Trigeminal neuralgia causes neurodegeneration in rats associated with upregulation of the CD95/CD95L pathway

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Abstract
Objectives: To explore the effects of trigeminal neuralgia on neurodegeneration in rats and the underlying mechanism.
Methods: Sixty adult male Sprague Dawley rats were divided randomly into Chronic Constriction Injury of the Rat's Infraorbital Nerve (ION-CCI) group and sham group (n = 30). Right suborbital nerve was ligated in ION-CCI group to establish a trigeminal neuralgia model. In sham group, suborbital nerve was exposed without ligation. Pain thresholds were measured before surgery and 1, 7, 15, and 30 days after surgery (n = 10). Morris water maze tests (n = 10) were conducted at 1, 15, and 30 days after surgery to evaluate the changes in learning and memory ability of rats. At 5, 19, and 34 days after surgery, serum S100β protein concentration and hippocampal Aβ1-42 protein expression were detected by enzyme-linked immunosorbent assay; total tau protein expression was detected by Western blotting; changes of neurons in hippocampus were observed by Nissl staining; and the expression of ser404-p-tau, cluster of differentiation (CD)95, CD95L, and cleaved caspase-3 proteins was detected by immunofluorescence and Western blotting.
Results: Hyperalgesia occurred in ION-CCI group, and mechanical pain threshold decreased significantly (P < 0.05). On the 15th and 30th days after surgery, ION-CCI group showed lower learning and memory ability than sham group (P < 0.05). Serum S100β protein concentration, hippocampal Aβ1-42, and ser404-p-tau protein expression increased in the ION-CCI group 19 and 34 days after surgery (P < 0.05), hippocampal CD95 expression increased in the ION-CCI group after surgery (P < 0.05), hippocampal CD95L expression increased at 19 and 34 days after surgery (P < 0.05), and cleaved caspase-3 expression increased at 5 and 19 days after surgery (P < 0.05). Nissl bodies in ION-CCI group decreased significantly at 15 days after surgery. The expression of cleaved caspase-3 protein in ION-CCI group was positively correlated with the expression of CD95 and CD95L (P < 0.05).
Conclusions: Trigeminal neuralgia may lead to neuronal inflammation and neuronal apoptosis associated with upregulation of CD95/CD95L expression, thus causing neurodegeneration.

Keywords
Trigeminal neuralgia, CD95/CD95L, neuroinflammation, cell apoptosis, neurodegeneration, hippocampus, cognitive dysfunction, Nissl staining

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Introduction

Recent studies have found that patients with chronic pain are often accompanied by cognitive dysfunction, which seriously affects the quality of life of patients.\textsuperscript{1-3} Existing studies suggested that changes in hippocampal synaptic structure, changes in the expression of neurotransmitters (cytokines, brain-derived neurotrophic factors), and overuse of opioids might be the mechanisms of cognitive dysfunction in patients.\textsuperscript{3,4} but the exact mechanism remains to be further studied.

Trigeminal neuralgia is a typical clinical chronic neuropathic pain. Meskal et al.\textsuperscript{5} suggested that patients with trigeminal neuralgia are at risk of cognitive defects. In addition, studies have reported that trigeminal neuralgia are at risk of cognitive defects.\textsuperscript{15-17} In addition, neuropathic pain animal models.\textsuperscript{18,19} These evidences suggest that hippocampus is involved in the pathogenesis of cognitive impairment caused by trigeminal neuralgia.

Based on the above results, this study intends to explore whether trigeminal neuralgia affects the neurodegeneration of rats through the CD95/CD95L pathway in the model rats of trigeminal neuralgia.

Methods

Animals

Sixty-seven healthy and clean adult male Sprague Dawley rats weighing 200 to 250 g were used in this experiment, provided by animal experiment center of Southwest Medical University. All animal procedures were approved by the Ethical Committee of Southwest Medical University, Luzhou, China. Additionally, all animal procedures in the experiment in strict accordance with the regulations of the People’s Republic of China on the management of experimental animals and the methods for the quality management of experimental animals.

