Abstract. Studies have shown that the cerebrospinal fluid-contacting nucleus (CSF-CN) may be associated with the transduction and regulation of pain signals. However, the role of the CSF-CN remains to be elucidated. Emerging evidence has suggested that neurokinin 1 receptor (NK1R) is important in the development of visceral pain and hyperalgesia, however, whether NK1R exists in the CSF-CN and its exact role in visceral pain remain to be fully elucidated. In the present study, double-labeled immunofluorescence staining and western blot analysis were performed to investigate this. It was revealed that NK1R was distributed in the CSF-CN. Following the induction of visceral pain by formalin instillation, NK1R in the CSF-CN was upregulated. In addition, by observing the behaviors of rats subjected to visceral pain, it was found that visceral pain was relieved by lateral intracerebroventricular injection of the NK1R antagonist, RP67580. These data provided a broader understanding of the role of NK1R in the CSF-CN and demonstrated that the CSF-CN was involved in acute visceral pain via the regulation of NK1R.

Introduction

Visceral pain refers to a type of pain, which occurs following noxious stimulation surrounding internal organs. Accompanied by ambiguous location and internal organ cramps, it causes severe pain and causes muscle pain in referred areas, which affects the quality of life of patients and substantially increases medical treatment burden. At present, how visceral pain occurs and the pathway of visceral noxious information transmission remain to be fully elucidated due to limitations in experimental techniques (1-4).

Materials and methods

Animals. All experiments were performed according to the regulations of The Committee for the Ethical Use of Laboratory Animal of Xuzhou Medical College. Male Sprague-Dawley rats provided by the Experimental Animal Center of Xuzhou Medical College (license number, SYXK 2002-0038) were 8-9 weeks old and weighed 250-300 g. The rats were randomly assigned into three groups, each containing six animals. The rats in the visceral pain group were subjected to lower colon instillation of 5% formalin; in the sham group, rats received instillation of saline; in the RP67580 group, a specific NK1R antagonist was injected into the lateral ventricle (LV) of rats 24 h prior to visceral pain induction. All animals were submitted to sacrifice by decapitation, and the brain was immediately harvested and postfixed in 4% paraformaldehyde for 24 h. Immunofluorescence staining and western blot analysis were used to detect the expression of NK1R in the CSF-CN. Substance P (SP), first identified in 1931, is a type of neurokinin. It is well documented that SP is an excitatory neurotransmitter released by the first stage of peripheral nociceptive afferent fibers and is involved in pain transmission. As its specific receptor, neurokinin 1 receptor (NK1R) has been demonstrated to be important in the development of pain and hyperalgesia. However, how NK1R functions when visceral pain occurs remains to be elucidated (5,6). The cerebrospinal fluid (CSF)-contacting nucleus (CSF-CN), the neurons of which contact the internal CSF, was first identified and named at the Jiangsu Province Key Laboratory of Anesthesiology, Xuzhou Medical College (Xuzhou, China). Using cholera toxin B subunit conjugated to peroxidase (CB-HRP), CSF-CN was successfully labeled and distributed in the ventral periaqueductal central gray matter of the brainstem (7,8). The CSF-CN is unique owing to its specific structure. The cell bodies of these neurons are in the brain parenchyma, whereas their projections are in contact with the CSF (9,10). Previous studies have demonstrated that there are subtle changes in the chemical composition of CSF when pain occurs (11,12). Considering the specific structure of the CSF-CN, the present study hypothesized that the CSF and the CSF-CN have certain indivisible connections. Evidence from previous studies has supported this hypothesis, and suggested that the CSF-CN is involved in the transduction and regulation of pain signals (13,14). However, the role of CSF-CN remains to be fully elucidated. The present study aimed to investigate the possible functions of NK1R within the CSF-CN under conditions of visceral pain in order to provide a feasible solution for the clinical treatment of visceral pain.
placed in separate rooms, in which conditions and light were controlled (23±1°C; 12/12 h dark/light cycle with light on at 8:00 a.m.), and food and water were available ad libitum.

