

Review Article

Prognostic Factors Affecting Ultrasound Guided Caudal Epidural Injection in Sciatica Patients

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Introduction

Sciatica results from spinal nerve root compression and produces pain in a dermatomal distribution. The pain is often lancinating shooting sharp in quality. It is frequently accompanied by numbness and tingling and may be associated with sensory or motor deficits. This should be differentiated from non -neurogenic sclerotomal pain [1].

The most common cause of sciatica is herniated intervertebral disc [2]. The herniated disc can cause nerve root impingement that leads to lumbosacral radiculopathy [2]. This is considered the mechanical component of sciatica. While there is in addition a biologic chemical component, including inflammation, vascular invasion, immune

responses and an array of cytokines [2].

Epidural corticosteroid injection is used mostly in subacute (>6 weeks) and chronic low back pain. It had gained popularity and the rationale beyond it that the genesis of radicular pain when a herniated disc impinges on a nerve root, is at least partly related to locally induced inflammation [3,4]. Caudal epidural injection is performed as diagnostic and therapeutic interventions in various lumbosacral pain syndromes. Caudal epidural injections is complicated by variations in sacral anatomy and the risk of inadvertent intravascular injection. Ultrasound guided injection was found to be as effective as fluoroscopic guidance without the risk of radiation exposure. However, the ultrasound-guided techniques are limited by lack of visualization inside the sacral canal thereby limiting the identification

of epidural spread and vascular spread [5].

Methods and Materials

This study was prospective study conducted at outpatient setting. All the included patients had signed informed consent prior to their participation after explanation of the benefits and risks of caudal epidural injection. The study included 320 patients with chronic back pain with sciatica more than 3 months exclusion criteria: severe motor weakness, previous back surgery, infection at site of injection

Intervention: all patients had detailed history taking, Body Mass Index (BMI), neurological & musckelo-skeletal examination, MRI lumbosacral spine, Nerve conduction-electromyography. Followed by ultrasound guided caudal epidural injection.

Technique of ultrasound guided caudal epidural

The patient was placed in prone position with abdomen resting on a pillow to relax the gluteal muscles, the patient was asked to turn his heels outward. Ethyl chloride was sprayed as local anesthetic for the whole sacral area after the skin overlying the sacrum and sacral hiatus was prepped with antiseptic solution. A curved ultrasound transducer was place over the lower sacrum after the application of a sterile gel. The transducer was placed in a transverse plane and slowly moved caudally until the sacral cornua are visualized. Sacral hiatus and sacrococcygeal ligaments are identified. The transducer then turned longitudinally and moved slowly cephalad until the inferior portion of the ultrasound transducer lies toward the top of the sacral hiatus. A 22-gauge 3-inch needle was inserted through the skin 1 cm below the inferior border of the transducer utilizing in plane approach and advanced with a 45-degree angle to skin through sacrococcygeal ligament in the caudal canal. After a negative aspiration, 40mg of triamcinolone together with 4 ml of lidocaine 1% was injected (1).

Main outcome measure

BMI, back pain by Visual Analogue Scale (VAS), Oswestry Disability Index before and 4weeks after injection [7,8].

Results

The demographic and clinical data of the studied patients. Mean age 56.2±12.74, 116 male (36.3%), 204 female (63.7%). Mean BMI 34.3±8.01, obese category was the highest (Table 1). The duration of pain varies from less than 1 year to more than ten years with the highest number of patients with duration 4-10 y. MRI findings of the studied patients (Table 2). The most frequent findings were L 4-L5 disc. MRI with one disc 64.1%, two disc 27.7%, three or more 11.3%. NCS-EMG finding revealed chronic radiculopathy; L4; 10.3%, L5; 45.9%, S1; 48.4%, bilateral radiculopathy 17.2%, axonal polyneuropathy 4.7%.

The change of VAS and Oswestry disability after caudal epidural injection. Both of them showed significant improvement (p=0.001) (Table 3). The Degree of VAS improvement and Oswestry disability index improvement after injection. Most of the studied patients falls among 50-75% category of improvement (Table 4). The number of injections received Most of the studied patients received only one injection (Table 5). The duration between injection (Table 6). Mean duration between injection; 2.82±2.01 months.