Three days before the experiment, the animals were trained to adapt the environment. The rats were knocked on the cage wall with a plastic rod, and the rats’ tails were lifted. After the rats were calm, the whisker pads were stimulated with a self-made brush for three times on each side. The bilateral stimulation was carried out alternately until the rats changed from initial exploration, fear curling, nose whipping, rapid escape, aggression, and other behaviors to silence. After three days of the above training, those who showed calmness and did not lose their whiskers were selected.

Rats were randomly divided into surgery group (Chronic Constriction Injury of the Rat’s Infraorbital Nerve (ION-CCI) group) and sham group, with 30 rats in each group and sham group as the control group. Chronic Constriction Injury of the Infraorbital Nerve (ION-CCI) was performed in the ION-CCI group to establish a trigeminal neuralgia model. In the Sham group, only the infraorbital nerve was exposed without ligating.
Surgical procedures

Anesthesia was given by intraperitoneal injection of sodium pentobarbital (40 mg/kg), a 1-cm incision was made along the upper margin of the right vibrissa pad of the rat to separate and expose the infraorbital nerve (Figure 1(a)), and then two chromium-enteric wires (4-0) were used to ligate the nerve at the proximal and distal ends, with an interval of about 2 mm. The standard of ligation was that the diameter of the infraorbital nerve became slightly thinner, but the conduction was not completely blocked; the blood circulation of the nerve outer membrane was unobstructed to prevent nerve ischemia and necrosis, and the incision was closed after surgery. The suborbital nerve was exposed in the sham group by the same method without ligation.

Behavioral testing

Mechanical pain threshold measurement was performed to evaluate whether the establishment of trigeminal neuralgia model is successful. Rats were placed in a mesh cage and waited until the rats were silent; the test subjects used the von Frey fiber filament to stimulate the rat’s surgical side whiskers (Figure 1(b)). Starting from the minimum stimulus intensity of 0.008 g, and gradually increasing, until the rats showed withdrawal and avoidance, aggression, and asymmetric face scratching behavior, it was considered as a positive reaction. Each stimulus intensity was tested 3 times/side, and when the positive number of the 3 stimuli was 2, the corresponding fibrofilament strength was the mechanical pain threshold of the rat. If the stimulation intensity was 26 g and the rats still did not have the above positive reaction, the test was stopped, and 26 g was taken as the threshold value of this test. The test time included 5 time points: before surgery and 1, 7, 15, and 30 days after surgery. At each time point, two groups of rats were randomly selected for pain threshold measurement (n = 10).

The Morris water maze (MWM) test was performed to assess learning and memory ability. Morris water maze consists of a black circular pool (160 cm diameter and 50 cm height), a cylindrical platform (12 cm diameter), and a computer camera analysis system. The water pool was covered by blue floodlight curtains and divided into four quadrants. Different markers were placed in each quadrant to provide visual clues for the rats to find the platform. An escape platform was placed in the middle of one of the quadrants, and the platform was 2 cm below the water surface. The temperature in the pool was maintained at 20°C-22°C, providing a stable environment for the rats to avoid external factors interfering with the behavior of the rats.

The rats trained continuously for four days, four times a day, 30 s apart. In each training, the rats face the inner wall of the pool and were put into the pool from the midpoint of the wall of each quadrant. The time for the rats to stand on the escape platform was the result of the time. If the platform was not found within 120 s, it will be recorded as 120 s. The average of four training sessions per day was used as the day’s escape latency. On the fifth day, the platform was removed, and the rats were put into the pool from the opposite quadrant of the original escape platform. The number of times the rats crossed the original platform and the percentage of time they spent in the target quadrant in 120 s were recorded.

At 1, 15, and 30 days after the surgery, the rats after the pain threshold measurement were subjected to the water maze experiment, which were recorded as Test 1, Test 2, and Test 3, respectively.

