Research Article

Low Frequency Electroacupuncture Alleviated Spinal Nerve Ligation Induced Mechanical Allodynia by Inhibiting TRPV1 Upregulation in Ipsilateral Undamaged Dorsal Root Ganglia in Rats

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Received 25 May 2013; Accepted 24 June 2013

Academic Editor: Yong-Qing Yang

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Neuropathic pain is an intractable problem in clinical practice. Accumulating evidence shows that electroacupuncture (EA) with low frequency can effectively relieve neuropathic pain. Transient receptor potential vanilloid type 1 (TRPV1) plays a key role in neuropathic pain. The study aimed to investigate whether neuropathic pain relieved by EA administration correlates with TRPV1 inhibition. Neuropathic pain was induced by right L5 spinal nerve ligation (SNL) in rats. 2 Hz EA stimulation was administered. SNL induced mechanical allodynia in ipsilateral hind paw. SNL caused a significant reduction of TRPV1 expression in ipsilateral L5 dorsal root ganglia (DRG), but a significant up-regulation in ipsilateral L4 and L6 DRGs. Calcitonin gene-related peptide (CGRP) change was consistent with that of TRPV1. EA alleviated mechanical allodynia, and inhibited TRPV1 and CGRP overexpressions in ipsilateral L4 and L6 DRGs. SNL did not decrease pain threshold of contralateral hind paw, and TRPV1 expression was not changed in contralateral L5 DRG. 0.001, 0.01 mg/kg TRPV1 agonist 6-IRTX fully blocked EA analgesia in ipsilateral hind paw. 0.01 mg/kg 6-IRTX also significantly decreased pain threshold of contralateral paw. These results indicated that inhibition of TRPV1 up-regulation in ipsilateral adjacent undamaged DRGs contributed to low frequency EA analgesia for mechanical allodynia induced by spinal nerve ligation.

1. Introduction

Neuropathic pain such as painful diabetic neuropathy, postherpetic neuralgia, trigeminal neuralgia, and poststroke pain is an intractable problem in clinical practice. It results from lesions or disease affecting the somatosensory nervous system either in the periphery or centrally and is characterized by spontaneous pain, allodynia, and hyperalgesia [1]. Up to now, the medication treatment of neuropathic pain is still unsatisfactory [2].

Abnormalities of channels or receptors in sensory nociceptors and dorsal root ganglia (DRG) are closely related to neuropathic pain [2, 3]. Transient receptor potential vanilloid type 1 (TRPV1), an important signal integrator in sensory nociceptors, plays a key role in neuropathic pain [4]. TRPV1 up-regulation contributes to mechanical allodynia and thermal hyperalgesia caused by various nerve injuries, while its antagonists can reverse the allodynia and hyperalgesia [5–7]. Furthermore, TRPV1 is coexpressed with various neuropeptides including calcitonin gene-related peptide (CGRP) in sensory ganglia and small sensory C and Aδ fibers [8, 9]. Activation of TRPV1 promotes CGRP release from nerve terminals [10, 11], which further aggravates neuropathic pain [12, 13].

Electroacupuncture (EA), a commonly used acupuncture method applying a pulsating electrical current to acupuncture needles for acupuncture stimulation, has been widely adopted for pain relief for decades [14]. Accumulating evidence shows that EA is effective in relieving neuropathic pain under multiple conditions like painful diabetic neuropathy,
nerve ligation, injury, and chronic constriction [15–19]. A former study showed that EA with low frequency reduced nociceptive response and normalized TRPV1 abnormalities in hind paw skin and DRG induced by nerve growth factor injection in rats [20]. However, little is known about the effect of EA with low frequency on TRPV1 under neuropathic pain condition, and whether this effect contributes to its alleviation of neuropathic pain.

For this sake, the rat neuropathic pain model induced by right L5 spinal nerve ligation (SNL) was used in the current study. The effects of EA with low frequency on mechanical allodynia and the expressions of TRPV1 and CGRP in ipsilateral L4–6 and contralateral L5 DRGs were systematically examined. A block study using TRPV1 agonist was also performed to further ascertain the involvement of TRPV1 in neuropathic pain relief by low frequency EA.