Formalin instillation. In the animal experiments, visceral pain was induced by rectal infusion with dilute formalin. In order to allow the rats to regain consciousness as soon as possible, a small quantity of halothane (induction at 3%, then 1.5% in a mixture of 2:3 nitrous oxide and 1:3 oxygen) was used. While under anesthesia, the animal was suspended by its tail (<5 min). Subsequently, a polyethylene tube, which was wrapped in surgical tape at ~30 mm from the edge, was inserted through the anus. A 5% formalin solution or saline (100 μl) was injected through the tube slowly to prevent leakage (15).

Visceral pain behavior score assessment. All animals were placed in separate observation boxes for adaptation 30 min prior to the behavioral assessments. On regaining consciousness, the rats were observed for 3 h to record pain-associated behaviors. The visceral pain-associated behaviors included licking of the abdomen (A), back stretching (B), contraction of flanks (C), and contraction of whole body (D). Based on the duration of the contraction, the contraction of the whole body was divided into three levels of <30 sec (W1), 30-60 sec (W2) and >1 min (W3). Using the following formula, the visceral pain score (PS) was calculated: PS=L+2B+3C+4W1+5W2+6W3 (16).

Nociceptive assessment. The animals were placed in grid-bottomed cages for 30 min for adaption prior to the evaluation of allodynia. Subsequently, to determine the hyperalgesia of the referred area, von Frey hairs (VFHs; Semmes-Weinstein Monofilaments, North Coast Medical, Inc., San Jose, CA, USA) were used. On stimulation of the abdomen with increasing force (0.16, 0.4, 0.6, 1, 1.4, 2, 4, 6, 8, 10 and 15 g), the rats showed withdrawal responses, which included sharp abdominal retraction, licking or scratching the site of application of the hair and jumping with all four paws off the floor (17). Each VFH test lasted for 6 sec, or until a withdrawal response was observed. Once a withdrawal reaction occurred, the next descending force of VFH was applied to retest the area where the former VFH was placed until no response was detected. The minimum force was recorded as the abdomen withdrawal threshold in grams. If the result of allodynia was ≤2 g VFH, it was considered innocuous to the rats.

Drugs administration. The animals were anesthetized with sodium pentobarbital (40 mg/kg, i.p.). When immobilized in stereotoxic instrument, the rat was injected with 3 μl of 30% CB-HRP (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) into the LV according to stereotaxic coordinates (Brega, -1.2±0.4 mm; depth, 3.2±0.4 mm; right to median sagittal plane, 1.4±0.2 mm) 48 h prior to the behavioral assessments. The NK1R antagonist, RP67580 (Tocris Biosciences, Bristol, United Kingdom) was dissolved in dimethyl sulfoxide, and 100 pmol was injected into the LV of rats in the RP67580 group 24 h prior to the behavioral assessments.

Immunohistochemistry and imaging. Following the intracerebroventricular injections, the rats were administered with the formalin instillation and visceral pain behavioral assessments were performed. Subsequently, the rats were deeply anesthetized by injection of pentobarbital sodium (50 mg/kg, i.p.) and then successively perfused with 150 ml of phosphate buffered saline (PBS; 0.01 M; pH 7.4) and 4% paraformaldehyde. Following isolation and overnight fixation, the brainstem was immersed in 30% sucrose solution for 2 days. On a cryostat microtome (Leica CM1900; Leica Microsystems GmbH, Wetzlar, Germany), all tissues were successively cut into 35 μm slices in the transverse plane. The sections were then incubated in donkey serum (1:100 dilution; cat. no. 566460; Merck KGaA) for 1 h at 37°C, following which the sections were treated with goat anti-CB (1:400 dilution; cat. no. 227040; Merck KGaA) and anti-rabbit NK1R (1:100 dilution; cat. no. BA3678; Wuhan Boster Biological Technology Ltd., Wuhan, China). Following reaction with goat anti-IgG coupled to Alexa 546 (1:400 dilution; cat no. A1056; Thermo Fisher Scientific, Inc., Waltham, MA, USA) and rabbit anti-IgG coupled to Alexa 488 (1:400 dilution; cat. no. A21206; Thermo Fisher Scientific, Inc.), the slices were thoroughly rinsed three times with 0.01 mol/ml PBS at the end of each step. The tissue sections were transferred onto slides and allowed to air dry, prior to sealing with glycerol. Finally, ultrathin sections (75x25x1 mm) were prepared and examined under a laser-scanning confocal microscope (TCS SP2; Leica Microsystems). Using Image-Pro Plus (version 6.0; Media Cybernetics, Inc., Rockville, MD, USA), images were captured, cropped and adjusted.