Sampling and sample preparation

After the completion of Test 1, Test 2, and Test 3 of the Water maze experiment, namely 5 days, 19 days, and
34 days after the operation, rats were sacrificed, and samples were obtained. At three time points, rats were randomly selected from two groups (n = 5) and anesthetized with sodium pentobarbital (40 mg/kg) to expose the heart and inferior vena cava. Blood was drawn from the inferior vena cava for 3-4 ml. Then, incubate with primary antibody overnight at 4°C (rabbit anti CD95, 1:300, Santa Cruz Biotechnology, Santa Cruz, CA, USA; rabbit anti-CD95L, 1:400, rabbit anti-cleaved caspase-3, 1:1000, rabbit anti-total tau, 1:1000, rabbit anti-ser404p-tau, 1:1000, Abcam, Cambridge, MA, USA). Then, the membranes were incubated with anti-rabbit secondary antibody (1:500, Proteintech, Chicago, IL, USA) at room temperature for 1 h. Finally, the bands were detected using enhanced chemiluminescence (Beyotime, Shanghai, China). Image J (National Institutes of Health, Bethesda, MD, USA) was used for quantitative band intensity: glyceraldehyde 3-phosphate dehydrogenase (GAPDH, 1:500, Abcam, Cambridge, MA, USA) was used as the sample control.

**Immunofluorescence test: expressions of CD95, CD95L, and cleaved caspase-3 proteins was detected**

The paraffin sections were de-waxed and gradient ethanol rehydrated, and then washed with distilled water. The sections were then placed in an antigen repair solution (Beyotime, Shanghai, China) and incubated in a water bath for 20 minutes. After the sections were naturally cooled to room temperature, the sections were removed and treated in 0.2% Triton X-100 solution (Amresco, USA) for 15 minutes. The sections were then sealed in 1% BSA for 30 min. They were then incubated with primary antibodies (CD95: 1:50, CD95L: 1:100, cleaved caspase-3: 1:200) Next, add different fluorescent-labeled secondary antibodies (Cy3: 1: 200, Fluorescein 5-isothiocyanate, FITC: 1: 100, CWBIO, Beijing, China) and incubate for 1 hour at room temperature in the dark. 4’-6-Diamidino-2-phenylindole (DAPI, Beyotime, Shanghai, China) was stained for 5 min. The hippocampus was viewed under an optical microscope (Olympus, Tokyo, Japan) (magnification 400).

**Nissl staining: observing the changes of neurons in hippocampus**

Paraffin slices were de-waxed to water routinely; then, reagent A (Methylene Blue Stain) was dripped for 10 min and then differentiated into reagent B (Nissl differentiation). The slices were observed under a microscope until the Nissl body was clear. Next, reagent C (Ammonium Molybdate Solution) was added to treat the slices for 4 min. The slices were quickly rinsed with distilled water, then dehydrated with anhydrous ethanol, then the slices were made transparent with xylene, and finally blocked slices with resinene.

**Statistical analysis**

SPSS19.0 statistical software was used. Experimental data were expressed as mean± standard deviation (x±s). Escape latency and swimming speed were analyzed by repetitive measure analysis of variance (ANOVA). The pain threshold, the number of times to cross the original target platform, the target quadrant time ratio, CD95, CD95L, Cleaved-caspase3, S100β, Aβ1-42, and Tau...
proteins were compared at different time points in the group using one-way ANOVA. Pairwise comparison within the group was performed using the Least Significant Difference (LSD) method, comparison between groups was performed using the Student’s t test, and analysis of correlation was performed using the Pearson analysis. \( P < 0.05 \) was considered statistically significant.

**Results**

**Changes of mechanical pain threshold in rats**

The pain threshold of ION-CCI group rats was higher one day after surgery than that before surgery \((P < 0.05)\), and the subsequent time points were significantly lower than that before surgery \((P < 0.01)\). This abnormal sensitivity to pain persisted during the 30-day observation, especially at the 15th day after surgery. The sham group showed no such change (Figure 2).