2. Materials and Methods

2.1. Animal Preparation. Male Sprague-Dawley rats (180–200 g body weight) were obtained from SLAC Laboratory Animal Co. Ltd., Shanghai, China. Rats were housed in temperature-controlled animal cages (25 ± 1°C) under a 12 h light and 12 h dark cycle, with free access to food and water. All animals were treated in accordance with the regulations of the State Science and Technology Commission for the care and use of laboratory animals (State Science and Technology Commission Order no. 2, 1988).

2.2. Experimental Design. Two experiments were conducted: (1) effects of low frequency EA on allodynia-like behavior and profiles of TRPV1 and CGRP in DRGs of rats with SNL-induced neuropathic pain and (2) effects of TRPV1 receptor agonist 6'-idoresiniferatoxin (6'-IRTX) on EA action. In experiment 1, rats were randomly divided into the following groups (n = 8 per group): normal, sham, SNL, and SNL + 2 Hz EA. In experiment 2, SNL-induced neuropathic pain rats were randomly divided into SNL + 2 Hz EA + vehicle, SNL + 2 Hz EA + 0.001 mg/kg 6'-IRTX, and SNL + 2 Hz EA + 0.01 mg/kg 6'-IRTX groups (n = 8 per group).

2.3. Spinal Nerve Ligation. L5 SNL was performed as previously described [21]. Briefly, rats were anesthetized with chloral hydrate (300 mg/kg, i.p.) and placed under a micro-surgical apparatus in a prone position. A midline incision was made at the L3–S2 level, and the dorsal vertebral column from L4 to S2 was exposed. The right L6 transverse process was carefully removed, and then the right L5 spinal nerve was carefully isolated and then tightly ligated with a 6-0 silk thread. Sham-operated animals were subjected to the same surgical procedure except that the isolated L5 spinal nerve was not ligated.

2.4. EA Treatment. Acupuncture needles of 0.25 mm in diameter were inserted approximately 5 mm deep into the ipsilateral acupoints Zusanli (ST36, 5 mm lateral to the anterior tubercule of the tibia) and Kunlun (BL60, at the ankle joint level and between the tip of the external malleolus and tendo calcaneus). The ends of the needles were attached to a pair of electrodes from an electrical stimulator (LH-202H, Huawei Co. Ltd., China). EA (2 Hz, 2 mA, 0.4 ms pulse width) was administered for 30 minutes once every other day from day 3 to day 15 after surgery. Since the analgesic effects of the two acupoints are well documented [10, 21], we did not carry out sham acupuncture for control. Animals were awake and calmed by placing the heads in black hoods with no physical restraint during EA treatment. Rats were subject to the same calming procedure in normal, sham, and SNL groups.

2.5. Drug Delivery. In experiment 2, 1 mg/mL 6'-IRTX (Sigma, USA) solution was made by dissolving 6'-IRTX in a 95% ethanol mixture which was used as the vehicle control. 6'-IRTX was intraperitoneally injected (0.001 mg/kg, 0.01 mg/kg, resp.) in a volume of 1 mL by adding saline 10 minutes before EA treatment on day 15.

2.6. Behavioral Testing. All tests were performed by an experimenter blinded to the treatment groups. After habituation, paw withdrawal threshold to a von Frey-like filament was measured to assess mechanical allodynia using a Dynamic Plantar Aesthesiometer (Ugo Basile, Italy) on day 0 (base), 1, 3, and 30 min after EA treatment on day 3, 7, 11, and 15 after surgery. The steel rod was pushed against the hind paw with linear ascending force (0–50 g) until a strong and immediate withdrawal occurred. Paw withdrawal threshold (PWT) was determined as the mean of three consecutive tests with intervals of 30 sec.

