Effect of Paroxetine on the Neuropathic Pain: A Molecular Study

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ABSTRACT

Background: Neuropathic pain, due to peripheral nerve damage, has influenced millions of people living all over the world. It has been shown that paroxetine can relieve neuropathic pain. Recently, the role of certain proteins like BDNF, GABAA receptor, and KCC2 transporter in the occurrence of neuropathic pain has been documented. In the current study, the expression of these proteins affected by paroxetine was evaluated. Methods: Male Wistar rats were allocated into two main groups of pre- and post-injury. Rats in each main group received paroxetine before nerve injury and at day seven after nerve damage till day 14, respectively. The lumbar spinal cord of animals was extracted to assess the expression of target genes and proteins. Results: In the preventive study, paroxetine decreased BDNF and increased KCC2 and GABAA gene and protein expression, while in the post-injury paradigm, it decreased BDNF and increased KCC2 genes and protein expression. In this regard, an increase in the protein expression of GABAA was observed. Conclusion: It seems that paroxetine with a change in the expression of three significant proteins involved in neuropathic pain could attenuate this type of chronic pain. DOI: 10.29252/ibj.24.5.301

Keywords: Brain-derived neurotrophic factor, Gamma-aminobutyric acid, Paroxetine

INTRODUCTION

Neuropathic pain occurs by damage to the somatosensory system, including peripheral and central neurons, through some diseases like diabetes mellitus, trauma, and cancer[1]. Although the aspects about the management of neuropathic pain have been changed, there are limited data on the pathogenesis of this pain[2,3]. In spite of the recent progress in neurosciences and pharmaceutical technology, no effective drug, with a clear mechanism, has been developed to manage the neuropathic pain[4].

Among different mechanisms involved in the neuropathic pain, the GABAergic system is the key player. GABA receptors located in pre- and postsynaptic terminals of primary afferent neurons are also found in the dorsal horn laminae I-IV[5]. Dorsal horn GABAergic interneurons play an important role in decreasing pain[6,7]. So far, the relationship between the GABAergic system and neuropathic pain has not been understood well[8]. It has been shown that GABAergic neurons transplanted to subarachnoid space attenuate hyperalgesia produced by nerve damage in rats[9]. Moreover, muscimol, as a GABAA receptor agonist decreases hyperalgesia caused by peripheral neurons injury[10]. The normal function of the GABAergic system is extremely dependent on cation-chloride cotransporters. The influx and efflux of chloride into and out of neurons are facilitated mostly by NKCC1 and KCC2, respectively[11,12]. Both NKCC1 and KCC2 expressed in spinal cord regulate intracellular chloride homeostasis. Various studies have indicated that the altered expression of these transporters can change neuropathic pain behavior[8,7,13,14]. On the other hand, the elevated concentration of intracellular chloride diminishes the inhibitory effect of GABA receptors[15].

New data show that glial cells in spinal cord cause hypersensitivity and continuation of pain in CNS[15].

List of Abbreviations:
BDNF, brain-derived neurotrophic factor; CNS, central nervous system; CCI, chronic constriction injury; DRG, dorsal root ganglion; KCC2, K+Cl- cotransporter 2; NKCC1, Na-K-2Cl cotransport 1; p38MAPK, p38 mitogen-activated protein kina; RT-PCR, reverse transcription PCR; SNL, spinal nerve ligation
Nerve injury not only increases specific microglia markers expression (Iba1) but also releases some painful mediators such as BDNF, prostaglandin E2, nitric oxide, and tumor necrosis factor-α. These factors produce hypersensitivity in CNS through rising the excitability and decreasing the inhibition of DRG neurons. Most of these mediators have a significant role in the production of chronic pain. Damage to nerve activates p38MAPK pathway in DRG and microglial cells. A recent study has suggested that the activation of the p38MAPK pathway by ATP and purinergic P2X4 receptor in microglia results in the production and release of BDNF. Investigations have also displayed that BDNF has a prominent function in neuropathic pain. Thermal hyperalgesia and mechanical allodynia are induced by intrathecal administration of BDNF. Other studies have revealed that nerve injury and peripheral inflammation change gene expression and production of BDNF. BDNF has ability to give rise to changes finally result in chloride efflux through GABAA and accordingly, depolarization of neurons. Based on evidence, alteration in KCC2 gene expression, thereby effluxing chloride from the neuron; these changes finally result in chloride efflux through GABAA and accordingly, depolarization of neurons. The GABAA receptor subtypes α2 and γ2 is expressed mostly in the spinal dorsal horn. Although γ2 subunit gene expression decreases after nerve injury, α2 subtype expression has no significant changes.