**Results of rat Morris water maze experiment**

In order to compare the effects of trigeminal neuralgia on cognitive function, we conducted three water maze experiments to observe the behavioral changes of rats, respectively, Test 1, Test 2, and Test 3. Each experiment included four consecutive days of positioning and navigation training and the fifth day of space exploration test. With the prolongation of learning time, the escape latency of rats was gradually shortened. In Test 1, there was no significant difference between the two groups in escape latency \((F = 0.153, P = 0.701)\) (Figure 3(a)) and swimming speed \((F = 1.553, P = 0.229)\) (Figure 3(d)), which means, 1 day after surgery, the learning function of the rats was not significantly impaired. In Test 2, the escape latency of ION-CCI group rats was significantly longer than that of sham group \((F = 101.315, P = 0.000)\) (Figure 3(b)). There was no statistical difference in swimming speed \((F = 0.485, P = 0.495)\) (Figure 3(e)), suggesting that trigeminal neuralgia impaired learning ability 15 days after surgery. In Test 3, the escape latency of ION-CCI group rats was longer than that of sham group \((F = 95.984, P = 0.000)\) (Figure 3(c)), and the swimming speed showed no statistical difference \((F = 0.010, P = 0.923)\) (Figure 3(f)), suggesting that trigeminal neuralgia has sustained damage to learning ability. On the fifth day of training, after the rats were removed from the platform, the space exploration experiment was conducted on the rats. Compared with sham group, the number of rats crossing the platform (Figure 4(a)) and the time percentage in the target quadrant of ION-CCI group were significantly reduced \((P < 0.01)\) (Figure 4(b)), while there was no significant difference between the two groups in Test 1 (Figure 4(a, b)).

**Expression of neurodegenerative-related proteins was increased in trigeminal neuralgia rats**

S100\( \beta \) protein is a specific protein of the nervous system, which is closely related to pain and cognitive function. Therefore, we used the ELISA to detect whether S100\( \beta \) protein in serum was upregulated. The results showed that there was no statistically significant difference in the concentration changes at postoperative time points in the sham group \((F = 0.602, P = 0.555)\), while the ION-CCI group was significantly higher at 19 and 34 days after surgery than the sham group \((P < 0.05)\), with the highest concentration at 19 days after surgery (Figure 5(a)). Excessive deposition of A\( \beta \) can lead to cognitive dysfunction. Therefore, we detected the expression of A\( \beta \)/42 protein in the hippocampus by ELISA. The results showed that there was no statistical difference in postoperative time points in the sham group \((F = 0.379, P = 0.692)\), and ION-CCI group was significantly higher than that in the sham group at 19 and 34 days after surgery \((P < 0.05)\), with the highest value at 34 days after surgery (Figure 5(b)). Tau protein phosphorylation level in the normal brain tissue is very low; when pathological changes occur, tau protein can be overphosphorylated at multiple sites, which affect cognitive function; therefore, we used the Western blotting to detect the tau protein expression in the hippocampus. The Western Blotting results showed that the changes of Total-Tau protein in the two groups of rats were not statistically different \((P > 0.05)\), while ser404-p-tau protein in the hippocampus of the ION-CCI group showed the same continuous increasing trend as A\( \beta \)l-42. Compared with the Sham group, it increased significantly at 19 and 34 days after surgery \((P < 0.05)\), and was highest at 34 days after surgery (Figure 5 (c) and (d)).

![Figure 2. Change of the mechanical threshold in two groups. Mechanical threshold. Data are expressed as mean ± SD (\( \#P < 0.05 \), compared to preoperative; \( ^{*}P < 0.05 \), compared to sham group). ION-CCI: Chronic Constriction Injury of the Rat’s Infraorbital Nerve.](Image)
Figure 3. Learning ability in rats on the water maze experiment: (a) escape latency in Test 1 \((F = 0.153, P = 0.701)\); (b) escape latency in Test 2 \((F = 101.315, P = 0.000)\); (c) escape latency in Test 3 \((F = 95.984, P = 0.000)\); (d) swimming speed in Test 1 \((F = 1.553, P = 0.229)\); (e) swimming speed in Test 2 \((F = 0.485, P = 0.495)\); and (f) swimming speed in Test 3 \((F = 0.010, P = 0.923)\). Data are expressed as mean ± SD (*\(P < 0.05\), compared to sham group).

ION-CCI: Chronic Constriction Injury of the Rat’s Infraorbital Nerve.

Figure 4. Probe trials in rats on the Morris water maze task: (a) the number of times to cross the original target platform in 3 Tests and (b) the target quadrant time ratio in 3 Tests \((P < 0.01)\). Data are expressed as mean ± SD (*\(P < 0.05\), compared to sham group).

ION-CCI: Chronic Constriction Injury of the Rat’s Infraorbital Nerve.
The expression of CD95/CD95L pathway was upregulated in the hippocampus of trigeminal neuralgia rats

Previous clinical studies have shown that CD95 plays an important role in neurodegenerative diseases, such as Alzheimer's disease. Therefore, we detected whether the CD95/CD95L pathway was associated with cognitive impairment induced by trigeminal neuralgia. We detected the expression of CD95, CD95L, and cleaved caspase-3 in the hippocampus of rats 5, 19, and 34 days after surgery by immunofluorescence and Western blotting.

