Evaluation of the L2 Spinal Nerve Root Infiltration as a Diagnostic Tool for Discogenic Low Back Pain

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Background: To assess whether unilateral L2 infiltration with local anesthetic can be used to identify patients who will have negative discograms and thus eliminate the need for the discogram. Discogenic low-back pain is considered to have afferent pathways in the sinuvertebral nerves, mainly originating from the ventral rami of the spinal nerves. There is evidence that pain arising from the lower lumbar intervertebral discs may be transmitted through the sympathetic afferent fibers contained in the L2 spinal nerve root. Provocative discography, within the context of other clinical data, is the current "gold standard" by which to diagnose discogenic low-back pain, but a far more invasive procedure than L2 infiltration.

Objective: To evaluate the correlation between unilateral second lumbar (L2) spinal nerve root infiltration with local anesthetic and provocative discography in patients with chronic low back pain.

Study Design: A prospective, observational study.

Methods: All patients scheduled for discography were asked to participate in having local anesthetic infiltration of the L2 spinal nerve root at least two weeks prior to discography, until forty subjects were enrolled. Discography was performed after the patient's pain level returned to baseline.

Results: Local anesthetic infiltration of the L2 spinal nerve root was predictive of provocative discography results in only 46.5% of the subjects (26% true positives, and 20.5% true negatives). In 53.5% of the subjects, L2 infiltration was not predictive of discography results (20.5% false positives, and 33% false negatives).

Conclusions: The results showed that unilateral L2 infiltration is not predictive of discogenic low-back pain when compared to discography, the current "gold-standard" for diagnosis.

Keywords: L2 spinal nerve infiltration, discogentic low back pain, provocative discography, sympathetic afferent pathways, selective nerve root block, sinuvertebral nerve.
grams secondary to having a non-disco- genic etiology of their low back pain. This would eliminate unnecessary diagnostic provocative discography in those patients, thereby reducing the risks and procedural related pain. The principle risk inherent with performing discography is the exacerbation of existing pain. Other risks of discography include bleeding, infection, discitis, damage to surrounding structures, and allergic reaction to agents used in performing the procedure. Discography would still be necessary in those patients with a pain relieving response to L2 nerve block, in order to identify and better characterize the involved disc level(s).

**METHODS**

The study was undertaken in the Pain Medicine Clinic of the Naval Medical Center Portsmouth, VA. The protocol was approved by the Institutional Review Board at Naval Medical Center, Portsmouth, VA. The design consisted of a prospective, observational evaluation.

**Informed Consent**

All patients were provided with the approved protocol and informed consent document, approved by the Institutional Review Board for this study. The informed consent document described the details of the evaluation.

**Inclusion and Exclusion Criteria**

All the patients who participated in the study were identified from the referral pool of the patients being referred to the Pain Medicine Clinic, Naval Medical Center. All patients had received diagnostic and therapeutic measures to determine the source of pain or provide relief, from orthopedic surgeons and neurosurgeons with failure to respond. All the patients were referred for provocative discography with findings of degenerative disc disease on MRI and failed conservative treatment including a trial of epidural steroid injections and medial branch blocks.

**Inclusion Criteria**

Patients included in the study had a history of chronic low back pain greater than six months, were between the ages of 18 and 65, were eligible for care, and able to give proper informed consent. All patients were without significant co-morbid disease, including overt psychosocial disease. When diagnostic and therapeutic measures failed to determine the source of pain or provide relief, orthopedists and neurosurgeons referred the patients for provocative discography. All subjects had magnetic resonance imaging (MRI) findings suggestive of degenerative disc disease and had failed conservative treatment including a trial of epidural steroid injection and medial branch blocks.

**Exclusion Criteria**

Patients with significant co-morbid disease, overt psychological disease, pregnant or lactating women, coagulopathy, inability to understand informed consent and protocol, and patients unwilling to participate in the study were excluded.

**Evaluation**

Evaluation consisted of collection of demographic data, routine physical and medical evaluation, information on previous treatments, and pain assessment.

