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***Retrospective Study***

**Effectiveness of pulsed radiofrequency on the medial cervical branches for cervical facet joint pain**

Chang MC *et al*. Pulsed radiofrequency for CFP

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**Abstract**

BACKGROUND

Cervical facet joint pain (CFP) is one of the most common causes of neck pain and headache. Persistent CFP deteriorates the quality of life of patients and reduces their productivity at work.

AIM

To investigate the effectiveness of pulsed radiofrequency (PRF) stimulation of cervical medial branches in patients with chronic CFP.

METHODS

We retrospectively included 21 consecutive patients (age = 50.9 ± 15.3 years, range 26-79 years; male: female = 8:13; pain duration = 7.7 ± 5.0 mo) with chronic CFP, defined as ≥ 4 on the numeric rating scale (NRS). We performed PRF stimulation on the cervical medial branches. The outcomes of the PRF procedure were evaluated by comparing the NRS scores for CFP before treatment and 1 and 3 mo after treatment. Successful pain relief was defined as a ≥ 50% reduction in the NRS score at 3 mo when compared with the pretreatment NRS score.

RESULTS

No patient had immediate or late adverse effects following PRF. The average NRS score for CFP decreased from 5.3 ± 1.1 at pre-treatment to 2.4 ± 0.6 at the 1 mo follow-up, and 3.1 ± 1.1 at the 3 mo follow-up. Compared to the NRS scores before PRF stimulation, those at 1 and 3 mo after PRF stimulation had significantly decreased. Eleven of the 21 patients (52.4%) reported successful pain relief 3 mo after the PRF procedure. PRF stimulation on cervical medial branches may be a useful therapeutic option to control chronic CFP.

CONCLUSION

PRF stimulation of the cervical medial branches may be used as an alternative treatment method in patients with CFP. PRF can effectively alleviate CFP, and is safe to perform.

**Key Words:** Pulsed radiofrequency treatment; Zygapophyseal joint; Chronic pain; Pain; Neck pain; Pain management

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**Core Tip:** This is a retrospective study to investigate the effectiveness of pulsed radiofrequency (PRF) stimulation of cervical medial branches in patients with chronic cervical facet pain (CFP). Eleven of the 21 patients (52.4%) reported successful pain relief 3 mo after the PRF procedure. Compared to the numeric rating scale scores for CFP before PRF stimulation, those at 1 and 3 mo after PRF stimulation had significantly decreased after 1-month and 3-month follow-up. PRF stimulation on cervical medial branches may be a useful therapeutic option to control chronic CFP.

**INTRODUCTION**

Cervical facet joint pain (CFP) is one of the most common causes of neck pain and headache[1,2]. Clinicians frequently encounter patients with CFP, the prevalence of which ranges from 36% to 55%[1]. If CFP persists and progresses to chronic pain, its management becomes difficult[3]. Persistent CFP deteriorates the quality of life of patients and reduces their productivity at work[4]. Furthermore, it can cause affective disorders, such as depression and anxiety, and sleep disturbance[5]. Therefore, clinicians should actively control CFP.

            Several treatments, such as facet joint injection of corticosteroids, oral medication, and physical therapy, have been used to control CFP[6-8]. However, despite these treatments, many patients complain of persistent CFP. Conventional radiofrequency (CRF) stimulation of the cervical medial branch has also been used to control CFP[9,10]. This involves continuous stimulation, which causes the ablation of nerves and tissues by frictional heat from a catheter needle[6,7]. Due to this characteristic of CRF, neuropathic pain following the ablation of nerves can occur, and electrical burns after the procedure have been reported[10,11]. In contrast to CRF, pulsed radiofrequency (PRF) is a useful tool to alleviate chronic pain. This uses a brief stimulation, followed by a long resting phase[12]. PRF exposes the target nerves and tissues to an electric field, and rarely damages these structures[12]. Because of this minimal tissue-destructive characteristic, PRF has been rapidly adopted in clinical practice for the treatment of several types of pain, including neuralgia, joint pain, and myofascial pain[12-16]. Recently, several studies have reported a positive effect of PRF on medial branches in the spine to manage facet pain[17,18]. However, little is known about its effect on the cervical medial branch in the management of CFP.

             In the current study, we evaluated the effectiveness of PRF stimulation of cervical medial branches in patients with chronic CFP.