Immunofluorescence results showed that the green fluorescence expression of CD95 and CD95L protein and the red fluorescence expression of cleaved caspase-3 protein were weak in the sham group, while the fluorescence expression of ION-CCI group was obvious (Figure 6(a) to (c)). Quantitative detection of hippocampal CD95, CD95L, and cleaved caspase-3 protein expression by Western blotting showed that no significant differences were found in CD95 ($F = 0.496, P = 0.621$), CD95L ($F = 2.054, P = 0.171$), and cleaved caspase-3 ($F = 2.463, P = 0.127$) expressions at various time points in the sham group. CD95 expression in ION-CCI group was significantly increased at each time point after surgery ($P < 0.05$), with the highest expression at 19 days after surgery (Figure 7(a) and (d)). CD95L showed a similar trend to that of CD95, which was significantly higher at 19 and 34 days after surgery than in the sham group ($P < 0.05$), with the highest expression at 19 days after surgery (Figure 7(b) and (e)). Cleaved caspase-3 expression was significantly higher at 5 and 19 days after surgery than in the sham group ($P < 0.05$), with the highest expression at 19 days after surgery (Figure 7(c) and (f)). We also measured the expression level of CD95, CD95L, cleaved caspase-3, total tau, and $\text{ser}^{404}$p-tau proteins in prefrontal cortex. To our surprise, we did not observe the same change trend in prefrontal cortex as in hippocampus, and they did not show a stable and consistent trend of change (data not shown).

The cleaved caspase-3 protein in hippocampus of trigeminal neuralgia rats was correlated with CD95/CD95L protein

The expression of cleaved caspase-3 protein in the hippocampal tissues of ION-CCI rats was positively correlated with the expression of CD95 protein ($r = 0.570, P = 0.027$) (Figure 8(a)), and the expression of cleaved caspase-3 protein was positively correlated with the
expression of CD95L protein \( (r = 0.650, \ P = 0.009) \) (Figure 8(b)).

Changes of neurons in hippocampus

In sham group, the structure of neurons in CA3 area of hippocampus was intact, nucleolus was obvious, and a large number of granular and tigroid Nissl bodies could be seen in the cytoplasm (Figure 9(a)). While in ION-CCI group, the neurons in CA3 area of hippocampus were mostly vacuolar, with no obvious nucleolus, and Nissl bodies dissolved or even disappeared (Figure 9(b)).

Discussion

Recent studies have found that head and facial pain is closely related to cognitive dysfunction.\textsuperscript{26–30} While trigeminal neuralgia is the most common chronic neuropathic pain in the head and face, its pathogenesis is complex, and the current mainstream is divided into central and peripheral mechanism; the central theory
of trigeminal neuralgia associated with voltage channel gating, and around the theory of trigeminal nerve, demyelination changes cause “short circuit” between adjacent nerves, and “short circuit” phenomenon of abnormal nerve impulses to the central cause sustained excitement. In order to simulate different pathogenesis, researchers have successfully established a variety of animal models of trigeminal neuralgia, including ION- 

Figure 7. Quantitative detection of hippocampal CD95, CD95L, and cleaved caspase-3 protein at various time points: (a) and (d) CD95 expression; (b) and (e) CD95L expression; (c) and (f) cleaved caspase-3 expression.

CD: cluster of differentiation; ION-CCI: Chronic Constriction Injury of the Rat’s Infraorbital Nerve; GAPDH: glyceraldehyde 3-phosphate dehydrogenase.

Figure 8. Correlation between cleaved caspase-3 protein and CD95/CD95L protein level in hippocampus in ION-CCI group. (a) The expression of cleaved caspase-3 and CD95 was positively related ($r = 0.570, P = 0.027$); (b) the expression of cleaved caspase-3 and CD95L was positively related ($r = 0.650, P = 0.009$); $P < 0.05$; there is statistical significance.