**Study Design and Investigation**

A single group of 40 patients received both L2 local anesthetic nerve root infiltration and discography. Discography was performed at a minimum of 2 weeks after L2 nerve root infiltration, but more importantly, only after the patient’s pain returned to baseline, which in no subject was greater than 8-weeks. Patients maintained a pain diary recording their pain by the Verbal Numeric Score (VNS) at 48 hours after L2 block and then weekly. Pain Clinic nurses called the patients weekly to ascertain each patient’s pain level and record the score.

The L2 nerve root infiltration was carried out on the patient’s predominantly painful side in the prone position. The skin was anesthetized with 1% Lidocaine (Xylocaine-MPF 1% AstraZeneca, Wilmington, DE). A 25g Quinke Spinal needle (Kimberly-Clark spinal tray, Roswell, GA) was advanced to the L2 foramen under fluoroscopic guidance (OEC Series 9600, GE Medical Systems, Raleigh, NC). Proper needle placement was confirmed in both anteroposterior and lateral views before and after injecting 1 mL of contrast (Omnipaque 300, Amersham, Princeton, NJ) to outline the L2 nerve root. The nerve root was then anesthetized by injecting 1.5 mL of 2% Lidocaine (Lidocaine-PF 2%, Abbott Labs, Chicago, IL). Pain relief was assessed by the VNS. The patient’s pain score was recorded immediately preceding L2 infiltration and again 20 minutes after the procedure. Pain relief was considered positive if there was at least a 50% decrease in VNS, otherwise negative.

The provocative discography procedure was initiated opposite the patient’s predominantly painful side in the prone position, in accordance with technique described elsewhere (4, 18). The skin was prepped with Betadine solution and drapes applied utilizing sterile technique. Sedation was performed with aliquots of intravenous Propofol (Diprivan 1%, AstraZeneca, Wilmington, DE). Skin was anesthetized with 1% Lidocaine. Under fluoroscopic guidance, 25g, 6” Quinke spinal needles were placed through 20g, 3.5” introducer needles (Nerve Root Block Kit, Kimberly-Clark, Roswell, GA) into the disc spaces that had corresponding degenerative changes on MRI. Once
Table 1. Demographic characteristics and L2 infiltration versus discography

<table>
<thead>
<tr>
<th>Gender</th>
<th>Male</th>
<th>Female</th>
<th>85% (34)</th>
<th>15% (6)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Race</td>
<td>African American</td>
<td>22% (9)</td>
<td>Caucasian</td>
<td>73% (29)</td>
</tr>
<tr>
<td></td>
<td>Hispanic</td>
<td>5% (2)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td>Mean ± SD</td>
<td>38 ± 8.7</td>
<td></td>
<td></td>
</tr>
<tr>
<td>VAS pain score</td>
<td>Baseline</td>
<td>Mean ± SD</td>
<td>5.7 ± 1.73</td>
<td></td>
</tr>
<tr>
<td></td>
<td>20 Minutes after L2</td>
<td>Mean ± SD</td>
<td>3.7* ± 1.93</td>
<td></td>
</tr>
<tr>
<td>L2 infiltration</td>
<td>Positive</td>
<td>47% (29)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Negative</td>
<td>53% (21)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Discogenic</td>
<td>Positive</td>
<td>59% (23)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Negative</td>
<td>41% (16)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Number of levels positive</td>
<td>Zero</td>
<td>41% (16)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>One</td>
<td>33% (13)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Two</td>
<td>23% (9)</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Three</td>
<td>3% (1)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* Indicates significant difference with Baseline

The patient was clearly no longer sedated, provocative discography was commenced by slowly injecting up to 1.5 mL of contrast (Omnipaque 300) into the disc spaces under fluoroscopy, utilizing a 20cc syringe with threaded plunger and manometric capability (Universal Fluid Dispensing Syringe, Merit Medical Systems, South Jordan, UT).

