Neuronal injury and roles of apoptosis and autophagy in a neonatal rat model of hypoxia-ischemia-induced periventricular leukomalacia

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Abstract. As research into periventricular leukomalacia (PVL) gradually increases, concerns are emerging about long-term neuron injury. The present study aimed to investigate neuronal injury and the relevant alterations in apoptosis and autophagy in a PVL model established previously. A rat model of hypoxia-ischemia-induced PVL was established. In the model group, Sprague-Dawley (SD) rats [postnatal day 3 (P3)] were subjected to right common carotid artery ligation followed by suturing and exposed to 6-8% oxygen for 2 h; in the control group, SD rats (P3) were subjected to right common carotid artery dissection followed by suturing, without ligation and hypoxic exposure. At 1, 3, 7 and 14 days following modeling, brain tissue samples were collected and stained with hematoxylin and eosin. Cellular apoptosis was detected by terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay and the protein and mRNA expression alterations of neuronal nuclei (NeuN), caspase-3 and Beclin 1 in the model group were detected by western blot analysis and reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analyses. Compared with the control group, the protein and mRNA expression levels of NeuN (a marker of mature neurons) were markedly reduced, the number of positive cells was increased as detected by TUNEL, and the protein and mRNA expression levels of caspase-3 and Beclin 1 were elevated in the model group. In the rat model of hypoxia-ischemia-induced PVL, oligodendrocyte injury and myelinization disorders were observed, in addition to neuron injury, a decrease in mature neurons and the co-presence of apoptosis and autophagy. However, apoptosis and autophagy exist in different phases: Apoptosis is involved in neuron injury, while autophagy is likely to have a protective role.

Introduction

White matter injury (WMI) is the most common type of brain injury and the major cause of sequelae of the nervous system in premature infants. In the USA, ~60,000 extremely low birth weight infants are born every year (1) and the incidence rate of WMI in these infants is 50% (2,3). In addition, ~90% of WMI patients survive with sequelae to varying extents, of which 5-10% suffercerebral palsy and 50% experience behavior and/or attention deficit (4,5). The majority of patients with serious sequelae demonstrate periventricular leukomalacia (PVL) as observed by magnetic resonance imaging (MRI), pathologically manifesting as local periventricular white matter necrosis, myelination disorders, and gliocyte proliferation and microglial cell activation in adjacent white matter (6,7).

The pathogenesis of WMI remains to be fully elucidated, although infection and hypoxia are recognized to be the primary inducing factors. Infection and hypoxia may lead to the release of glutamates, oxygen-derived free radicals and inflammatory cytokines, and thus activate microglial cells. Microglial cells may further release glutamates, oxygen-derived free radicals and inflammatory cytokines which may aggravate WMI; activated microglial cells may induce the death of oligodendrocyte precursor cells (OPCs) and astrocytes, which has been previously demonstrated (8,9). Additionally, OPCs account for 90% of the total number of oligodendrocytes in the white matter of premature infants. Animal experiments have demonstrated that, based on the maturation-dependent vulnerability of oligodendrocytes, immature oligodendrocytes are more vulnerable during a specific window and are easily injured by glutamates, oxygen-derived free radicals and inflammatory cytokines, thus leading to apoptosis, necrosis and myelination delay (10-12).

However, in the recent years, follow-ups of patients with PVL have demonstrated that these patients frequently exhibit such sequelae as all-round cognitive deficit, attention deficit, human communication disorders, language retardation and autism, which cannot be reasonably explained by WMI alone;
these sequelae are closely associated with neuronal/axonal injury. With the development of neuropathology and imaging techniques, the focus of previous research has been directed to gray matter injury which has been long-neglected; a number of investigators have proposed that the pathological alteration sand neurodevelopmental outcomes of PVL may be not only limited to oligodendrocyte absence and myelination disorders (13-15). MRI volumetric analysis indicates that the volume of gray matter in the cerebral cortex and subcortex is decreased in premature infants with PVL. Andiman et al (13) demonstrated by autopsy that the density of granular neurons of patients with PVL was markedly decreased compared with that of controls. In 2005, Volpe et al (14) from Harvard University (Cambridge, MA, USA) first proposed a concept of ‘encephalopathy of prematurity (EP)’, and they hypothesized that brain injury may not be accurately defined by WMI in premature infants and that the role of neuronal injury in brain injury was relevant. However, there are few preclinical studies into neuron injury in PVL. In the present study, neuronal alterations were investigated using a previously established hypoxia-induced PVL model (15) to clarify the impact of neuronal injury and its mechanism.