Western blot analysis. Following the behavior assessments, the rats were sacrificed and the CSF-CN region of rat brain was removed and stored at -80°C. The tissues were homogenized by an electric homogenizer for 20 sec at room temperature. Following centrifugation at 13,523 x g for 15 min at 4°C, supernatant collection and denaturation, the lysates were electrophoresed on 8% SDS-polyacrylamide gel. According to the analysis, the semi-dry western blot transfer method was used, and the polyvinylidene difluoride membranes were blocked with 5% skim milk for 1 h at room temperature. The membranes were then incubated with anti-rabbit NK1R (1:500 dilution; cat. no. BA3678; Wuhan Boster Biological Technology Ltd.) and anti-rabbit GAPDH (1:5,000 dilution; cat no. A21206; Thermo Fisher Scientific, Inc.), the slices were thoroughly rinsed three times with 0.01 mol/ml PBS at the end of each step. The tissue sections were transferred onto slides and allowed to air dry, prior to sealing with glycerol. Finally, ultrathin sections (75x25x1 mm) were prepared and examined under a laser-scanning confocal microscope (TCS SP2; Leica Microsystems). Following exposure to goat anti-IgG coupled to Alexa 546 (1:400 dilution; cat no. A1056; Thermo Fisher Scientific, Inc., Waltham, MA, USA) and rabbit anti-IgG coupled to Alexa 488 (1:400 dilution; cat. no. A21206; Thermo Fisher Scientific, Inc.), the slices were thoroughly rinsed three times with 0.01 mol/ml PBS at the end of each step. The tissue sections were transferred onto slides and allowed to air dry, prior to sealing with glycerol. Finally, ultrathin sections (75x25x1 mm) were prepared and examined under a laser-scanning confocal microscope (TCS SP2; Leica Microsystems). Using Image-Pro Plus (version 6.0; Media Cybernetics, Inc., Rockville, MD, USA), images were captured, cropped and adjusted.

Statistical analysis and image analysis. Quantitative immunohistochemistry was used to assess the expression of NK1R in the CSF-CN. Image analysis software (Image-Pro Plus Version 6.0; Media Cybernetics, Inc.) was used for digital image analysis. On examination of every tissue section, details were recorded on the number, area and density of immunoreactive cells. Dual labeling of neurons with CB-HRP and NK1R was performed using CB-HRP/NK1R antibody (1:1,000 dilution; cat. no. A0545; Sigma-Aldrich) and goat anti-Boster Biological Technology Ltd. (Wuhan, China). Following reaction with goat anti-IgG coupled to Alexa 546 (1:400 dilution; cat no. A1056; Thermo Fisher Scientific, Inc., Waltham, MA, USA) and rabbit anti-IgG coupled to Alexa 488 (1:400 dilution; cat no. A21206; Thermo Fisher Scientific, Inc.), the slices were thoroughly rinsed three times with 0.01 mol/ml PBS at the end of each step. The tissue sections were transferred onto slides and allowed to air dry, prior to sealing with glycerol. Finally, ultrathin sections (75x25x1 mm) were prepared and examined under a laser-scanning confocal microscope (TCS SP2; Leica Microsystems). Using Image-Pro Plus (version 6.0; Media Cybernetics, Inc., Rockville, MD, USA), images were captured, cropped and adjusted.
Using SPSS v.13.0 (SPSS, Inc., Chicago, IL, USA), all data were analyzed using one-way analysis of variance or Student’s t-test. P<0.05 was considered to indicate a statistically significant difference.