Each disc was interrogated sequentially by a principle investigator, with notation made of 1) opening pressure, in pounds per square inch, 2) whether or not pain was produced, 3) pressure (over opening) which caused pain, 4) concordance of the pain compared to the patient’s typical pain, 5) severity of pain produced. The test was considered negative if the patient’s typical back pain was not reproduced, and positive if all of the patient’s pain, or a component of their pain in cases of multilevel involvement, was reproduced with any amount of contrast less than or equal to 1.5 mL and at a change in pressure of less than 50 pounds per square inch over the opening pressure, at any level tested. Antibiotics were administered peri-procedurally as infection prophylaxis, especially intradiscal. Ancef (Cefazolin, Glaxo Smith Kline, Research Triangle Park, NC) 1 g iv administered as a slow infusion 30 minutes prior to the procedure. Ancef in a concentration of 0.5 mg/mL was included in the contrast. Postprocedurally, Tequin (Gatifloxacin, Bristol-Myers Squibb, Princeton, NJ) 400 mg was taken enterally once a day for five days.

Statistical Methods

The SPSS version 9.0 statistical packages were used to generate the frequency tables and chi-squared statistic was used. A paired t-test was used to compare baseline and post VAS pain scores for individual patients. Results were considered statistically significant if the P value was less than 0.05.

RESULTS

The patient population was homogeneous with regard to age and co-morbidities. The patient population was, however, predominantly male (34 of 40), reflecting the predominantly male population in the Armed Services (Table 1). One patient withdrew from the study after undergoing L2 infiltration, but before discography, when he moved out of the area. There were no complications of either provocative discography or L2 infiltration. Measured outcomes of the L2 spinal nerve root infiltration and subsequent provocative discography are displayed by subject number in Table 1. Chi square analysis, as depicted in Table 2, was essentially random, and thereby demonstrates no correlation.

DISCUSSION

In contrast to the results of Nakamura et al (15) the results of this study do not support the effectiveness of unilateral L2 infiltration as a therapeutic modality in the relief of discogenic low-back pain. Nor do the results support the use of unilateral L2 infiltration as a diagnostic tool in the evaluation of discogenic low-back pain.

Ohtori et al (19) published an animal study a few years after the Nakamura et al study, in the same lab, which may help explain our study’s results. They found that the posterior portion of the lumbar discs in rats were innervated by two distinct pathways: segmentally by the sinusvertebral nerves, and by non-segmental nerve fibers through the paravertebral sympathetic trunks. High variability in the anatomic innervation of the human disc may explain the variability of response to unilateral L2 nerve block when discogenic pain is present as proven by discography.

Of note, the current study took over two and a half years to collect 40 patients. During that time, there were seven different investigators at different levels of experience, some fellows in training, some staff pain physicians, performing the procedures and collecting the data. That may have allowed for some variation in technique and interpretation of clinical results.

The current study establishes the need for further research to determine more precisely the pathophysiology of discogenic low-back pain, as well as the continued need to develop conservative and cost effective means of diagnosing and treating discogenic low back pain. The authors are considering repeating the
study employing bilateral L2 infiltration, rather than only unilateral infiltration. Many patients report bilateral low-back pain. Additionally, there is the possibility that there may be some right-left crossover in the innervation of the posterior lumbar disc.

**CONCLUSION**

Unilateral L2 infiltration is not predictive of discogenic low-back pain when compared to discography, the current “gold-standard” for diagnosis. Whether this same conclusion would hold true utilizing bilateral L2 segmental nerve block is unknown.

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the above listed authors were employees of the Federal Government at the time the research was conducted, specifically the Naval Branch of the Department of Defense. This work was prepared as part of the authors' official duties. Title 17 U.S.C. 105 provides that "Copyright protection under this title is not available for any work of the United States Government." Title 17 U.S.C. 101 defines a United States Government work as a work prepared by military service members as part of their official duties.

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**REFERENCES**

16. Suseki K, Takahashi Y, Takahashi K, Chiba T, Yamagata M, Moriya H. Sensory nerve fibers from lumbar intervertebral discs...


