time, that miR-380-3p transcriptionally regulated by Nrf2, and inhibited the translation of Sp3 mRNA, suggesting that Nrf2 regulates Sp3 posttranscriptionally via miR-380-3p. This seems to counter the guardian-like portrait of Nrf2. Noteworthy, the fact that miR-380-3p is encoded within a miRNA cluster, suggests that other miRs, such as miR-379, miR-411, miR-329, to name a few, within the cluster are likely to be transcriptionally regulated by Nrf2; all these miRs are probably functional. For example, miR-380-5p, a miR sharing precursor with miR-380-3p, was found to target p53, a potent molecule known as the guardian of the genome, and function in cellular survival of neuroblastoma (Swarbrick et al., 2010), implying that Nrf2 may repress the activity of p53 via miR-380-5p. Additionally, a particular miRNA in vertebrates can regulate the translation of as many as hundreds of different mRNAs (Krek et al., 2005), thus, it is feasible that some other mRNAs, such as Cu5, Cln5, Mdg5a, Tceb3, Wapal, and Zbtb20, predicted by sequence-based analysis, might be targeted by miR-380-3p and contribute to its effects. However, the Nrf2/miR-380-3p/Sp3 pathway has yet to be determined as specific for PQ or widely involved oxidative stress-induced neurotoxicity.

PQ was demonstrated to attenuate the expression of miR-380-3p. In earlier studies, oxidative stress was shown to increase the expression of miR-153 and miR-146a, and these miRs subsequently inhibited the translation of Nrf2 (Jiao et al., 2015; Narasimhan et al., 2014; Smith et al., 2015). These effects may represent possible mechanisms by which impairs the expression of miR-380-3p. We also explored other possible pathways by which PQ might decrease miR-380-3p levels, such as changes in methylation in regions regulating the expression of miR-380-3p, namely the intergenic differentially methylated regions (IG-DMRs). The IG-DMRs were invariably highly methylated in response to treatment with various concentrations of PQ, suggesting that changes in methylation are unlikely to account for mechanisms by which PQ reduces the expression of miR-380-3p in N2a cells. However, the high methylation level may explain the restriction of miR-380-3p abundance, because high methylation level in the IG-DMR silenced the miRNA gene cluster encoding miR-380-3p (Seitz et al., 2004). The underlying mechanisms of PQ-induced neurotoxicity remain to be fully understood. Our future work should include further exploration on how PQ attenuates miR-380-3p expression, and a better characterization of the function of miR-380-3p.

AUTHOR CONTRIBUTIONS

H.L. and S.W. for conceptualized and supervised the study; Z.C. and F.Z. performed most of the experiments and analyzed the data; Y.D. contributed to experiments; Z.C. wrote the original draft; Y.Z., R.G., and J.L. contributed to discussion; F.Z., M.A., and Q.Z. reviewed and edited the manuscript. All authors have read and approved the final manuscript.

DECLARATION OF CONFLICTING INTERESTS

The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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