Comparison of Two Methods of Epidural Steroid Injection in the Treatment of Recurrent Lumbar Disc Herniation

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Study Design: Prospective study.

Purpose: We compared the effects of two methods of epidural steroid injection in patients with recurrent disc herniation.

Overview of Literature: To our knowledge, there is no previous report of such a comparison in these patients.

Methods: The study was performed with 30 patients with relapsed lumbar disc herniation whose pain was not relieved by conservative remedies. The patients were divided into two groups, each of 15 patients, and entered the study for caudal or transforaminal injections. The degree of pain, ability to stand and walk, and the Prolo function score were evaluated in both groups before the injection and 2 months and 6 months after the injection.

Results: The degrees of pain reduction in the caudal injection group in the second and sixth months were 0.6 and 1.63, respectively, and in the transforaminal injection group were 1.33 and 1.56, respectively. The difference between the two methods was not statistically significant. Similarly, no other evaluated criterion showed a significant difference between the methods.

Conclusions: In the current study, the caudal and transforaminal steroid injection methods showed similar outcomes in the treatment of relapsed lumbar disc herniation. However, more detailed patient categorizing may help in finding possible subgroups with differences.

Keywords: Lumbar spine; Recurrent disc herniation; Epidural steroid; Caudal; Transforaminal

Introduction

Lumbar and leg pain are among the most common causes of disability in Western societies. Such disabilities may have several causes but lumbar disc herniation alone accounts for about 50% of cases [1]. Epidural steroid injection (ESI) is the most commonly used intervention in treatment of the pain of disc herniation [2-4]. Epidural steroid injections are performed in three ways: caudal, interlaminar, and transforaminal (TF). The TF method is favored because less drug is needed for the injection and because of the proximity of the injection site to the nerve root [5]. Furthermore, diffusion of the drug to the anterior part of the sac is more feasible [6,7]. The analgesic properties of the interlaminar method have been demonstrated in many studies [8]. Although some
side effects, such as paraplegia, increased blood pressure [8], and even death (Smuck), have been reported, the procedure is considered safe because these complications are very rare [8,9]. The caudal method has been demonstrated to have acceptable analgesic effects and is easier to perform than the other two methods [5].

After discectomy, however, symptoms may relapse in 5% to 40% of cases [10], and in cases of prolonged follow-up, the relapse rate will be even higher [10]. The main reasons are relapsed disc herniation and nerve root compression, epidural fibrosis, foraminal or lateral canal stenosis, segmental instability, facet joints, and myofascial pain syndromes [11-13]. The presence of adhesions will increase the probability of nerve root damage and rupture of the dura mater in a second operation [10]. Because a second operation has less favorable outcomes, alternative treatment methods may be more attractive [14].

Some researchers have studied the effects of epidural steroid injections in patients with relapsed disc herniation [15,16], but to our knowledge, there is no report comparing the effectiveness of the different methods of ESI in these patients. Adhesion at the site of a laminectomy increases the rate of dural puncture during an interlaminar epidural injection, so we sought to compare the transforaminal and caudal methods in this study.

Materials and Methods

The Medical Ethics Committee of the Kerman Neuroscience Research Center approved the study protocol. Those who entered the study received information about the side effects and probable complications. Written informed consent was obtained from each participant. The trial is registered in the Iranian Registry of Clinical Trials (IRCT138903084047N1).

Sampling was done during a 17-month period, from June 12, 2009 to November 22, 2010. The inclusion criteria were as follows: 1) a previous episode of successful lumbar open discectomy due to lumbar disc herniation, 2) recurrent radicular pain that was not responsive to at least 6 weeks of continuous treatment, 3) age of 18 or older, and 4) recurrent disc herniation at the same level, ipsilateral to the side of the previous operation, as verified by lumbar spine magnetic resonance imaging (MRI) (Fig. 1) [17-19].

We excluded patients with severe central disc herniation, based on MRI findings, significant or progressive neurologic deficits, patients who had undergone previous epidural steroid injections, patients with coagulation disorders or known allergies to analgesics or corticosteroids, and pregnant women.

The patients were randomly assigned to the caudal and transforaminal groups in a 1:1 ratio, using block randomization. All procedures were performed in Bahonar Hospital, which is supervised by Kerman University of Medical Sciences. Before they entered the operating room, 0.5–2 mg of midazolam had been injected intravenously. The patients were placed in a prone position and, after disinfection with betadine, the skin was anesthetized with 1% lidocaine.

In the transforaminal method, a 20-gauge LP needle was placed anterosuperiorly towards the foramen under fluoroscopy guidance (Fig. 2), and after confirmation of the correct positioning of the needle, 40 mg of depomedrol, 1 mL of bupivacaine, and 1 mL of 2% lidocaine were injected.