2.7. Immunofluorescence. Rats were deeply anesthetized by an intraperitoneal injection of 10% (w/v) chloral hydrate (3.5 mL/kg) and transcardially perfused with 150 mL cold sterilized saline followed by 500 mL cold, fresh 4% (w/v) paraformaldehyde in 0.01 M phosphate-buffered saline (PBS, pH 7.4). Contralateral L5 and ipsilateral L4, L5, and L6 DRGs were harvested, postfixed in the same fixatives for 2 h, and then consecutively immersed in 15% (w/v) and 30% (w/v) sucrose solution overnight at 4°C. DRGs were embedded in OCT (Bayer Corp., Elkhart, IN), frozen, and then cut in 10 μm sections. Sections were mounted on glass slides, rinsed in PBS (pH 7.4), and blocked for 1 h at 37°C in 0.01 M PBS containing 5% (v/v) normal donkey serum and 0.3% (v/v) Triton X-100. Sections were incubated overnight at 4°C with a primary antibody (sheep anti-rat TRPV1, 1:4000; Abcam, USA) dissolved in PBS containing 0.3% (v/v) Triton X-100 and 5% (v/v) donkey serum. After being washed in PBS, sections were incubated for 1 h at 37°C with Alexa Fluor 594-conjugated donkey anti-sheep IgG (1:400; Jackson Immunoresearch, USA). After immunostaining, sections were rinsed in PBS and cover-slipped with 50% (v/v) glycerol and 2.5% (w/v) triethylenediamine (antifading agent) in 0.05 M PBS. Images were obtained using a fluorescence microscope (Olympus IX71; Olympus, Japan) equipped with Image-Pro Insight 8.0 software (Media Cybernetics, USA).

TRPV1-immunoreactive (IR) analysis was performed as previously described [22, 23]. The observer was blinded for
the treatment groups. DRG neurons with a clear nuclear profile were counted using Image-Pro Plus 6.0 (Media Cybernetics, USA). TRPV1-IR ratio was calculated by dividing the number of TRPV1-IR neurons by the total number of neurons in each section. The mean ratio was determined from three every tenth DRG sections for each DRG.

2.8. Western Blotting. Rats were deeply anesthetized by an intraperitoneal injection of 10% (w/v) chloral hydrate (3.5 mL/kg). Contralateral L5 and ipsilateral L4, L5, and L6 DRGs were harvested, sonicated on ice in RIPA Lysis Buffer (Beyotime, China), and centrifuged at 10,000 × g for 10 min at 4°C, and then the supernatants were collected. Protein concentrations were determined by the bicinchoninic acid method. Protein extracts of animals in each group were equally pooled according to their concentrations. Samples were mixed with an equal volume of 2 × sample loading buffer and denatured by boiling at 100°C for 5 min. Proteins (30 μg/lane) were separated by an 8% SDS-PAGE gel for TRPV1 and 15% SDS-PAGE gel for CGRP and then transferred to 0.45 μm and 0.22 μm PVDF membranes, respectively (0.45 μm, Millipore, USA; 0.22 μm, Bio-Rad, USA). After being blocked in 0.01 M TBS with 0.1% Tween 20 and 5% dehydrated skim milk, the membranes were incubated overnight at 4°C with sheep anti-rat TRPV1 (1:2000; Abcam, USA), rabbit anti-rat CGRP (1:2000; Abcam, USA), or horseradish peroxidase-conjugated mouse anti-rat β-actin (1:10,000; Kangcheng, China). After being washed, the membrane for β-actin was visualized by chemiluminescence (ECL Plus; Amersham), while the membranes for TRPV1 and CGRP were incubated with the species-specific secondary antibodies for 2 h at room temperature and then washed and visualized by chemiluminescence (ECL Plus; Amersham, USA). Bands were detected by an Image Quant LAS 4000 System (Fujifilm, Japan) with Image Quant TL 7.0 software (GE Healthcare, USA). Three independent experiments were carried out for western blotting analysis. Target protein levels were normalized against β-actin levels and then expressed as relative fold changes compared to the normal control group [24, 25].

2.9. Statistical Analysis. All data were expressed as mean ± SEM. Statistical analysis was performed by one-way analysis of variance (ANOVA) followed by post hoc test of the least significant difference (LSD) for multiple comparisons. P < 0.05 was set as the level of statistical significance.

3. Results

3.1. EA Alleviated Mechanical Alldynia Induced by SNL. We first determined bilateral PWTs of rats to assess the extent of mechanical allodynia induced by SNL. Rats subjected to L5 SNL surgery developed mechanical allodynia in ipsilateral hind paw, as shown by a drastic reduction of ipsilateral PWLs (Figure 1(a)). 2 Hz EA significantly increased ipsilateral PWLs of rats subjected to SNL from day 3 after treatment, compared to the PWLs of SNL-controlled rats (P < 0.05, P < 0.01). Sham operation had no significant effect on ipsilateral PWLs during the whole period. Moreover, there was no significant difference in rats’ contralateral PWTs among normal, sham-operated, SNL, and SNL + EA groups (Figure 1(b)).