Materials and methods

Experimental animals and grouping. Ethical approval for the present study was provided by Shengjing Hospital of China Medical University Ethics Committee (Shenyang, China). A total of 80 Sprague-Dawley (SD) rats [postnatal day 3 (P3); weight, 10.2-12.1 g] were randomly assigned to the control group (normoxic exposure; n=40) or the model group (hypoxic exposure; n=40). The rats were raised under the following conditions: The temperature was 18-26°C, the humidity was 47-70%, day light for 10 h and dark for 14 h, food and water was supplied ad libitum, and the food and water was supplied ad libitum. The body weight at the end of the experiment was monitored. The body weight of rats was 11.5±1.3 g in the control group (male, 22) and 11.4±1.4 g in the model group (male, 21) and no statistical difference was identified between the weight and sex of the two groups. All rats were provided by the Experimental Animal Department, Shengjing Hospital of China Medical University. A hypoxia-ischemia-induced PVL rat model established previously by our research group was used and the modeling method was same as that for a hypoxia model established by Mizuno et al (16). The neonatal SD rats (P3) were weighed and numbered prior to surgery. Animals were anesthetized by isoflurane inhalation and fixed on the bench in a supine position. Following conventional dissection, a 0.7-0.8 cm incision was made close to the tracheal midline under a dissecting microscope, and the right common carotid artery was exposed and ligated. Rats were postoperatively sent to the female rat cages with a room temperature and normoxic exposure for 2 h recovery, and were subsequently subjected to 2 h hypoxic exposure (6-8% oxygen). Rats in the control group were only subjected to the dissection of the right common carotid artery, without ligation and hypoxic exposure. At 1, 3, 7 and 14 days post-surgery, the brain tissues of experimental rats were collected.

Hematoxylin and eosin (H&E) staining. The brain tissues were fixed with 4% paraformaldehyde for 48 h at room temperature, dehydrated with gradient ethanol (70% ethanol for 2 h, 80% ethanol over night, 90% ethanol for 2 h, 100% II ethanol for 1 h, 100% II ethanol for 1 h) vitrified with xylene, waxed, embedded with paraffin and conventionally sliced into 5-µm tissue sections. Subsequently, the sections were stained with hematoxylin (ZLI-9609; Beijing Zhongshan Goldenbridge Biotechnology, Beijing, China) for 10 min, washed with water for 15 min, then eosin for 1 min (ZLI-9612; Beijing Zhongshan Goldenbridge Biotechnology), and washed with water for 10 sec. A total of six tissue sections from each group were selected randomly at various time-points, and were observed in 5 random fields under a light microscope at magnification, x400.

Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assay. The paraffin-embedded sections of brain tissues were conventionally dewaxed, treated with 50 µl 0.1% Triton X-100 (prepared with 0.1% sodium citrate) and preserved at room temperature for 8 min for vitrification, washed with PBS for 5 min three times, treated with 3% H2O2, at room temperature for 10 min to block peroxidase and washed three times with PBS for 5 min. TUNEL reaction solution (cat. no. 11684817910; Roche Diagnostics GmbH, Mannheim, Germany) was prepared with enzyme solution and label solution at a dilution of 1:9 (freshly prepared on ice). The sections were wiped dry, 50 µl TUNEL reaction solution was added, and sections were incubated in a humidified atmosphere away from light at 37°C for 60 min, followed by washing with PBS for 5 min three times. Thereafter, the sections were wiped dry, 50 µl converter-POD was added, and sections were incubated in a humidified atmosphere at 37°C for 30 min, and washed with PBS for 5 min three times. The sections were removed and wiped dry, 50 µl diaminoenzidene substrate (DAB; 1:20; zli-9017; OriGene Technologies, Inc.) was added at room temperature for 10 sec, and sections were rapidly placed into water to terminate the reaction when the color had just become deep. The sections were subsequently counterstained with hematoxylin at room temperature for 10 min, dehydrated with ethanol, vitrified with xylene and mounted. Instead of TUNEL reaction solution, PBS was used in the negative controls, and all procedures were same as the above. The cells with brown-yellow particles deposited in the cytoplasm were assessed to be positive cells. A total of six tissue sections from each group were selected randomly at various time-points, and they were observed in 5 random fields under a light microscope at magnification, x500.

Double-labeling immunofluorescence staining. The frozen sections (10-µM thick; frozen at -80°C) of brain tissues were defrosted, fixed with acetone at 4°C for 30 min, and washed with PBS for 5 min three times. Antigen retrieval was performed by boiling in antigen retrieval solution I at 100°C for 5 min, and washed with PBS for 5 min three times. Sections were blocked with bovine serum albumin (BSA; cat. no. bs043a; Biosharl, Shandong, China) at room temperature for 1 h. When the blocking solution was removed, the sections were incubated with primary antibodies against Beclin 1 (rabbit anti-rat antibody; 1:1,000; cat. no. ab55878) and neuronal nuclei (NeuN; mouse anti-rat antibody; 1:1,000; cat. no. ab104224) (both from Abcam, Cambridge, UK) at 4°C
overnight. The following day, the sections were rewarmed at room temperature for 1 h and washed three times with PBS for 5 min. Secondary antibodies [cat. no. ta130022; fluorescein isothiocyanate (FITC) conjugated goat anti-rabbit IgG, 1:64; and ta130012, Cy3-conjugated goat anti-mouse IgG, 1:50; OriGene Technologies, Inc.] were added and incubated away from light, at room temperature for 4 h, followed by washing in the dark with PBS for 5 min three times. Nuclear staining was performed with DAPI (OriGene Technologies, Inc.) for 5 min, and sections were washed in the dark with PBS for 5 min three times. Tissues were observed and photographed under a fluorescence microscope at magnification, x400 (MTC-600; Bio-Rad Laboratories, Inc., Hercules, CA, USA).