Factors significantly affect VAS improvement; Age p=0.013, BMI

Table 1: Distribution of the studied patients group regarding their demographic and basic clinical data.

| | Number "n= 320" | Percent |
|-------------------------------|--------------------|---------|
| Age | | |
| <40 | 32 | 10 |
| 40-60 | 145 | 45.3 |
| 60* | 143 | 44.7 |
| Range | 23.0-84 | |
| Mean | 56.2 | |
| S.D | 12.74 | |
| Sex | | |
| Male | 116 | 36.3 |
| Female | 204 | 63.7 |
| BMI | | |
| Normal | 55 | 17.2 |
| Over weight | 120 | 37.5 |
| Obese | 145 | 45.3 |
| Range | 21.0-55.3 | |
| Mean | 34.3 | |
| S.D | 8.01 | |
| History of diabetes | 77 | 24.1 |
| Pain duration | | |
| <1 year | 72 | 22.5 |
| 3-Jan | 95 | 29.7 |
| 10-Apr | 114 | 35.6 |
| >10 | 39 | 12.2 |
| Side of sciatica | | |
| Right | 195 | 60.9 |
| Left | 94 | 29.4 |
| Bilateral | 31 | 9.7 |
| Objective Sensory Exam | | |
| Positive | 209 | 65.3 |
| Negative | 111 | 34.7 |
| Objective Motor Exam | | |
| Positive | 70 | 21.9 |
| Negative | 250 | 87.1 |

Table 2: MRI findings among the studied patients.

| MRI findings | Number "n= 320" | Percent |
|--------------|--------------------|---------|
| L3-L4 Disc | 64 | 20.0 |
| L4-L5 | 80 | 25.1 |
| L5-S1 | 75 | 23.4 |
| Facet | 25 | 7.8 |
| S. Stenosis | 52 | 16.3 |

Table 3: Comparison between pre and post-injection VAS and Oswestry Disability Index.

| | Pre Injection | Post- Injection | t-test | P value |
|----------------------------------|---------------|--------------------|--------|---------|
| VAS | | | | |
| Range | 5.0-10.0 | 0-10 | 8.22 | 0.001* |
| Mean | 8.75 | 4.20 | | |
| S.D | 1.36 | 2.34 | | |
| Oswestry Disability Index | | | | |
| Range | 60-93.0 | 8.0-75.0 | 5.98 | 0.001* |
| Mean | 77.4 | 40.1 | | |
| S.D | 8.9 | 12.2 | | |

p=0.022, diabetes p=0.003, objective positive sensory & motor exam p=0.046, 0.04 respectively, presence of more than one- disc p=0.0021, spinal stenosis p=0.003 (Table 6).

Discussion

This study was carried out to evaluate the efficacy of ultrasound-guided caudal epidural injection among sciatica patients and the factors that affect the degree of improvement. All the studied patients

Table 4: Distribution of the studied patients regarding the degree of improvement of VAS and Oswestry Disability Index.

| Degree of Improvement | Number "n= 320" | Percent |
|----------------------------------|-----------------|---------|
| VAS | | |
| >75.0% | 65 | 20.3 |
| 50.0-75.0% | 122 | 38.1 |
| <50.0% | 81 | 25.3 |
| No change | 52 | 16.3 |
| Oswestry disability index | | |
| >75.0% | 85 | 26.6 |
| 50.0-75.0% | 124 | 38.8 |
| <50.0% | 72 | 22.5 |
| No change | 39 | 12.2 |

Table 5: Number of injection received among the studied patients group.

| Number of injection received | Number "n= 320" | Percent |
|------------------------------|-----------------|---------|
| 1 | 252 | 78.8 |
| 2 | 46 | 14.4 |
| 3 | 4 | 1.3 |
| 4 | 16 | 5.0 |
| 6 | 2 | 0.6 |

Table 6: Duration between each injection received among the studied patients group.

| Duration between each injection (months) | Number "n=320" | Percent |
|--|----------------|---------|
| 1 month | 12 | 3.8 |
| 2-3 months | 165 | 51.6 |
| 4-5 months | 109 | 34.1 |
| 6* | 34 | 10.6 |
| Range | 1-9 | |
| Mean | 2.82 | |
| S.D | 2.01 | |

presented with sciatica more than three months and had responded significantly to corticosteroid caudal epidural injection with variable degree of improvement of pain score and Qwstery disability index based on the tested variables.

Revising literature regarding the efficacy of caudal epidural injection in the management of sciatica, Watts & Silagy [9] carried a meta-analysis on the efficacy of epidural corticosteroid in the treatment of sciatica and documented that it is very effective in the management of lumbosacral radicular pain [9]. In addition, other authors reported the significant efficacy of caudal epidural in treating sciatica patients [10,11].