**MATERIALS AND METHODS**

***Study design***

This study was conducted retrospectively. We consecutively recruited patients who received PRF stimulation of the cervical medial branches under fluoroscopy in a pain clinic at a single university hospital from January 2014 to December 2019. The inclusion criteria were as follows: (1) PRF stimulation of cervical medial branches performed to control CFP; (2) aged between 20 and 79 years; (3) ≥ 3 mo history of axial cervical pain without radicular symptoms; (4) ≥ 80% temporary pain relief following a diagnostic cervical medial branch block with 0.5 mL of 1% lidocaine for each level prior to PRF stimulation of cervical medical branches; (5) ≥ 4 points on the Numeric Rating Scale (NRS, 0 = no pain, 10 = worst pain imaginable) prior to PRF stimulation of the cervical medial branches; and (6) no procedure to treat CFP performed ≥ 3 mo prior to the PRF stimulation. Each patient underwent cervical spine magnetic resonance imaging. We excluded patients who experienced cervical radicular pain due to disc herniation or foraminal stenosis and neck pain due to cervical canal stenosis. We retrospectively reviewed the medical records of 90 patients and included 21 patients (age = 50.9 ± 15.3 years, range 26-79 years; male: female = 8:13; pain duration = 7.7 ± 5.0 mo) in the analysis. A putatively painful cervical facet joint was selected on the basis of the distribution of pain and the location of tenderness[12]. All the included patients agreed to undergo PRF stimulation of cervical medial branches prior to the procedure. The Institutional Review Board of Yeungnam university hospital approved this study, and the need for written informed consent was waived due to the retrospective design of the study.

***Procedure***

An aseptic technique was adopted for PRF stimulation of the cervical medial branches using a posterior approach. For the procedure, patients were placed in a prone position, with the chest supported by a pillow, and the head slightly bent. Under the guidance of C-arm fluoroscopy (Siemens), a 22-gauge cannula (SMK Pole needle, 100 mm with a 10 mm active tip, Cotop International BV) was inserted in a posterior to anterior direction, and its tip was placed around the cervical medial branches, just lateral to the posteroanterior center of the C2-3 facet joint for the superficial medial branch of the third cervical spinal dorsal ramus (third occipital nerve), waists of the articular pillars of C3-C6 for C3-6 medial branches, and the apex of the superior articular process of C7 for the C7 medial branch (Figure 1). PRF stimulation of the superficial medial branch of the third cervical spinal dorsal ramus was conducted to control the C2-3 facet joint pain (third occipital nerve). For C3-4, C4-5, and C6-7 facet joint pain, the two vertically adjacent spinal medial branches, the C3 (deep medial branch of the third cervical spinal dorsal ramus) and C4, C4 and C5, and C6 and C7 medial branches were stimulated, respectively (Table 1)[19]. Once the needle tip was at the target site of the medial cervical branch, the needle was repositioned until the patient reported pain or a pressure sensation similar to the pain they usually experienced at less than 0.5 V to confirm the proximity to the medial cervical branch. An electrode was connected to the cannula, and the thoracic medial branch was stimulated (G4 radiofrequency generator; Cosman Medical Inc., Burlington, MA, United States). PRF treatment was administered at 5 Hz, with a 5-millisecond pulsed width for 360 s at 45 V under the condition that the electrode tip temperature did not exceed 42°C.

***Outcome measures***

Pain intensities were assessed using the NRS pain scores before and 1 and 3 mo after PRF treatment. Successful pain relief was defined as ≥ 50% reduction in the NRS score at 3 mo as compared with the pretreatment NRS score. Changes in NRS scores were also calculated as the difference between the pretreatment and 3 mo post treatment scores, to validate the degree of change in pain reduction [change in NRS (%) = (pretreatment score – scores at 3 mo post treatment)/pretreatment score × 100]. After 3 mo, the patient global perceived effect (GPE) was assessed using a 7-point Likert scale (Table 2)[20,21]. Patients reporting very good (score = 7) or good results (score = 6) were considered to be satisfied with the procedure.

***Statistical analysis***

Statistical analysis was performed with SPSS, version 23.0 (IBM Corporation, Armonk, NY, United States) for Windows (Microsoft Corporation, Redmond, WA, United States). The overall change in NRS scores over time was evaluated using a repeated-measures one-factor analysis. Multiple comparison results were obtained with Bonferroni correction. Statistical significance was set at *P* < 0.05.