3.2. EA Inhibited Ipsilateral Undamaged DRG TRPV1 Upregulation Resulted from SNL. To determine the contribution of TRPV1 to the mechanical allodynia resulted from SNL, we performed immunofluorescence study and western blotting analysis to test TRPV1 expressions and levels in ipsilateral L4, L5, and L6 and contralateral L5 DRGs of rats. Immunofluorescence showed that TRPV1 positive neurons were mainly small-to-medium DRG cells (20–50 μm; Figure 2). TRPV1 expression in ipsilateral L5 DRG in SNL group (70 ± 1.0%) was significantly reduced as compared to that in normal group (14.7 ± 2.8%, P < 0.01).
Ipsilateral L5 DRG
Normal 
SNL 
SNL + EA

Ipsilateral L4 DRG
Normal 
SNL 
SNL + EA

Ipsilateral L6 DRG
Normal 
SNL 
SNL + EA

Contralateral L5 DRG
Normal 
SNL 
SNL + EA

Figure 2: Immunofluorescence of TRPV1 expressions in ipsilateral L5, L4, and L6 and contralateral L5 DRGs in normal, SNL, and SNL + EA groups. TRPV1-IR is mainly seen in small-to-medium DRG neurons (20–50 μm). Scale bar, 50 μm for all.

of TRPV1 expression in ipsilateral L5 DRG was observed between SNL and SNL + EA groups. TRPV1 expressions in ipsilateral L4 and L6 DRGs in SNL group (18.1 ± 1.8%, 15.3 ± 1.0%) were significantly increased as compared to those in normal group (10.2 ± 1.3%, 9.4 ± 0.6%; P < 0.001, P < 0.001, resp.). The increases were inhibited by 2 Hz EA (7.3 ± 0.9%, 5.5 ± 0.7%; P < 0.001, P < 0.001, resp., compared to SNL group). Besides, there was no significant difference in TRPV1 expressions in rats’ contralateral L5 DRG among normal, SNL, and SNL + EA groups.

The changes of TRPV1 protein levels as revealed by western blotting analysis were consistent with the changes of TRPV1 expressions. TRPV1 protein level in ipsilateral L5 DRG of rats in SNL group was significantly decreased as compared to that in normal group (P < 0.01), and no significant difference was found between SNL and SNL + EA groups (Figure 3(a)). TRPV1 levels in ipsilateral L4 and L6 DRGs in SNL group were significantly increased as compared to that in normal group (P < 0.05, P < 0.05, resp.). EA treatment with 2 Hz frequency fully inhibited SNL-induced increases of TRPV1 protein in ipsilateral L4 and L6 DRGs (P < 0.001, P < 0.01, resp., compared to SNL group) and even lower than normal TRPV1 levels (P < 0.01, P < 0.05, resp., compared to normal group; Figures 3(b) and 3(c)). No significant difference of TRPV1 levels in contralateral L5 DRG was observed in rats among normal, SNL, and SNL + EA groups (Figure 3(d)).

3.3. EA Inhibited Ipsilateral Undamaged DRG CGRP Upregulation Resulted from SNL. Since TRPV1 activation promotes CGRP release, which further strengthens TRPV1 role in pain sensation, we also performed western blotting analysis to detect CGRP protein levels. The changes of CGRP levels in ipsilateral DRGs were consistent with the changes of TRPV1 levels. CGRP protein level in ipsilateral L5 DRG in SNL group was significantly decreased as compared to that in normal group (P < 0.05), and no significant difference in CGRP protein level was found between SNL and SNL + EA groups.
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3.4.6-IRTX Blocked the Antiallodynic Effect of EA. To testify EA with 2 Hz frequency alleviated SNL-induced mechanical allodynia, at least in part, by inhibiting ipsilateral undamaged and adjacent DRG TRPV1 up-regulation, we performed block study via using TRPV1 ultrapotent agonist 6'-IRTX. Intraperitoneal injection of 0.001 and 0.01 mg/kg 6'-IRTX greatly reduced ipsilateral PWTs in EA-treated SNL rats (P < 0.001, P < 0.01, resp., compared to EA treatment-controlled group; Figure 5(a)). The two doses of injection blocked the antiallodynic effect of 2 Hz EA in ipsilateral hind paw, and 0.01 mg/kg 6'-IRTX was more potent than 0.001 mg/kg 6'-IRTX in counteracting 2 Hz EA action (P < 0.05). 0.001 mg/kg 6'-IRTX did not significantly change contralateral PWLs of SNL rats that received EA treatment, while 0.01 mg/kg 6'-IRTX significantly reduced contralateral PWL as compared to SNL + EA + vehicle and SNL + EA + 0.001 mg/kg 6'-IRTX groups (P < 0.05, P < 0.05, resp., Figure 5(b)).