Nandi J and Chowdhery A studied 47 patients with sciatica receiving caudal epidural corticosteroid injection, In comparison to placebo, there was significant improvement after 4 weeks but at 12 weeks, there was no difference between groups and they concluded that caudal epidural provide no additional improvement over placebo in long term natural history of lumbosacral sciatica, however it can be an important component of short term management of painful sciatica [12]. In Contradiction to our results, Iversen T et al., concluded in their study that no difference between caudal steroid or saline injection in treating chronic lumbar radiculopathy. Their study compared injection between saline and steroid injection and they found short-term improvement for both groups but on long-term basis after 52 weeks, follow up there was no improvement. Each

of the tested patient groups was composed of 41 patient's only [13]. In our study, we had large population sample that can lead to more accurate statistical results and we had measured VAS and Owstery disability index after 4 weeks i.e. short term period as we believe that it is unlikely that the improvement of caudal steroid injection will persist forever so to test the patient after 52 weeks, in our opinion it is very long periods that is unlikely the effect of steroid will be persistent during this whole period.

In this study, the degree of improvement of VAS and Owstery disability index after caudal epidural injection were variable and the highest category falls among 50-75% improvement. In the previous literature, it was documented the improvement of VAS and Owstery index, however the degree of improvement was not listed in the literature [14].

In this study, we had studied the variable factors that can influence the outcome of the caudal epidural injection, namely the presence of DM, BMI, age and presence of objective sensory and motor findings, MRI findings.

As regard the presence of DM. It significantly negatively influenced the degree of improvement of VAS after epidural injection. Although the diabetic patients still showed significant improvement of pain compared to prior injection but the degree of improvement is significantly less compared to non-diabetic patients.

Diabetes mellitus is associated with low back pain and spinal pain, however direct causal link between diabetes and back pain was not established [15]. The association with chronic back pain are more stronger for severe cases of pain.

The association of DM with the severity of pain and the frequency of its chronisation and recurrence has been established. The most likely mechanism of such association is the lesion of intervertebral discs mediated by the accumulation of advanced glycation end products (EGP). In DM the concentration of EGP increases significantly, they initiate ectopic calcification, a decrease in cell density in the end plate and changes in vertebrae. Cells of pulposus nuclei begin to produce pro inflammatory cytokines and chemokines that trigger the process of angio and neurogenesis [16].

Won Ho Kim et al reported significant improvement of pain after epidural injection for diabetic patients using either 20 or 40 mg triamcinolone without significant difference between doses. However, in their study they did not compare the results with none diabetic patients to see the influence of diabetes on the degree of pain improvement [17]. In this study, the majority of the studied patients falls among the obese body mass index. All of the studied patients including the obese category have significant improvement of VAS and Owstery disability index which denotes the effectiveness of ultrasound guided epidural injection even among the obese patients, but when we compared the degree of improvement of VAS between the obese and non- obese patients, we found that obesity inversely affected the degree of improvement.

Conducted a study aiming to find the association between caudal epidural steroid injection and BMI. They concluded that caudal epidural injection improved all body weight and they noted that the limitation of the study was due to small number of obese patients [18].

Table 7: Multivariate analysis of different risk factors, which may affect the degree of improvement by VAS.

| | Improvement regarding VAS | | | | | | | | Total | P value |
|-------------------------------|---------------------------|------|--------|------|------|------|-----------|-------|-------|---------|
| | >75% | | 50-75% | | <50% | | No change | | | |
| | No. | % | No. | % | No. | % | No. | % | | |
| Age | | | | | | | | | | |
| <40 | 22 | 68.8 | 6 | 18.8 | 2 | 6.3 | 2 | 6.3 | 32 | 0.013* |
| 40-60 | 32 | 22.1 | 63 | 43.4 | 50 | 34.5 | 0 | 0.0 | 145 | |
| 60+ | 11 | 7.7 | 53 | 37.1 | 29 | 20.3 | 50 | 35.0 | 143 | |
| BMI | | | | | | | | | | |
| Normal | 32 | 58.2 | 18 | 32.7 | 3 | 5.5 | 2 | 3.6 | 55 | 0.022* |
| Over weight | 20 | 16.7 | 69 | 57.5 | 15 | 12.5 | 16 | 13.3 | 120 | |
| Obese | 13 | 9.0 | 35 | 24.1 | 63 | 43.4 | 34 | 23.4 | 145 | |
| History of diabetes | 6 | 7.8 | 12 | 15.6 | 19 | 24.7 | 40 | 51.9 | 77 | 0.003* |
| Pain duration | | | | | | | | | | |
| <1 year | 45 | 62.5 | 12 | 16.7 | 10 | 13.9 | 5 | 6.9 | 72 | 0.069 |
| 1-3 yrs | 10 | 10.5 | 52 | 54.7 | 30 | 31.6 | 3 | 3.2 | 95 | |
| 4-10 yrs | 6 | 5.3 | 50 | 43.9 | 34 | 29.8 | 24 | 21.1 | 114 | |
| >10 yrs | 4 | 10.3 | 8 | 20.5 | 7 | 17.9 | 20 | 51.3 | 39 | |
| Objective Sensory Exam | | | | | | | | | | |
| Positive | 2 | 1.0 | 20 | 9.6 | 62 | 29.7 | 209 | 100.0 | 209 | 0.046* |
| Negative | 63 | 56.8 | 102 | 91.9 | 19 | 17.1 | 27 | 24.3 | 111 | |
| Objective Motor Exam | | | | | | | | | | |
| Positive | 4 | 5.7 | 13 | 18.6 | 18 | 25.7 | 35 | 50.0 | 70 | 0.040* |
| Negative | 61 | 24.4 | 109 | 43.6 | 63 | 25.2 | 17 | 6.8 | 250 | |
| Number of disk | | | | | | | | | | |
| One | 62 | 30.2 | 52 | 25.4 | 42 | 20.5 | 49 | 23.9 | 205 | 0.0021* |
| Two | 3 | 3.8 | 52 | 65.8 | 23 | 29.1 | 1 | 1.3 | 79 | |
| Three or more | 0 | 0.0 | 18 | 50.0 | 16 | 44.4 | 2 | 5.6 | 36 | |
| S. Stenosis | 3 | 5.8 | 16 | 30.8 | 9 | 17.3 | 24 | 46.2 | 52 | 0.003* |