**RESULTS**

None of the patients presented immediate or late adverse effects following the PRF procedure. The average NRS score for CFP declined from 5.3 ± 1.1 at pre-treatment to 2.4 ± 0.6 at the 1 mo follow-up and 3.1 ± 1.1 at the 3 mo follow-up. The NRS scores significantly changed over time (*P* < 0.001; Figure 2). Compared to the NRS scores before PRF stimulation, those at 1 and 3 mo after PRF stimulation were significantly decreased (*P* < 0.001). Eleven of the 21 patients (52.4%) reported successful pain relief (≥ 50%) at 3 mo after PRF stimulation.

On the 7-point Likert scale, Good (score = 6) and fairly good results (score = 5) were observed in 11 (52.4%) and 5 patients (23.8%), respectively. However, no change in results (score = 4) was observed in 5 patients (23.8%). Accordingly, 11 patients (52.4%) were satisfied with the results 3 mo after the PRF procedure. Very good (score =7), fairly bad (score = 3), bad (score = 2), and very bad (score = 1) scores were not reported. These findings demonstrated that PRF stimulation was effective at alleviating CFP, and more than half of patients who received the treatment were satisfied with the results of this treatment.

**DISCUSSION**

In the current study, we found that PRF simulation of the cervical medical branches could effectively control chronic CFP. After undergoing PRF stimulation of the cervical medial branch, significant pain relief was observed in patients with CFP, and approximately half of the patients reported successful pain relief (≥ 50% pain reduction); this effect lasted for at least 3 mo. Furthermore, about half of the patients reported successful pain relief and satisfaction with the results following PRF stimulation.

Facet joints are true synovial joints. It is assumed that the production of inflammatory cytokines and matrix-degrading enzymes disturbs chondrocyte metabolism, leading to cartilage degradation, as in other osteoarthritic joints[22]. Repetitive chemical and mechanical stress on cervical facet joints causes inflammation and narrowing of the capsule, resulting in osteoarthritis and chronic CFP[23]. Additionally, facet joint injury can occur due to whiplash injury following a sudden acceleration-deceleration force, which is a common cause of chronic CFP[19].

Medial branch nerves are very small nerve branches that carry pain signals from facet joints to the brain. There are various treatment methods for CFP. Physical therapy, manipulation, mobilization, oral medication, and cognitive behavioral therapy may all be applied, but their pain-reducing effect is controversial[24]. Three types of interventions for the treatment of CFP include intraarticular facet injections, medial branch blocks (MBBs), and neurolysis of medial branch nerves using radiofrequency[25]. The MBB is performed with corticosteroids and local anesthetics to reduce CFP. This may provide pain relief by suppressing nociceptive discharges and blocking the axonal transport and sympathetic reflex arc, thereby exerting anti-inflammatory effects[17]. However, local anesthetics can cause various adverse effects, such as hypotension, dizziness, nausea, seizures, and cardiac arrest[26]. Moreover, repeated corticosteroid injections can cause hyperglycemia, suppression of the hypothalamic-pituitary-adrenal axis, and osteoporosis[27]. To avoid the side effects of local anesthetics and corticosteroids, PRF stimulation was suggested as an alternative treatment method for CFP. No previous study has yet directly compared the effect of PRF stimulation to the cervical medial branches with other treatment methods for non-traumatic facet pain. Therefore, this study aimed to investigate whether PRF stimulation was effective in the management of chronic CFP.

PRF stimulation is a minimally neuro-destructive treatment applied in clinical practice to treat pain related to the facet joint, without inducing any significant complications[9]. The main advantages of PRF stimulation are that the procedure is painless and does not induce thermal damage to the tissues. PRF produces an electric field, which exerts a local or regional effect on immune cells, thus preventing progression to chronic pain[28,29]. The nociceptive inputs may be reduced along the pain pathways, and the electrical fields produced by PRF may alter the synaptic signal transmission[12]. Furthermore, PRF stimulation is reported to decrease microglia activity in the spinal dorsal horn[28]. Because microglia release several cytokines and chemokines that are associated with progression to chronic pain, the down-regulation of microglia activity can control pain[28]. Additionally, PRF stimulation may cause microscopic damage to unmyelinated C fibers that transfer the pain sensation[30].