Results

Behavior in formalin-induced visceral pain. Prior to surgery, the PS of visceral pain behavior at baseline was determined in untreated rats, which was 0.8±0.7. Following formalin instillation, a series of discrete behavioral episodes were observed, including licking of the upper abdomen, stretching of the whole body and arching of the back against the floor. Compared with the PS of rats in the sham group, the PS of rats in the visceral pain group presented with a typical biphasic time course. The first peak occurred at ~15 min post-instillation (PS=87.8±5.4), which was 15 times higher, compared with that in the saline instillation group (P<0.05). The second PS peak occurred at 60 min post-instillation (PS=36.7±3.3), which was 12 times higher, compared with that of the sham group (P<0.05). The PS of the RP67580 group was significantly decreased at 60 min post-formalin instillation, compared with that of the visceral pain group (P<0.05; Fig. 1A).

Mechanical allodynia in rats following formalin instillation. In the present study, the von Frey assessment was used to measure the abdomen withdrawal threshold. The values of the abdomen withdrawal threshold in rats of the visceral pain group were significantly lower 60 min post-formalin instillation, compared with that of the sham group (P<0.05). The values were significantly upregulated in rats pretreated with RP67580 (P<0.05; Fig. 1B).

Immunohistochemical analysis of NK1R in the CSF-CN of rats following formalin instillation. The CSF-CN was first identified by injecting CB-HRP, as a fluorescent tracer, into the LV of the rats, which was used as a reliable method to label CSF-CN. As shown in the images in Fig. 2, the neurons labeled by CB-HRP were predominantly located around the midline of the aqueduct midbrain and ventral periaqueductal gray matter of the brainstem. In order to determine whether the CSF-CN expressed NK1R, a double-labeled immunofluorescence technique was used. The results indicated that NK1R-immunoreactive neurons (green) were distributed in the CSF-CN. The total CSF-CN sample of the visceral pain group did not differ from that of the sham group (P>0.05). The number of simple labeled NK1R-containing neurons counted in the visceral pain group was significantly higher, compared with that in the sham group (P<0.05). Compared with the sham group, then number of dual-labeled CB-HRP/NK1R neurons counted in the visceral group was significantly upregulated (P<0.05; Fig. 2; Table I).

Western blot analysis of NK1R in the CSF-CN of rats experiencing visceral pain. Based on the above-mentioned findings, the present study investigated the role of NK1R in visceral pain. Western blot analysis was used to examine the protein expression levels of NK1R, the result of which indicated that NK1R was upregulated 60 min following formalin instillation, compared with that of the sham group (P<0.05).
Table I. Expression of CB-HRP, NK1R and CB-HRP/NK1R in the sham group and visceral pain group.

<table>
<thead>
<tr>
<th>Group</th>
<th>CB-HRP</th>
<th>NK1R</th>
<th>CB-HRP/NK1R</th>
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</thead>
<tbody>
<tr>
<td>Sham</td>
<td>398±16</td>
<td>114±7</td>
<td>70±3</td>
</tr>
<tr>
<td>Visceral pain</td>
<td>407±12</td>
<td>604±12</td>
<td>353±7*</td>
</tr>
</tbody>
</table>

CB-HRP is indicative of cerebrospinal fluid-contacting nucleus. Data are presented as the mean ± standard deviation. *P<0.05. NK1R, neurokinin 1 receptor; CB-HRP, cholera toxin B subunit conjugated to peroxidase.