Excess body weight causes extra stress on the disc. Disc is a soft rubbery pad between the vertebrae, which carries the body weight. When the disc herniates, the nucleus protrudes and presses the nerve through the spinal canal [19].

Baysal and Friends compared the epidural steroid injection between the obese and non-obese patients. They did not experience any difference between the groups [20].

Klocke in another study stated that ultrasound guided epidural injection is safe and effective in over weight and obese patients [21]. In this study, the majority of the patients are from 40-60 y old age group. All the studied patients showed significant improvement of VAS and Oswestry disability index. However, older age, above 60 y showed significantly less degree of improvement in VAS.

The frequency of disc herniation increases with age. The peak frequency of herniation at L5-S1, L4-L5 levels is between the age of 44 & 50 year with a progressive decline in frequency thereafter [22]. In our study, most of the patients lie within the above-mentioned category of patients i.e. 40-60 y, which goes with the reported literature. However, the explanation why the older age achieved significantly less degree of improvement of pain after caudal epidural was not mentioned. May be because with older age, the pathology of degenerative disc disease become more advanced and long-standing, which can influence the degree of pain relief after epidural injection.

Among the variables, that we studied whether it affects the degree of improvement of VAS after caudal epidural was the presence of sensory and/or motor deficit. We found that the presence of such deficit negatively significantly affected the degree of VAS improvement. The suggested explanation for this that the presence of either sensory and or motor deficit indicate more grave compression

on the nerve root with more permanent effect. In contradiction to this result, Billy GG et al., denied the effect of sensory and or deficit on the results of epidural [23].

We disagree with this result because during the natural course of spinal nerve root entrapment, first stage there is only sensory complaint presenting the sciatica with no objective clinical findings, the more the compression on the spinal nerve root is persistent, then first the sensory fibers of the nerve root become affected manifested clinically as an objective sensory deficit. Further compression will affect the motor component of the nerve root and then will be manifested clinically as motor deficit so this consequence reflects the stages and the degree of nerve root compression so our findings that proves that the presence of sensory and or motor deficit affects the degree of improvement and response to caudal epidural injection although they still got significant improvement compared to their baseline.

In our study we had tried to analyze the degree of improvement of VAS and Oswestry disability index after caudal epidural injection with MRI findings, we found that that presence than more than one herniated disc negatively affected the degree of improvement of VAS and Oswestry disability index although those group still showed significant improvement compared to their base line. In addition, those patients with spinal stenosis in MRI findings showed less significant improvement compared to patient with only one herniated disc but still they experienced significant improvement of pain and disability score compared to their base line.

Revising the literature for these particular findings, we did not find a study that specifically analyze the degree of improvement after caudal epidural with lumbosacral MRI. But almost all the studies documented the general improvement after caudal epidural

injection as mentioned before. Carried out meta-analysis for efficacy of epidural injections in managing [24].

Chronic spinal pain and reported that the evidence is level II for caudal and lumbar interlaminar epidural injection with level III evidence for lumbar transforaminal epidural for lumbar spinal stenosis. The evidence is level III for axonal or discogenic pain without facet arthropathy [24].

In Conclusion; ultrasound guided caudal epidural injection is very effective and safe tool for short-term pain relief in patients with chronic back pain associated with sciatica without any obvious side effects. The degree of improvement of pain and function is negatively influenced by the age, body mass index, diabetes mellitus, the presence of sensory and or motor deficit as well as the presence of multiple herniated disc and spinal stenosis in lumbosacral MRI. However, patients having those variables still had a significant improvement of their symptoms compared to their base line.

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