The effect of PRF stimulation on the management of patients with CFP was documented in two studies. Mikeladze *et al*[30] investigated the effect of PRF on patients with cervical or lumbar facet joint pain. More than half of the patients (68 out of 114 patients) reported pain relief of 50% or more after PRF stimulation at 42°C for 120 s. Liliang *et al*[31] enrolled patients with whiplash-related chronic CFP, and showed that PRF stimulation of the cervical medial branches relieved pain and reduced medication requirement. Our study included patients with only CFP, and the enrolled patients were not confined to those with a history of trauma. In line with these previous studies, the results of our study support the fact that PRF stimulation is safe and might effectively relieve CFP. In our study, PRF simulation was performed by a single physician with approximately 20 years of spinal intervention experience. Therefore, the risk of operator bias is low. Five patients in our study showed no improvement in CFP after PRF stimulation. This may be due to different underlying mechanisms involved in the development of chronic pain, which may be varied and complex[32]. Individualized treatment plans are required for the appropriate management of CFP.

However, there are several limitations to this study. First, the sample size was small. Second, this study lacked a placebo-controlled group. However, there are ethical considerations regarding the use of placebo in a controlled trial with patients who suffer from moderate to severe pain. Third, this study was conducted retrospectively. Fourth, the level of the origin of CFP was determined on the basis of distribution of pain, potentially adding a subjective component to our study. Fifth, we did not measure a beneficial effect on the quality of life. Further studies, including randomized controlled trials, are needed to compensate for these limitations. Authors should discuss the results and how they can be interpreted from the perspective of previous studies and of the working hypotheses. The findings and their implications should be discussed in the broadest context possible. Future research directions may also be highlighted.

**CONCLUSION**

In conclusion, we found that CFP was significantly reduced at 1 and 3 mo after PRF stimulation. The rate of successful pain relief and patient satisfaction at 3 mo after PRF stimulation was found to be 52.4%. In the current study, we showed that PRF stimulation of the cervical medial branches may be used as an alternative treatment method in patients with CFP. PRF may alleviate CFP effectively and is safe to perform.

**ARTICLE HIGHLIGHTS**

***Research background***

Cervical facet joint pain (CFP) is one of the most common causes of neck pain and headache. Persistent CFP deteriorates the quality of life of patients and reduces their productivity at work.

***Research motivation***

In order to investigate the effectiveness of pulsed radiofrequency (PRF) stimulation of cervical medial branches in patients with chronic CFP.

***Research objectives***

The authors aim to investigate the effectiveness of PRF stimulation of cervical medial branches in patients with chronic CFP.

***Research methods***

The authors retrospectively included 21 consecutive patients (age = 50.9 ± 15.3 years, range 26-79 years; male: female = 8:13; pain duration = 7.7 ± 5.0 mo) with chronic CFP, defined as ≥ 4 on the numeric rating scale (NRS). The authors performed PRF stimulation on the cervical medial branches.

***Research results***

The outcomes of the PRF procedure were evaluated by comparing the NRS scores for CFP before treatment and 1 and 3 mo after treatment. Successful pain relief was defined as a ≥ 50% reduction in the NRS score at 3 mo when compared with the pretreatment NRS score.

***Research conclusions***

PRF stimulation of the cervical medial branches may be used as an alternative treatment method in patients with CFP. PRF can effectively alleviate CFP, and is safe to perform.

***Research perspectives***

PRF stimulation on cervical medial branches may be a useful therapeutic option to control chronic CFP.

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**Footnotes**

**Institutional review board statement:** The Institutional Review Board of Yeungnam university hospital approved this study, and the need for written informed consent was waived due to the retrospective design of the study.

**Conflict-of-interest statement:** The authors declare no conflict of interest.

**Informed consent statement:** This study was conducted retrospectively, and there was the need for written informed consent was waived.