Western blotting. Brain tissues were collected from 3 rats in each group at each time-point. Tissues were added into radio immunoprecipitation assay lysis buffer (Beyotime Institute of Biotechnology, Haimen, China) at a dilution of 1:10, sheared into pieces using scissors, broken up with ultrasonication by 20 kHz frequency on ice for 5 min, and centrifuged at 4°C at 12,000 x g for 30 min. The supernatant was removed to separate the proteins, which were transferred into a 0.5 ml centrifuge tube. The protein concentration was measured using a bicinchoninic acid assay. The samples were adjusted to the same concentration, sample buffer was added, and samples were boiled for 5 min and preserved at -80°C. Proteins were mixed with 7.5% SDS-PAGE sample loading buffer and boiled for 5 min to achieve protein denaturation. The protein were loaded for 40 μg per lane in a preset sequence, and the electrophoresis apparatus was connected to the power source (80 V; the current was directed towards the anode) and powered off when bromphenol blue was migrated to a position 0.5 cm away from the bottom of the separation gel. The gel glass plate was unloaded from the electrophoresis apparatus and washed clean with deionized water. Two filter papers and a polyvinylidenedifluoride (PVDF) membrane were prepared. The cut PVDF membrane was soaked in methanol for 10 sec and transferred into the buffer solution with filter papers for soaking. A piece of sponge was placed on a splint, and two soaked filter papers were placed on the sponge and aligned accurately. The rinsed gel was placed on the top filter paper and air bubbles were removed with a glass tube. Thereafter, the PVDF membrane was placed on the gel aligning accurately, and 3 filter papers were placed on PVDF membrane. A piece of sponge was placed on the top filter paper, and the splint was clamped tightly when the air bubbles had been removed. One side of the gel was attached to the cathode and one side of filter membrane to the anode, and 100 V electrotransfer was conducted for 80 min. Subsequently, the filter membrane was stained with ponceau at 12,000 x g for 5 min. The supernatant was discarded, and the sediment was added with 1 ml precooled 75% alcohol, mixed evenly and centrifuged at 4°C (7,500 x g; 5 min). The supernatant was discarded, and the sediment was dried at room temperature for 10-15 min, added with 0.02 ml RNase Free water for dissolution, and cryopreserved at -80°C for use. The random primers and RNA template were transcribed using the Prime Script<sup>™</sup> RT reagent kit with gDNA Eraser (Perfect Real Time) (cat. no. RR047A; Takara Bio, Inc.) at 37°C for 1.5 h, inactivated at 70°C for 10 min and fixed in an ice bath for 5 min, and cDNA was preserved at -20°C for use. In order to control the sample loading errors during experimental procedures and the difference of amplification efficacy among various reaction tubes, GAPDH was set as internal reference, and the cycle was controlled in the linear phase of amplification reaction. The primers were as follows: GAPDH forward primer, 5'-GCAAGTTCAACGGCACA-3' and reverse primer, 5'-CATTGTGAATGCCGAGGAT-3'; solution, sealed, and slowly shaken on the shaking table at 4°C overnight. Following discarding of the blocking solution containing primary antibody, the PVDF membrane was placed into a hybridization bag containing 10 ml horseradish peroxidase-labeled secondary antibody (goat anti-rabbit IgG 1:5,000; cat. no. zb-2301; OriGene Technologies, Inc., for the PVDF membrane incubated with Beclin 1 antibody, caspase-3 antibody, and β-actin antibody; goat anti-mouse IgG 1:5,000; zb-2305; OriGene Technologies, Inc., for NeuN antibody) diluted with the blocking solution, sealed, and slowly shaken on the shaking table at room temperature for 1 h. Following discarding of the blocking solution containing secondary antibody, the PVDF membrane was rinsed with Tris-buffered saline and Tween-20 (TBST) for 15 min four times. Solution A and solution B of the enhanced chemiluminescence (ECL) reagents (Santa Cruz Biotechnology, Inc., Dallas, TX, USA) were mixed 1:1 in a dark room (as calculated by 0.125 ml ECL mixture/1 cm² membrane). The PVDF membrane was gently dried with filter paper and placed on a fresh-keeping film with the side containing protein facing upward. ECL mixture was evenly added onto the PVDF membrane to react for 1 min, and the PVDF membrane was lifted, dried gently with a filter paper and covered with the fresh-keeping film. Protein bands were scanned using Chemi Imager 5500 v2.03 software (ProteinSimple, San Jose, CA, USA), and the results were analyzed using a computerized image analysis system (FluorChem 2.0, ProteinSimple; Bio-Techne, Minneapolis, MN, USA). β-actin was used as the internal reference for comparison among groups. A higher relative gray scale indicated that the protein content was greater.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis. A total of 1 ml TRIzol (Takara Biotechnology Co., Ltd., Dalian, China) was added into 1-2 g frozen tissue and homogenized. Subsequently, the tissue homogenate was treated with 0.2 ml chloroform, heavily shaken, preserved at room temperature for 30 min, and centrifuged at 4°C for 1 h (12,000 x g; 15 min). The supernatant was discarded, and the sediment was added with 1 ml precooled isopropanol and mixed evenly, and the sample was preserved at room temperature for 10 min, followed by centrifugation at 4°C (12,000 x g; 10 min). The supernatant was discarded, and the sediment was added with 1 ml precooled 75% alcohol, mixed evenly and centrifuged at 4°C (3,000 x g; 5 min). The supernatant was discarded, and the sediment was dried at room temperature for 10-15 min, added with 0.02 ml RNase Free water for dissolution, and cryopreserved at -80°C for use. The random primers and RNA template were denatured at 70°C for 5 min, the reaction system was reverse transcribed using the Prime Script<sup>™</sup> RT reagent kit with gDNA Eraser (Perfect Real Time) (cat. no. RR047A; Takara Bio, Inc.) at 37°C for 1.5 h, inactivated at 70°C for 10 min and soaked in an ice bath for 5 min, and cDNA was preserved at -20°C for use. In order to control the sample loading errors during experimental procedures and the difference of amplification efficacy among various reaction tubes, GAPDH was set as internal reference, and the cycle was controlled in the linear phase of amplification reaction. The primers were as follows: GAPDH forward primer, 5'-GCAAGTTCAACGGCACA-3' and reverse primer, 5'-CATTGTGAATGCCGAGGAT-3';
caspase-3 forward primer, 5'-GAAGAGTTGGAGCAGCTGTA GCAC-3' and reverse primer, 5'-TGGATCGTAGACCCCTGT CG-3'; and Beclin 1 forward primer, 5'-ACTGATGGCCTGT AACGGTTCA-3' and reverse primer, 5'-CCAAGCATGGA CACAGAG3'. SYBR® Premix Ex Taq™ (TliRNase H Plus) (RR420A; Takara Bio, Inc.) was used for DNA amplification. The total volume of the mixture was 20 µl, and the reaction conditions were as follows: 95°C for 15 sec; 60°C for 1 min; 45 cycles in total. The expression of target genes was calculated using the 2^ΔΔCq method (17).