Discussion

In previous studies, acute colitis has been used as a model to simulate visceral pain of pelvis organs, including the lower colon, rectum and bladder (16,17). Through the rectal infusion of formalin, inflammatory pain can be successfully simulated in regions of regions of the pelvic organs (18). In the present study, this method was used to induce visceral pain. The results showed that the visceral pain behavior scores of rats in the visceral pain group were significantly increased at each time point, compared with those of rats in the sham group, which suggested establishment of the visceral pain model had been successful. The present study demonstrated that the intestinal perfusion of formalin mediated a biphasic reaction of visceral pain, which was similar to the reaction of formalin-induced somatic pain (19). The specific mechanism underlying this biphasic pain remains to be fully elucidated and requires further investigation.

Our previous investigations demonstrated that the CSF-CN participated in the transmission of information within the brain, with chemical messages transferred in a cell-to-cell manner or in the CSF (20). It has been found that the neurons of the CSF-CN have a specific cell structure, with cell bodies located within the brain parenchyma and projections extending into the CSF through ependymal barriers (7). In addition, signal transmission by the CSF-CN may affect the composition of the CSF, which may lead to neuromodulation or neuroendocrine regulation (21).

NK1R is a G protein-coupled receptor, which can specifically combine with SP. It has been well documented that SP is involved in the pain regulation process (6). Several studies have suggested that CSF-CN is involved in the transmission of pain signals by the expression of SP, transient receptor potential cation channel, subfamily C, member 6 and extracellular-signal-regulated kinase 5 (10,22,23). In the present study, using a double-labeled immunofluorescent technique, it was observed that NK1R was expressed in the CSF-CN when visceral pain occurred, and that there were alterations in the PS and abdomen withdrawal threshold in rats exposed to formalin instillation, compared with those of naïve rats. To assess the involvement of NK1R of the CSF-CN in the behavioral noceptive symptoms due to visceral inflammation and to semi-quantitatively analyze variations in the expression of NK1R in visceral pain, western blot analysis was used. The results showed that, at 60 min post-formalin instillation, when the second phase of visceral pain occurred, the protein expression level of NK1R within the CSF-CN was upregulated. In addition, RP67580, a specific NK1R antagonist, significantly alleviated visceral pain in rats and reduced the expression of NK1R, suggesting that the upregulation of NK1R may be crucial in the process of establishing inflammation-induced visceral pain. Based on these findings, it was hypothesized that NK1R was involved within the CSF-CN in visceral pain.

Previous clinical and experimental studies have confirmed the presence of NK1R on postsynaptic dorsal column (PSCD) neurons in inflammation of the colon. These findings may explain why DC lesions can relieve visceral pain in patients (16). Studies on NK1R gene-deficient mice have shown that NK1R can mediate the central noceptive and peripheral inflammatory response. It can cause neurogenic inflammation and regulate the peripheral inflammatory responses to noxious stimuli (24). Evidence has suggested that the perception of visceral pain may be completed by an ascending excitatory pathway in the DC, particularly under
the conditions of peripheral inflammation. There is increasing evidence that this pathway may contain an amplification loop, which enhances the responsiveness of spinal cord neurons through a descending facilitatory pathway, possibly originating in the rostroventral medulla. PSDC and other projection neurons may be affected by this amplification loop and lead to potentiation (25). As NK1R is expressed in PSDC neurons under conditions of visceral pain, PSDC neurons may be involved in the amplification mechanism mediated by the NK1R. NK1R may regulate visceral pain through the transfer of information between the brain parenchyma and CSF, or be involved in the DC pathway via its expression in PSDC neurons. Additionally, the results of the present study showed that the visceral pain was not completely relieved following pre-injection of RP67580, which suggested that there may be other mechanisms for regulating visceral pain. A substantial number of neurons of the CSF-CN did not express NK1R, and other neurotransmitters or signaling pathways may affect the regulation of visceral pain.

In conclusion, the present study provided novel evidence that NK1R was expressed in the CSF-CN, and that NK1R within the CSF-CN may be involved in the regulation of visceral pain. However, the way in which NK1R in the CSF-CN affects the regulation of visceral pain remains to be elucidated in future investigations.

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References