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**Peer-review report’s scientific quality classification**

Grade A (Excellent): 0

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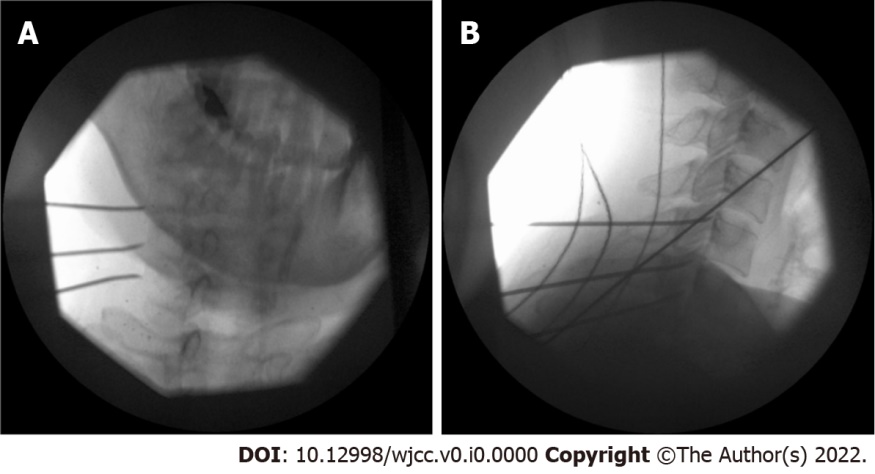
Grade C (Good): C

Grade D (Fair): 0

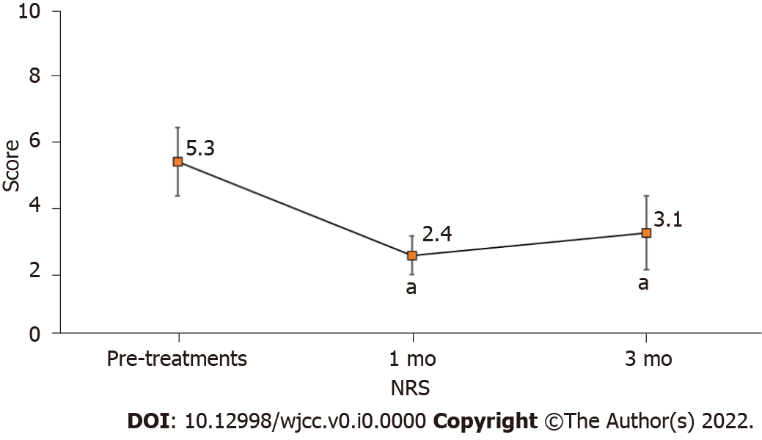
Grade E (Poor): 0

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**Figure Legends**

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**Figure 1 Under fluoroscopic guidance, the catheters were inserted for pulsed radiofrequency stimulation of the Lt. C5, 6, and 7 medial branches.** The catheter tips were placed around the Lt. C5, 6, and 7 cervical medial branches. A: posterior-anterior view; b: lateral view.



**Figure 2 Changes in numeric rating scale score for cervical facet joint pain over the assessment period.** The numeric rating scale scores significantly reduced from 5.3 prior to treatment to 2.4 at 1 mo, and 3.1 at 3 mo after pulsed radiofrequency treatment. a*P* < 0.05 indicate a significant result. NRS: numeric rating scale.

**Table 1 The cervical medial branches on which pulsed radiofrequency was applied**

|  |  |
| --- | --- |
| **Patient** | **Stimulated level** |
| 1 | Lt. C4, 5, 6 |
| 2 | Rt. C4, 5, 6 |
| 3 | Rt. TON, C3, 4, 5 |
| 4 | Both C3, 4, 5 |
| 5 | Rt. C3, 4 |
| 6 | Rt. TON, C3, 4 |
| 7 | Lt. C3, 4, 5 |
| 8 | Rt. C3, 4, 5 |
| 9 | Rt. TON, C3, 4, 5 |
| 10 | Lt. C3, 4, 5 |
| 6 | Rt. TON, C3, 4 |
| 7 | Lt. C3, 4, 5 |
| 8 | Rt. C3, 4, 5 |
| 9 | Rt. TON, C3, 4, 5 |
| 10 | Lt. C3, 4, 5 |
| 11 | Rt. TON, C3, 4, 5, 6 |
| 12 | Lt. C5, 6, 7 |

TON: third occipital nerve.

**Table 2 Global perceived effect according to a Likert scale**

|  |  |  |
| --- | --- | --- |
| **Score** | **% Change** | **Description** |
| 7 | ≥ 75 improvement | Very good |
| 6 | 50-74 improvement | Good |
| 5 | 25-49 improvement | Fairly good |
| 4 | 0-24 improvement or worse | Same as before |
| 3 | 25-49 worse | Fairly bad |
| 2 | 50-74 worse | Bad |
| 1 | ≥ 75 worse | Very bad |