Statistical analysis. SPSS 13.0 software (SPSS, Inc., Chicago, IL, USA) was used for the processing and statistical analysis of data. The data are presented as the mean ± standard deviation. Statistical analysis was performed by two-way analysis of variance, followed by multiple comparison least significant difference (LSD) tests. P<0.05 was considered to indicate a statistically significant difference.

Results

Morphological alterations. Cells were observed under a light microscope and it was demonstrated that in the control group, cells were arranged tightly and regularly, with a normal structure, had a morphology same as that of brain tissues from healthy rats, and appeared normal. In the model group, cell karyopyknosis and necrosis were observed at day 1; at day 3, cells were unclear in hierarchy and structure, and periventricular white matter had a porous, malacotic structure; at day 7 periventricular white matter was loose and demonstrated net necrosis, and there was gliosis; at day 14, cerebral white matter and the corpus callosum became thinner, and fibers were disordered. However, no obvious gray matter nuclei and apparent cerebral cortex injury was observed by H&E staining (Fig. 1).

Decreased expression of NeuN. In the control group, NeuN mRNA expression was increased over the experimental time period. Compared with the control group, NeuN mRNA expression in the model group exhibited no significant change at day 1, although it was significantly decreased and reached a trough at day 3, followed by an increase at days 7 and 14; however, it remained lower compared with the respective control group (P<0.05; Fig. 2A). In the model group, NeuN protein expression was decreased compared with the control group. The trend observed was consistent with the mRNA expression (Fig. 2B).

Apoptosis of neurons
Alterations in neuronal apoptosis detected by TUNEL. In the model group, the nuclei of TUNEL-positive cells were stained in a scattered spot pattern and shaped irregularly. Neuronal TUNEL-positive cells were notably increased and widely present in periventricular white matter, corpus callosum, hippocampus and cerebral cortex. A small number of positive cells were observed in the control group. At days 1 and 3, the number of stained positive cells in the model group was markedly increased compared with the control group; at days 7 and 14, no difference was identified between the two groups (Fig. 3).