CD: cluster of differentiation.
CCI model, dental pulp perfusion model, and suborbital nerve injection model of cobra venom. This study is to replicate Liu et al.31 through the zygomatic line down to the road of ION-CCI model of trigeminal neuralgia. ION-CCI group rats showed a low response period one day after surgery, which may be due to axonal degeneration after nerve ligation, and the nerve impulse could not be transmitted through the ligation site, resulting in increased pain threshold. However, the specific mechanism remains to be further studied. The pain threshold was the lowest 15 days after surgery, which was considered to be the most severe myelin damage during this period. The pain threshold was slightly increased 30 days after surgery, this may be due to gradual absorption of the catgut, and the nerve was gradually repaired. This is consistent with the results reported in previous studies, indicating that the modeling is successful.

In this study, the Morris water maze experiment was conducted at three time points (1 day, 15 days, and 30 days after the surgery) from the acute stage, pain sensitive stage, and chronic stage after nerve injury in combination with the aforementioned changes in pain threshold, to verify that chronic trigeminal neuralgia would lead to spatial learning and memory dysfunction in rats. However, we found that at 30 days after surgery, the learning and memory ability of rats showed an opposite trend to the change of pain threshold, which may be due to the gradual recovery of the injury of the infraorbital nerve after catgut absorption, but the effect of pain on the central nervous system did not recover. Previous studies often choose to test the water maze behavior at a point in time, the data are relatively simple, and this experiment observed the three time points, more fully reflect the dynamic change of learning and memory ability of rats, so this study can reflect the trend of trigeminal neuralgia effect on cognitive function. However, whether the learning and memory abilities continue to decline or improve after 30 days remains to be further explored.

S100β protein is a specific protein of the nervous system. When the central nervous system is injured, S100β protein exudes from the damaged site to the cerebrospinal fluid and then enters the blood through the blood–brain barrier. In addition, studies have reported increased serum S100β protein concentration in AD patients32; therefore, serum S100β protein is often used as an indicator of the extent of reactive nerve injury and the prediction of neurodegeneration. The results of this study are consistent with previous studies showing that pain can lead to the upregulation of S100β protein levels.33–35 In particular, serum S100β protein levels were significantly upregulated 19 and 34 days after surgery, suggesting severe nerve damage in rats during this period. At the same time, the upregulation of S100β protein also suggested neurodegeneration in rats.

Excessive deposition of Aβ is an important pathological feature of AD; Aβ includes Aβ1-40 and Aβ1-42. Because Aβ1-42 is hydrophobic and it is much more likely to be the main ingredient in the SPs, this study

![Figure 9. Nissl staining: the changes of neurons in hippocampus: (a) morphology of hippocampal neurons in sham group and (b) morphology of hippocampal neurons in ION-CCI group.](image-url)
aims to test the expression of Aβ1-42. Abnormal phosphorylation of tau protein can reduce the binding ability of tau protein to microtubules, the combination of tau protein and microtubule cytoskeleton damage, leading to the formation of NFTs, thus causing nerve degeneration; Ser404 is considered to be a reliable marker of the transition from mild cognitive impairment to AD; moreover, phosphorylation of this site was crucial for the formation of paired helical filaments, so this study selected Ser404 site to detect the tau protein phosphorylation level. In our study, Aβ1-42 and ser404 p-tau level are raised after surgery, prompt degeneration of rat nerve may have happened when the pain threshold began to recover gradually, Aβ1-42 and ser404 p-tau level is still on the rise, the tip of the central nervous system may be irreversibly damaged, and its mechanism and the interaction is Aβ and p-tau synergies related to each other.36,37 Further research is needed to confirm this. 

$\alpha$-Synuclein protein, Aβ protein, and p-tau protein are neurodegenerative protein-related proteins. The expression of all three was upregulated in the hippocampus of trigeminal neuralgia rats, in combination with the water maze rats behavior change, further confirmed the trigeminal neuralgia can cause nerve degeneration assumption; this is consistent with previous studies that neuropathic pain can lead to cognitive dysfunction; in this study, we further explore its possible mechanism.