4. Discussion

In the current study, the correlation of DRG TRPV1 with low frequency EA analgesia was for the first time investigated in rat neuropathic pain model induced by L5 SNL of the right
The rats’ ipsilateral but not contralateral PWTs were decreased after SNL surgery. It was interesting to find that L5 SNL caused reductions of TRPV1 and CGRP in ipsilateral L5 DRG, but up-regulations in ipsilateral undamaged L4 and L6 DRGs. Low frequency EA alleviated mechanical allodynia and inhibited TRPV1 and CGRP upregulations. Furthermore, TRPV1 agonist 6-IRTX fully blocked the antiallodynic effect of low frequency EA in ipsilateral hind paw.

Neuropathic pain may arise from nerve compression and trauma, diabetic neuropathy, chemotherapy-caused peripheral neuropathy, spinal cord injury stroke, and viral infections. Despite obvious differences in etiology, many of these pain conditions share common clinical phenomena: spontaneous pain together with allodynia and hyperalgesia [26, 27]. Nerve injury caused by SNL is a classic method to study neuropathic pain [28]. In the current study, rats subjected to L5 SNL developed prominent mechanical allodynia in ipsilateral hind paw. The signs of spontaneous pain such as guarding, licking, and lifting of ipsilateral hind paw were also observed. These results showed that neuropathic pain model was successfully made. Considering that mechanical allodynia was not further serious on day 3 after SNL surgery, we administered 2 Hz EA from day 3 and found that it effectively alleviated mechanical allodynia. Neuropathic pain relief by EA may be frequency specific. A former study showed that 2 Hz EA significantly attenuated hyperalgesia, whereas 120 Hz EA did not exert beneficial effect in diabetic neuropathic pain model [16]. The effectiveness of low frequency EA is also documented in other neuropathic pain models [17–19].

Changes in phenotype of primary sensory neurons following peripheral nerve injury contribute to allodynia and hyperalgesia in neuropathic pain [29]. TRPV1 is a member of TRP family channels which are ion channels that
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Ipsilateral paw withdrawal threshold (g)

| Base | D15 (pre-EA) | D3 (pre-EA) |
|------|--------------|-------------|
| SNL + EA + vehicle | SNL + EA + 0.001 mg/kg RTX | SNL + EA + 0.01 mg/kg RTX |

Contralateral paw withdrawal threshold (g)

| Base | D15 (pre-EA) | D3 (pre-EA) |
|------|--------------|-------------|
| SNL + EA + vehicle | SNL + EA + 0.001 mg/kg RTX | SNL + EA + 0.01 mg/kg RTX |

Figure 5: TRPV1 agonist 6'-IRTX blocked the antiallodynic effect of 2 Hz EA. 6'-IRTX was intraperitoneally injected 10 min before 2 Hz EA treatment on day 15. Data are presented as mean ± SEM, n = 8 per group. *P < 0.05, **P < 0.001, SNL + EA + 0.001 mg/kg 6'-IRTX group or SNL + EA + 0.01 mg/kg 6'-IRTX group versus SNL + EA + vehicle group; #P < 0.05, SNL + EA + 0.01 mg/kg 6'-IRTX group versus SNL + EA + 0.001 mg/kg 6'-IRTX group.

In conclusion, our data indicate that compensatory TRPV1 up-regulation in ipsilateral adjacent undamaged DRGs, with accompanied increased CGRP, may contribute to SNL-induced mechanical allodynia. EA with low frequency may alleviate SNL-induced mechanical allodynia through inhibiting TRPV1 up-regulation. Our study shows that EA with low frequency may be a useful approach for the management of neuropathic pain.

Conflict of Interests

There is no financial or commercial conflict of interests.

Acknowledgments

This study was supported by the Zhejiang Provincial Natural Science Foundation of China (LY12H27015, Z2100979), the National Natural Science Foundation of China (81072855).
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