Increased expression of caspase-3. In the control group and model group, caspase-3 mRNA expression was significantly increased at day 3 compared with day 1 (P<0.01), although it was decreased at days 7 and 14 compared with day 1. At days 1, 3 and 7, caspase-3 mRNA expression in the model group was increased compared with the control group (P<0.05); at day 14, caspase-3 mRNA expression in the model group was increased compared with the control group, although the difference between the two groups was not statistically significant (P>0.05; Fig. 4A). In the control group and the model group, cleaved caspase-3 protein expression was significantly increased at day 3 compared with day 1 (P<0.01), and significantly decreased at days 7 and 14 compared with day 1. At all time-points, cleaved caspase-3 protein expression in the model group was greater when compared with the control group (P<0.05; Fig. 4B).

Neuronal autophagy
Double-labeling immunofluorescence staining of Beclin 1 and NeuN. Beclin1 was expressed in the cytoplasm, while NeuN...
was primarily expressed in the nuclei, exhibiting limited expression in the cytoplasm. At day 7, cells were arranged regularly in the control group and irregularly in the model group; the co-expression of Beclin 1 and NeuN in the same cells was observed in the two groups, and the number of cells with such co-expression was increased in the model group compared with the control group (Fig. 5).

**Increased expression of Beclin 1.** In the control group, Beclin 1 mRNA expression was significantly increased at days 7 and 14 compared with days 1 and 3 (P<0.05). Compared with the control group, Beclin 1 mRNA expression in the model group was significantly increased at days 1, 3 and 7 (P<0.01; Fig. 6A). In the model group, Beclin 1 protein expression at days 7 and 14 was markedly increased compared with day 3. Data were obtained by densitometry and were normalized using β-actin as a loading control. Values are expressed as relative optical density and are presented as the mean ± standard deviation. For each column, n=3. *P<0.05, **P<0.01 vs. the normoxia group. N, normoxia; M, model; NeuN, neuronal nuclei.

**Discussion**

Perinatal brain injury, one of the most common diseases in neonates and, particularly, premature neonates, may lead to altered development of the neonatal nervous system and result in varying degrees of sequelae. The most frequently reported manifestation is PVL (local or diffuse), which is the primary cause of cerebral palsy and may induce cognitive dysfunction, motor dysfunction or visual impairment (18,19). PVL is most frequently observed in premature infants and maybe caused by a number of factors, including perinatal infection and asphyxia. Previous studies of PVL have focused on oligodendrocyte involvement, myelinization disorders and microglial cell activation. OPCs are the primary targets organ for PVL (20,21). In general, PVL occurs in premature infants at a gestational age...
of 23-32 weeks; OPCs account for ~90% of the total number of oligodendrocytes in the cerebral white matter of these infants, although they are markedly decreased in >32-week infants, which is consistent with a reduction in the incidence rate of PVL. A number of previous studies have demonstrated the increased vulnerability of immature oligodendrocytes during a specific window (22,23). Animal experiments have confirmed that, based on the maturation-dependent vulnerability of oligodendrocytes, OPCs are easily injured by glutamates, oxygen-derived free radicals and inflammatory cytokines, thus leading to apoptosis, necrosis and myelinization delay (24,25).

Hypoxia increases the phagotrophic and oxidative stress injury-promoting functions of microglial cells through mediation of inflammatory cytokines, including γ-interferon, tumor necrosis factor-α, interleukin-1 and lipopolysaccharide causing microglia cells to release excitatory glutamates (26,27). Accordingly, the expression of α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid and N-methyl-D-aspartic acid receptors (NR1 and NR2A-D) has been observed to be increased in microglial cells (28). Activated microglial cells may induce the death of OPCs and astrocytes, which has been widely confirmed (29,30).