Recent studies have demonstrated the importance of the CD95/CD95L pathway to the nervous system, which is involved in the occurrence and development of a variety of neurological diseases such as AD,48–51 but whether it is associated with neuropathic pain is not clear. The main function of CD95/CD95L is to participate in the immune response, etc. And a number of studies suggest that pain may affect the immune response of the body.42–44 In this study, we found that trigeminal neuralgia resulted in upregulation of CD95/CD95L expression in the hippocampus of rats. The processing of pain information by the nervous system needs to be completed through a complex neural network. The upregulation of CD95/CD95L induced by trigeminal neuralgia may be related to the upward conduction of pain information. When pain information from the periphery is transmitted to the central nervous system, it may activate the hippocampus to participate in the processing of pain information, but the exact mechanism remains to be determined by further research.

The prefrontal cortex is also involved in the regulation of spatial learning and memory abilities, but in our study, no upregulation of these proteins was found in the prefrontal cortex, which suggests that CD95/CD95L pathway may be one of the molecules specifically expressed in the hippocampus in cognitive impairment caused by trigeminal neuralgia. Although hippocampus and prefrontal cortex have many similar functions, their specific molecules may not be consistent. Studies have shown that in inbred Roman High-Avoidance (RHA-1) strain rats with poor learning and memory ability and cognitive impairment, some synaptic-related molecules show different trends in hippocampus and prefrontal cortex.45 Of course, whether there are also molecular pathways related to cognitive impairment caused by trigeminal neuralgia in prefrontal cortex can be the direction of our future research.

Upregulation of CD95/CD95L may affect the occurrence of neurodegeneration through multiple pathways. It can promote the secretion of cytokines such as IL-6, IL-8, and IL-1β or the migration of immune cells such as neutrophils to cause neuroinflammatory reactions, which are caused by long-term chronic neuroinflammatory reactions. At the same time, CD95 is also a signaling molecule upstream of the caspase apoptosis pathway, which activates the cascade reaction of the caspase family after binding to CD95L, among which the activation shearing of caspase-3 is the most critical, and eventually leads to the apoptosis of cells due to the degradation of DNA. Cleaved caspase-3 is an iconic protein of cell apoptosis. The increased expression of cleaved caspase-3 protein after ION-CCI surgery suggested that trigeminal neuralgia increased the apoptosis of hippocampal cells, which was consistent with the results of previous animal studies that pathological pain increased the apoptosis of hippocampal cells in rats.8,19 Abnormal increase of apoptosis leads to excessive loss of neurons, which can lead to neurodegeneration. This study found that the expression of cleaved caspase-3 protein was positively correlated with the expression of CD95/CD95L, suggesting that the CD95/CD95L apoptosis pathway might be involved in the apoptosis of hippocampal cells in rats with trigeminal neuralgia. In addition, CD95/CD95L can affect Aβ and tau proteins by stimulating the activation of downstream caspase-3, and studies have found that caspase-3 can regulate the production of Aβ.46 The specific mechanism is: γ-secretase hydrolyzes amyloid precursor protein (APP) to produce Aβ, γ-secretase activating protein (GSAP) is the key protein that regulates this process, and GASP is regulated by Caspase3, so the activation of Caspase3 can promote the production of Aβ, and the activated Caspase3 can also cleave Tau protein and make Tau protein more likely to aggregate by exposing phosphorylation sites.

Nissl bodies exist in neuron bodies or dendrites and are basophilic granules. It can reflect the changes of morphological structure and function of neurons.47 In this study, the obvious histological changes of neurons in rats with trigeminal neuralgia were observed by Nissl staining, which is another direct and strong evidence that trigeminal neuralgia causes neurodegeneration in rats. In this study, the above experiments indicated that trigeminal neuralgia may lead to increased
neuroinflammation and neuronal apoptosis by upregulating the expression of CD95/CD95L, thus causing neurodegeneration. However, further studies are needed to determine the exact mechanism of upregulation of CD95/CD95L.

Conclusions
This study verified that trigeminal neuralgia can cause learning and memory dysfunction in rats and revealed that the upregulation of the hippocampal CD95/CD95L pathway may be involved in relevant mechanisms, providing a theoretical basis for the prevention and intervention of cognitive dysfunction in patients with trigeminal neuralgia.

Limitations
Trigeminal neuralgia causes neurodegeneration in rats associated with upregulation of the CD95/CD95L pathway, but the upstream impact factors of this pathway remains to be elucidated. The exact mechanism of the CD95/CD95L pathway needs to be further determined using knockout rats.

Declaration of Conflicting Interests
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