Considering P2X4 receptors role in the release of important mediators involved in neuropathic pain, inhibition of this receptor can be helpful to understand the mechanism of neuropathic pain. Up to now, no selective inhibitor of the P2X4 receptor has been presented. Antidepressant drugs, specifically tricyclic antidepressants have been widely used to manage neuropathic pain. It has been demonstrated that some antidepressants and antiseizure drugs are applied to treat neuropathic pain and inhibit the P2X4 receptor. Among the antidepressants, paroxetine has a significant inhibitory effect on P2X4 receptors. In the current study, we aimed to find out any possible changes in the expression of some proteins involved in the neuropathic pain (Iba1, BDNF, KCC2, and GABAA/γ2) affected by paroxetine.

**MATERIALS AND METHODS**

**Animals**

Rats (male Wistar, 150-200 g) used in the study were housed in an environment with controlled temperature (23 ± 2 °C). Food and water were available to animals without any limitation. At least one week before surgery, all the rats were permitted to be adapted to the housing facilities.

**Surgery and drug preparation**

The left sciatic nerve close to trifurcation was tied loosely (4 ligatures) by chronic gut suture, and thus, a model of neuropathic pain so-called CCI was created. Except for the sham group, the left sciatic nerve was tied in both drug and control groups. After ligation, the wound was closed. All surgical procedures were under sterile condition. Ketamine (60 mg/kg) and xylazine (10 mg/kg) were administered for the induction of anaesthesia. Paroxetine hydrochloride (Sigma, USA) was dissolved in DMSO 5%.

**Drug administration**

Animals were placed to pre- and post-injury groups. In each group, the rats were divided into CCI vehicle-treated (control), sham, and CCI paroxetine-treated groups. CCI- and sham-operated animals received the vehicle. Paroxetine was administered i.p. to the drug-treated group before and after surgery. In the preemptive paradigm, 10 mg/kg of paroxetine was injected to rats one hour prior to surgery and then daily after surgery until day 14. Animals in the post-injury group received the drug the same dose as the preventive group at day seven post-injury and then daily until day 14.

**Tissue collection for RT-PCR and Western blot analysis**

After euthanizing by CO2 asphyxiation, rats were decapitated immediately on day 14 post surgery. The spinal cord displaced by the normal saline from the vertebral column was frozen in dry ice. For evaluating gene and protein, the lumbar spinal cord segment was isolated from the intact frozen cord.

**Gene expression study**

Isothiocyanate-phenol-chloroform protocol was used for the isolation of total RNA, using RNX+ reagent (Cinaclon, Iran), according to the instructions provided by the manufacturer. Based on the manufacturer's protocol, 2 μg of total RNA, Oligo(dT) primer (Fermentas, USA), and M-MuLV reverse transcriptase (Fermentas) was used for the synthesis of cDNA. As shown in Table 1, designing of primer sequences (CinnaGen, Iran) was performed as per sequences in the GenBank. The PCR was carried out using the synthesized cDNA, the specific primers, and Taq DNA Polymerase MasterMix. In the beginning, the PCR was run for 10 min (95 °C), then continued by amplification cycles (25 or 26), each including 1-min denaturation (95 °C), annealing(45 s, 59 °C), and extension (45 s, 72 °C) steps. PCR products were
Paroxetine and Neuropathic Pain

Effect of prophylactic administration of paroxetine on the gene expression

Paroxetine and control groups showed a significant difference in the Iba1 expression ($p < 0.01$ and $p < 0.001$, respectively) relative to the sham group (Fig. 1). Compared to the control group, paroxetine decreased the expression of BDNF ($p < 0.001$). In comparison to the sham and control groups, the expression of KCC2 decreased and increased significantly in the paroxetine groups ($p < 0.05$) and paroxetine ($p < 0.05$) groups. However, paroxetine decreased the gene expression of GABAa/$\gamma$2 ($p < 0.05$) compared to the sham group.

Effect of paroxetine on gene expression after nerve injury

As depicted in Figure 2, a significant rise was seen in the expression of Iba1 in the control ($p < 0.001$) and paroxetine ($p < 0.01$) groups compared to the sham group. The paroxetine-treated group showed a significant decline in the expression of BDNF compared to the control ($p < 0.05$) and sham ($p < 0.01$) groups. KCC2 gene expression showed a significant reduction in the control group in comparison to the sham group $p < 0.01$). Compared to the control, the paroxetine group showed a significant rise ($p < 0.05$) in KCC2 gene expression. Moreover, no significant decrease was observed in the expression of GABAa/$\gamma$2 in the drug-treated group relative to other groups.

Statistical analysis

Statistical analyses were performed by SPSS 18 software using ANOVA, followed by Tukey’s post hoc test. Statistical significance was indicated by $p < 0.05$.

Ethical statement

The above-mentioned sampling and treatment protocols were approved by the Research Ethics Committee of Shahid Beheshti University of Medical Sciences, Tehran, Iran (ethical code: SBMU.REC1393.327).