However, neuronal injury has also been frequently observed in PVL as the understanding of PVL has been increasing. Volpe et al (14), first proposed the concept of EP, and they observed non-negligible neuronal injury among brain injuries in premature infants. Andiman et al (13), reported a
38% decrease in the density of pyramidal neurons in the fifth layer of the cerebral cortex of premature infants with PVL. In addition, neuropathologists have previously demonstrated that the time at which neurons in the germinative zone migrated towards the cerebral cortex via the periventricular white matter coincided with the time when periventricular WMI occurred, and the toxicities of inflammatory cytokines and excitatory amino acids may injure these migrating neurons while damaging periventricular white matter (31). Such neurons (for example, subplate neurons) may exhibit the same pathogenesis as OPCs in white matter (32), express a number of glutamate receptors, and demonstrate selective vulnerability similar to OPCs (33,34). In the present study, neuronal injury in premature infants following hypoxia-ischemia was investigated using a hypoxia-ischemia-induced PVL model. The model was successfully established in a previous PVL study completed by the present research group (15). NeuN is a soluble nucleoprotein and a marker of mature neurons, and its molecular weight is 46-48 kDa. In the present study, NeuN expression was reduced and mature neurons were reduced in the model group, which indicated that the number of mature neurons was decreased in the rat model of hypoxia-ischemia-induced PVL; the presence of neuronal injury in PVL was confirmed in the animal models. NeuN expression reached a trough in the model group at day 3, suggesting that neuronal injury was most severe at day 3 following hypoxic exposure. NeuN protein expression was increased at day 7 to exceed the level at day 1; however, it remained low compared with the level observed in the control group, which demonstrated that a certain repair mechanism to achieve neuronal regeneration or transformation following neuronal injury was induced by hypoxia in premature infants, although NeuN protein expression was not able to be recovered to the same level as prior to the injury. The trough of NeuN expression in the model group was observed at day 3, perhaps due to the repair mechanism not having been initiated while tissue injury was ongoing. The present study further investigated the apoptosis and autophagy of injured neurons, in an attempt to elucidate the injury and repair mechanisms.

The following types of cell death have been recognized: Necrosis, apoptosis (type I apoptosis) and autophagic apoptosis (type II apoptosis) (35). Necrosis is a type of non-apoptotic cell death caused by external injury, and its morphological characteristics are different from those of apoptosis; generally, necrosis is accompanied by inflammation (36). Caspase-3 is a member of the caspase family and an important protein to detect in apoptosis, as there are caspase-3-dependent and caspase-3-independent forms of apoptosis. Cleaved caspase-3 is an active form of sliced caspase-3, and its protein expression elevation may indicate apoptosis increase (37). In the present study, caspase-3 expression was increased and reached a peak at day 3 (when NeuN expression reached at rough) in the model group, and thereafter it was decreased (mean while NeuN expression was increased). Therefore, it may be inferred that caspase-3-dependent apoptosis may be an important cause of neuronal injury. A number of studies have demonstrated that hypoxia may lead to an increase in neuronal apoptosis and result in brain injury (38-40). The peak expression of caspase-3 in the control group was additionally observed at day 3 (6 days following birth). There are previous studies investigating hypoxia-ischemia-induced brain injury in P5-7 rats (equal to full-term or nearly full-term human infants) (41,42). Therefore, it was considered that P5-7 maybe a window for neuronal injury and P5-7 rats may demonstrate the most marked apoptosis and the least tolerance to hypoxia. The TUNEL assay results in the present study further confirmed that neuronal apoptosis in the cerebral cortex was increased in the hypoxia-ischemia-induced PVL model. The results of the present study demonstrated that in the hypoxia-ischemia-induced PVL model, in addition to injury to oligodendrocytes, there was a decrease in the...
number of neurons and an increase in neuronal apoptosis, which validates the hypothesis that neuronal apoptosis is an important cause of neuronal injury in this model.