In the caudal method 40 mg of depomedrol, 1 mL of 2% lidocaine, and 1 mL bupivacaine 0.25% and 7 mL of distilled water (total, 10 mL) were injected via a 22-gauge needle from the sacral notch. After the injection, patients were observed in the hospital for 30–60 minutes and then discharged.

All patients underwent treatment with tizanidine 6–12 mg per day for 7 days to relieve muscle spasm, celecoxib...
100–200 mg per day for 7 days to relieve pain, and nor-triptyline 25–100 mg per day for 21 days. The severity of the pain sensed by the patients had been asked and registered according to the NPS table before the procedures.

The patients were then followed via phone at 2, 4, and 8 weeks and 6 months later [20]. After asking patients about their ability to do their daily affairs, using the Prolo scales table [21], their function was recorded. A walking tolerance test was performed according to the ability of the patients to walk without feeling pain, and patient responses were recorded using the ‘foot’ scale. A standing tolerance test was performed according to the ability of the patients to stand without feeling pain, and patient responses were recorded using the ‘minute’ scale. The number of rest days was recorded according to the days in which patients could not do their routine daily affairs because of back pain, and patient responses were recorded using the ‘day’ scale.

After descriptive statistical analyses were performed using the SPSS software (ver. 10.0), comparisons between the two groups were made using Student’s t-test for parametric and the Mann-Whitney U-test for non-parametric data. The χ² statistic was used to test differences in proportions and a paired t-test was used to compare the pre- and post-treatment results of average pain. Results were considered statistically significant if the p-value was less than 0.05.

Results

In the 17-month period, 32 patients entered the study. Two cases needed early operations because of neurological deficits. In total, 30 patients met our criteria with an average age of 48±10.51 years, 14 (46.6%) male and 16 (53.4%) female cases, who had undergone discectomies 18±6 months earlier. In 22 (73.3%) cases, prior surgeries had been performed at the level of the disc between L4 and L5, and in 6 (20%) cases, at the level of L5–S1. A foraminotomy had been performed in all 30 patients. These patients’ pain scores had been established 14.4±6.5 weeks earlier. The severity of pain, according to the NPS scale, was 6.23±1.48, the standing ability score was 17.52±13.70, and the walking ability score was 304±149.10 feet.

There was no significant difference in age, gender, level of involvement, or time passed since surgery between two groups. Patient details are provided in Table 1. Table 2 compares the severity of pain and other criteria evaluated in the patients.

Patients in the caudal group underwent ESIs via the caudal method and the transforaminal group via the transforaminal method. At 2 months after the injections,

![Fig. 2. The location of the needle in the foramina.](image-url)
the pain score in the first group reached 5.53±1.51, from 6.13±1.64, and in the second group reached 5.0±0.93, from 6.33±1.35. Thus, the mean pain reduction with the caudal method was 0.6 and 1.33 with the transforaminal method; this difference was not statistically significant (p=0.184).

Over the same time period, the patients’ tolerance to walking in the first group reached 249.26±129.0, from 342.20±166.8, and in the second group reached 267.4±169.23, from 273.33±131.40. Thus, during this period, walking tolerance decreased by 37.80 units in the caudal group and increased by 57.87 units in the transforaminal group; however, this difference was also not statistically significant (p=0.30).

The patients’ tolerance for standing in group caudal reached 20.17±17.53, from 22.57±18.27, and in group transforaminal reached 14.67±11.57, from 9.13±9.05. Thus, during this period, standing tolerance decreased by 0.80 units in the caudal group and increased by 5.54 units in transforaminal group; this difference was not statistically significant (p=0.10).

The mean rest days in the caudal group increased by 1.6 days and decreased by 1.11 days in the transforaminal group; this difference was also not significant (p=0.37). All criteria were then re-evaluated at the end of a 6-month period; these results are presented in Table 3.

A comparison between the groups according to the Prolo scoring system before treatment and at the end of the second and sixth months is shown in Table 4.

One patient suffered temporary paraparesis in group caudal. We did not see any other complication.

**Discussion**

Although inflammatory disorders are among the important causes of low back pain, inflammation has been shown to be present even with mechanical etiologies of low back pain. Many animal studies have revealed that some cytokines, such as tumor necrosis factor alpha, which are synthesized by macrophages in the nucleus pulposus, on reaching the nerve root, can start pathological changes that induce pain sensations and sensory deficits [5,22,23]. The role of inflammatory and chemical mediators in pain induction explains the success of steroid use in the treatment of low back pain and radicular pain caused by lumbar disc herniation [24,25].