RESULTS

Western blot analysis

For protein expression assay, Western blotting was used. Tissue samples were lysed in RIPA buffer (150 mM of NaCl, 1% NP-40, 50 mM of Tris pH 8.0, 1% SDS, 0.5% sodium deoxycholate, and 1 mM of EDTA and protease inhibitor cocktail) and centrifuged at 20,000 ×g at 4 °C for 20 min. After adding the SDS sample buffer to the aliquots of tissue extracts, the samples were placed in a water bath at 100 °C for 5 min. Proteins were separated on 10% SDS-PAGE, and then transferred to the blot. Blot membranes were incubated with 1:500 dilution of specific primary polyclonal rabbit antibodies against BDNF, KCC2, GABAa/$\gamma$2, and Iba1 and 1:1000 dilution of GAPDH (all from Abcam, USA) in TBS-T for 18 h. Blots were then incubated separately with secondary anti-rabbit (1:500 dilution; Abcam) in TBS-T for 90 min. KCC2, GABAa/$\gamma$2, BDNF, and Iba1 immune-reactive proteins were detected using chemiluminescence kit (Enhanced Chemiluminescence, Amersham Biosciences, UK). The signal intensity of the detected bands was measured by an image analysis system (Image j, version 1.46r).

Table 1. Primer sequences for the PCR amplification of genes of interest

<table>
<thead>
<tr>
<th>Target gene</th>
<th>Primer name</th>
<th>Sequence 5’ to 3’</th>
<th>Primer length (bp)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Iba1</td>
<td>F-iba1</td>
<td>5’ ACAAGACACTCTCCTCGATGATC 3’</td>
<td>21</td>
</tr>
<tr>
<td></td>
<td>R-iba1</td>
<td>5’ GCAACTCAGAAATAGCCTTTTCTG 3’</td>
<td>23</td>
</tr>
<tr>
<td>BDNF</td>
<td>F-bdnf</td>
<td>5’ GCTGCGCCCATGAAAGAACG 3’</td>
<td>20</td>
</tr>
<tr>
<td></td>
<td>R-bdnf</td>
<td>5’ GACCCCTCAGCATGTTTCTG 3’</td>
<td>21</td>
</tr>
<tr>
<td>KCC2</td>
<td>F-kcc2</td>
<td>5’ AGGAGGAGATGGACACGGCAGCC 3’</td>
<td>22</td>
</tr>
<tr>
<td></td>
<td>R-kcc2</td>
<td>5’ GCCTAGATGGCCAGGCACTG 3’</td>
<td>20</td>
</tr>
<tr>
<td>GABAa/$\gamma$2</td>
<td>F-gaba-a</td>
<td>5’ AGATATTAGCTTCTAATAAAAC 3’</td>
<td>22</td>
</tr>
<tr>
<td></td>
<td>R-gaba-a</td>
<td>5’ CACCATATTGCTTTCAATCG 3’</td>
<td>21</td>
</tr>
<tr>
<td>GAPDH</td>
<td>F-gadph</td>
<td>5’ GTTACCAGGGGCTGCTTCTTG 3’</td>
<td>23</td>
</tr>
<tr>
<td></td>
<td>R-gadph</td>
<td>5’ GTGTTGAGATGCAATGGCTGAC 3’</td>
<td>23</td>
</tr>
</tbody>
</table>
Effect of prophylactic injection of paroxetine target proteins

As it is clear from Figure 3, compared to the sham group, the expression level of the microglia marker (Iba1) rose in the control (p < 0.001) and paroxetine (p < 0.01) groups. Paroxetine could decrease the BDNF protein level significantly as compared to the sham (p < 0.01) and the control (p < 0.001) groups. Compared to the control group, KCC2 showed a significant rise in the paroxetine group (p < 0.05). On the other hand, a significant difference in GABAA/γ2 levels (p < 0.001) was found between paroxetine and control group.

Effect of paroxetine on the expression of target proteins after nerve injury

There was a significant rise in Iba1 protein level in the control and paroxetine groups as compared to the sham group (p < 0.001; Fig. 4). On the other hand, compared to the control group, the expression of BDNF protein decreased significantly in...
the paroxetine-treated group ($p < 0.01$). Paroxetine increased the KCC2 expression compared to the control group ($p < 0.05$). GABAA/γ2 protein levels showed no change in the drug-treated group in comparison with the control group.