Autophagic programmed cell death is embodied by the appearance of numerous vacuolar structures, encapsulating the cytoplasm and organelles within the cytoplasm, and the degradation of their contents by lysosomes. In autophagic programmed cell death, intermediate filaments and microfilaments are redistributed, although not degraded, and actin remains in a polymeric form, so the cytoskeletal system is well-maintained. Conversely, in apoptotic programmed cell death, the depolymerization of actin and the degradation of intermediate filaments occur early on, and the cytoskeletal system is consequently damaged. Autophagy exerts an important role in the growth and development of cells, and the occurrence of diseases. A number of studies have reported that autophagy and apoptosis may antagonize or promote one another in certain cases, and they may occur successively or co-exist in the same cells (35,43-45). Beclin1 was the first autophagy-associated gene identified in mammals, and the Beclin 1 protein is able to bind to type III phosphatidylinositol 3-kinase (PI3K) to regulate autophagy (46). The results of the present study demonstrated that Beclin 1 expression was increased in the model group, with a trend coincident to the increase of NeuN expression and opposite to the decrease of caspase-3 expression. Therefore, it was hypothesized that autophagy may be involved in the inhibition of apoptosis and even the repair process of neuronal injury, which requires validation in subsequent experiments.

Balduini et al (38) suggested that autophagy may be part of a pro-survival signal containing the PI3K-Rac-α serine/threonine protein kinase-protein kinase mTOR axis, and that its activation mechanisms included drug effects, infection and hypoxia-ischemia. Carloni et al (47) demonstrated that rapamycin administration prior to hypoxia-ischemia markedly increased the expression of Beclin 1 and microtubule-associated proteins 1A/1B light chain 3B (autophagy-associated proteins), and decreased neuronal injury. These previous results indicated that facilitating autophagy prior to hypoxia-ischemia may have a certain protective effect on neurons, while interrupting autophagy may inhibit such protection. It was previously reported that blocking autophagy at 3 h post-ischemia/reperfusion had a neuroprotective role (48). Therefore, autophagy may be used as a target for the protective treatment of the nervous system. This opposite result demonstrates the double role of autophagy: Autophagy is protective in the early stage sand harmful in the late stages of nerve degeneration, which depends on the time of drug administration. Therefore, autophagy has a double effect on cells and neurons: In hypoxia-ischemia-induced injury, it may protect cells and prevent cell death, or may become one of the factors causing cell death, specifically depending on different phases of disease progression, alterations to the pericellular environment, and different intervention measures. The present study demonstrated that Beclin 1 was expressed in neurons, and that Beclin 1 expression was increased in the model group, close to the level of the control group at day 14 following hypoxia-induced injury; the time of the Beclin 1 expression increase coincided with the time window of repair following neuronal injury. Therefore, it may be inferred that cellular autophagy may exert a protective effect on neurons in the hypoxia-ischemia-induced PVL rat model, which requires confirmation in further experiments.

In conclusion, there are alterations to oligodendrocytes and neurons in a hypoxia-ischemia-induced PVL model. In the model group, the trough of neuronal cell expression was observed at the time-point when cellular apoptosis was in the most active state, which proves that neuronal apoptosis is an important cause of neuronal injury. Neuronal autophagy is additionally involved in this pathological process, although it is increased at the later stages of injury. Cellular autophagy may exhibit protective effect in neurons, although its specific role in this process has not been confirmed. The present study may promote a novel target for the treatment of PVL.

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Availability of data and materials

The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

Authors' contributions

LQ: Conceptualization of the study, performance of assays, data curation, formal analysis, investigation, validation, writing at the original draft and review and editing stage. JF: Conceptualization of the study, data curation, performance of assays, validation and writing at the review and editing stage. XX: Conceptualization of the study, data curation, performance of assays, validation and supervision. YS: Conceptualization of the study and performance of assays. LY: Conceptualization of the study, performance of assays, validation and writing at the review and editing stage. YT: Conceptualization of the study, data curation, formal analysis, investigation, software operation. YD: Performance of assays and writing of the original draft. YP: Performance of assays and writing of the original draft.

Ethics approval and consent to participate

Ethical approval for the present study was provided by Shengjing Hospital of China Medical University Ethics Committee (Shenyang, China).

Consent for publication

Not applicable.
Competition interests
All authors declared that they have no conflict of interest with regard to this study.

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