Many studies have investigated the effects of the epidural steroid injections, but few have compared different methods of the injection in primary lumbar disc herniation. The superiority of the transforaminal method has been supported by some RCTs [8]. However, this remains controversial. Mendoza-Lattes et al. [26] compared the caudal method to the transforaminal method in patients with primary lumbar disc herniation. The transforaminal method showed better results, but the difference was not statistically significant. Despite the controversy, it is commonly accepted that the transforaminal injection has better results because of better distribution of the drug to
of our cases had facet joint involvement; this may explain our lower response rate. We also did not use fluoroscopy during caudal injections. Manchikanti et al. [30] reported that injection without fluoroscopy guidance was associated with an inappropriate site of the injection in 30% of cases, which can decrease the distribution of the drug to the site of the inflammation and nerve compression. This may also explain our lower response rate. Revel et al. [16] showed that a forceful caudal injection had a 49% response rate versus 19% in a routine injection; the difference was statistically significant. The results for the caudal method in our study resemble the results of the caudal injection without force in Revel et al.'s study. It is possible that if we had used a forceful injection, our results would have been better, but further investigations are required to assess this.

One limitation of our study is the absence of a control group. The presence of a control group could help in interpreting the results. However, our major goal was to compare the two methods of the epidural injection, not the effect of the epidural injection on relapsed disc herniation. Second, the presence of a control group could impair our randomization. It should not be forgotten that not all the causes of recurrent pain are related to disc herniation or epidural fibrosis and other etiologies, such as degeneration of the adjacent disc, facet joint involvement, and fibrosis of the nerve root can also be causes of relapse pain [11]. Among these etiologies, relapsed discs and canal stenosis respond partially to epidural injections [13,15]. Epidural fibrosis, which occurs in 20% to 36% of “failed back syndrome,” may respond to epidural injection, but the response is very short-term [13,15]. Using a single treatment remedy for a heterogeneous group of diseases with similar symptoms may explain treatment failure.

Although epidural steroids have a short-term effect, their use along with other modalities of treatment may help relieve pain and enhance patient function. It would seem that the relatively low response rate could be improved by better categorizing patients and enhancing the injection methods. However, generally increased knowledge about the pattern of epidural adhesion may also help in choosing the most appropriate injection method.

**Conclusions**

Epidural steroid injection via the caudal and transforaminal epidural injection to evaluate the distribution of the drug. Attempts to quantify the amount and location of epidural fibrosis after lumbar discectomy may help in choosing the most appropriate method for the epidural injection.

Manchikanti et al. [30] showed that the caudal injection decreased chronic low back pain significantly and radicular pain in 65% of the patients with recurrent disc herniation, which was higher than our result. In that study, cases with facet joint involvement were excluded. It has been reported that pain originates from the facet joints in 17% to 24% of the patients with FBSS, and it is recommended to perform a facet block test to diagnose facet joint involvement [13]. However, this test is not very sensitive or specific. We did not perform the test, and probably some the anterior part of the sac [6,7].

In our study, the transforaminal epidural injection did have better results than to the caudal injection method; however, none of the differences were statistically significant. Although different studies have reported up to 84% efficacy for the transforaminal injection method in the treatment of the primary lumbar disc herniation [27], our response rate with the transforaminal method was only 40%, considerably lower than some of those studies. Buenaventura et al. [8] also reported a low response rate after transforaminal steroid injection in patients with recurrent disc herniation, which may have been due to epidural fibrosis. A peridural scar is capable of binding the dura and nerve roots to the circumferentially surrounding structures and in animal models of post-lumbar laminectionomy nerve root adherence to the underlying disc and pedicle has been demonstrated [28]. The pattern of epidural fibrosis in humans is not well known; however, it is supposed that fibrosis would appear at the junction of the dura mater with a hematoma [28]. In our study, all the patients underwent a foraminotomy in their previous operation. It is possible that adhesion in the foramina, which was already compromised by the herniated disc, restricted the distribution of the drug to the anterior part of the sac through the foramina. However, it is known that innervation to the annulus and its peripheral ligaments is multi-segmental, and in some studies it has even been stated that pain transferring via sympathetic routes will not be relieved by nerve block below the level of L2 [29]. This mechanism may make the transforaminal method more vulnerable to fibrosis and adhesion.

We did not use a contrast material during the transforaminal epidural injection to evaluate the distribution of the steroid. Attempts to quantify the amount and location of epidural fibrosis after lumbar discectomy may help in choosing the most appropriate method for the epidural injection.
nal methods in patients with relapsed disc herniation showed similar effects in terms of pain and patient function. Better categorizing of patients may help in achieving better outcomes.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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