**DISCUSSION**

In the present study, the effect of paroxetine on the expression of certain important mediators involved in neuropathic pain was investigated. Our data indicated an altered expression of BDNF and KCC2 upon administration of paroxetine. This was also the case for GABAA/γ2 proteins when the drug was injected before nerve damage. Microglial cells remain in resting state under physiological conditions and have appendages morphologically. Nerve injuries and lesions activate microglial cells and change their morphology with losing the appendages as well as swelling. After nerve injury, the gene expression level of P2X4 receptor and the Iba1 protein levels increased in microglial cells. It has been shown that Iba1 gene is specifically expressed in microglial cells in the CNS but not in other cells (neurons, astrocytes, and oligodendrocytes). Studies have evidenced the significant role of Iba1 in the migration and phagocytic activity of microglial cells. Moreover, the expression of this protein and mechanical allodynia-elevates after SNL in the rat. As noted previously, paroxetine has the most effect on the inhibition of the P2X4 receptor among various antidepressants and antiepileptic drugs used in neuropathic pain. A previous study has indicated that the expression of this receptor enhances in microglia after the nerve injury, and the neuropathic pain symptoms are also observed more frequently afterward. However, there is no change in the expression of this receptor in nerve cells or astrocytes. Another report has suggested that the P2X4 ionotrophic receptor is expressed only in microglia and its expression increases after neuropathy. Consistent with our data, some behavioral and biochemical findings have demonstrated that the activity of P2X4 receptors expressed in dorsal horn microglia is necessary for the induction of mechanical allodynia. A number of investigations have revealed that BDNF neurotrophin has a critical role in neuropathic pain and its expression boosts in the spinal cord dorsal horn and DRG. The activation of the P2X4 receptor in microglia is essential for the expression and release of BDNF after peripheral nerve injury, which in turn leads to the increased pain transfer in neurons. It has also been suggested that BDNF indirectly facilitates the release of GABA from the spinal cord interneurons, and BDNF expression increases in the spinal cord (dorsal horn) 24 hours after SNL, and this elevation continues for several days. In the present study, while after CCI, the expression of the BDNF gene enhanced significantly, paroxetine reduced the expression of this neurotrophin both before and after the nerve injury. Our previous surveys demonstrated that the prophylactic injection of paroxetine could diminish neuropathic pain. Considering the key role
of BDNF in neuropathic pain, this change in pain behavior is probably due to a shift in BDNF expression, which is reduced by paroxetine when administered before and after nerve damage.

KCC2 is a transporter for potassium and chloride ions and contributes to the regulation of anion gradients on both sides of the membrane. This transporter plays a very important role in regulating GABAA receptor activity[33]. The post-synaptic activity of GABAA receptors in the adult nervous system leads to the opening of the chloride channel, neuronal hyperpolarization and as a result, its inhibitory activity[34,45]. The role of KCC2 has been proven in chloride ion homeostasis in the spinal interneurons. KCC2 actively pumps the ion chloride to the outside of the neuronal cell to support conditions for the inhibitory activity of GABA receptors[46,47]. Moreover, the expression of KCC2 can contribute to the neuropathic pain induced by nerve injury[6,48,49]. In this study, paroxetine increased the expression of KCC2 as compared to the control group when used before or after nerve injury. Further researches are needed to find out whether paroxetine directly affects the expression of this protein, or secondarily, by blocking the purinergic receptor that causes this change. The GABAA receptors existing at the end of the primary afferent neurons are responsible for their synaptic inhibition[50,51]. It has been proven that GABAA/γ2 expression decreases significantly in the DRG after the nerve injury[25]. However, after the nerve injury, the expression of BDNF rises in DRG neurons, but the γ2 subunit of GABAA receptors decreases concurrently[52]. As mentioned before, KCC2, as a key protein in regulating the equilibrium potential of anions, is crucial for the GABAA inhibitory effect[53]. Reduced expression of the KCC2 protein in dorsal horn of the spinal cord neurons leads to the elimination of the inhibitory function of GABAA receptor in case of chronic neuropathy[23,54]. However, considering what mentioned before, the inhibitory effect of GABAA receptor depends on two important factors: first, the expression of the receptor itself and its subunits, and the second, the post-synaptic activity of the membrane protein of KCC2. The findings of this study showed that only pre-injury injection of paroxetine resulted in the increased protein level of GABAA/γ2 receptors.

In conclusion, it seems that paroxetine with change in the expression of some important proteins involved in neuropathic pain (BDND, KCC2, and GABAA) attenuates neuropathic pain. This effect was observed either when paroxetine was administered before nerve injury or when it was injected after damage to the nerve.

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![Graph showing effect of paroxetine on Iba1, BDNF, KCC2, and GABAA/γ2 protein levels administered after nerve injury. GAPDH was used as a loading control. Arbitrary unit: proteins optical density/GAPDH optical density. Data are expressed as means ± SEM. “p < 0.05, “p < 0.01, and ““p < 0.001 indicate a statistically significant difference when compared to the sham group, and “p < 0.05, “p < 0.01 show significant difference compared to the control group.](image)
Medical Sciences, Faculty of Medicine (Tehran, Iran) for their kind collaboration.

CONFLICT OF INTEREST. None declared.

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