Lumbar Epidural Injections



DEFINING APPROPRIATE COVERAGE POSITIONS



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Introduction

North American Spine Society (NASS) coverage policy recommendations are intended to assist payers and members by proactively defining appropriate coverage positions. Historically, NASS has provided comment on payer coverage policy upon request. However, in considering coverage policies received by the organization, NASS believes proactively examining medical evidence and recommending credible and reasonable positions may be to the benefit of both payers and members in helping achieve consensus on coverage before it becomes a matter of controversy.

Methodology

The coverage policies put forth by NASS use an evidence-based approach to spinal care when possible. In the absence of strict evidence-based criteria, policies reflect the multidisciplinary and non-conflicted experience and expertise of the authors in order to reflect reasonable standard practice indications in the United States.

NASS Coverage Policy Methodology

Background Information

Lumbar epidural steroid injections can be performed via an interlaminar or caudal approach (CPT code 62311) or a transforaminal approach that includes the use of fluoroscopic or CT-guidance, which is bundled into the procedure (CPT codes 64483-64484). Fluoroscopic-guidance (CPT code 77003) is not bundled into CPT code 62311 and can be billed separately when performed with an interlaminar epidural steroid injection. Interlaminar and transforaminal epidural steroid injections using ultrasound guidance (CPT codes 0030T-0031T) are not recommended for coverage by NASS.

Scope and Clinical Indications

Therapeutic lumbar epidural steroid injections (ESIs) are indicated for the following diagnoses with qualifying criteria, when appropriate.

- 1. Lumbar radicular pain in which the following criteria are met:
 - a. the pain is severe enough to cause some degree of functional deficit
 - b. failure of at least four weeks of noninvasive care (see below*)
 - c. imaging demonstrating a correlative region of nerve impingement
- 2. **Neurogenic claudication** in which the following criteria are met:
 - a. the pain is severe enough to cause some degree of functional deficit

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- b. failure of at least four weeks of noninvasive care (see below*)
- c. imaging demonstrating a correlative region of nerve impingement
- 3. Low back pain without lower extremities symptoms ONLY in the following clinical scenarios:
 - a. High-level athletes during a competitive season
 - b. Pregnant women with intractable low back pain unresponsive to other treatments

*It is known that the majority of back and radicular pain will improve over 4 weeks. It is therefore reasonable to recommend failure of four weeks of non-surgical, noninvasive care. Appropriate non-surgical, non-injection treatments should be considered along with a rationale for interventional treatment. Exceptions to waiting 4 weeks can exist but should be carefully documented and should be reviewed on a case-by-case basis. These include, but are not limited to:

- a. At least moderate pain with significant functional loss at work and/or home
- b. Severe pain unresponsive to outpatient medical management
- c. Inability to tolerate non-surgical, non-injection care due to co-existing medical condition(s) (e.g. cardiac disease)
- d. Prior successful ESI for the same condition

Diagnostic selective nerve root blocks (DSNRBs) use a small amount of anesthetic via a **transforaminal approach** to anesthetize a specific spinal nerve and share the same CPT codes as therapeutic transforaminal ESIs (64479-64484). DSNRBs are used to evaluate a patient's anatomical level and/or source of radicular pain and are often used in surgical planning and decision-making. The following must be documented:

• Post-injection assessment of the percentage of pain relief and/or change in visual or numerical analog score (VAS/NAS).

Contraindications to Lumbar Epidural Injections and DSNRBs

Lumbar ESIs and DSNRBs are **NOT** indicated in cases that do not fulfill the above criteria. Of note, lumbar epidural steroid injections are not indicated in the following scenarios:

- Cancer:
 - New onset low back pain with a history of cancer, multiple risk factors for cancer, or strong clinical suspicion for cancer in the absence of advanced imaging studies (to rule out local cancer involvement)
 - Epidural injections may be considered if cancer is ruled-out or if the patient's pain is felt to be unrelated to their cancer AND they meet one of the above criteria lists (Items 1, 2, or 3)

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- Infection:
 - New onset of low back pain with fever in the absence of advanced imaging studies (to rule out local infection)
 - History of active intravenous drug use
 - History of recent or ongoing systemic bacterial or fungal infection
 - o Immunosuppression
- Cauda equina syndrome
 - o New onset urinary retention, fecal incontinence, or saddle anesthesia
 - o Rapidly progressing (or other) neurological deficits
- Axial Low Back Pain without lower extremity symptoms
- Co-existing medical conditions that would preclude the safe performance of the injection or be a contraindication to the intervention (e.g. bleeding disorder, presence of an epidural mass, or central nerve system (CNS) disorders[#] such as transverse myelitis or other demyelinating disorder)

*Note that if a CNS process is present, but the pain or neurologic deficit is clearly unrelated, an ESI may still be indicated if the patient meets one of the above criteria lists (Items 1, 2, or 3)

Procedural Requirements, Utilization, and Restrictions:

Lumbar epidural steroid injections, regardless of approach or indication, are subject to the following requirements and restrictions:

- Contrast enhanced fluoroscopy or CT guidance.
 - For transforaminal ESIs, live contrast-enhanced fluoroscopy or digital subtraction angiography is preferred, though contrast-enhanced CT guidance may be performed with the understanding that this form of visualization might not detect intravascular flow leading to potential complications, especially if particulate steroids are used.
 - Exceptions to the use of contrast are considered in patients who have a significant history and/or are at high risk for an adverse event if contrast material is used (e.g. contrast allergy).
 - In these cases, physicians should consider using a test-dose injection prior to injecting any particulate steroids and/or use only non-particulate steroid solutions.
 - The reasons for not using contrast should be documented in the procedure report.

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- Injections are performed independently based on the patients' symptoms and response to prior injections and approach (if performed). There is no role for a routine "series of 3" ESIs.
- If a prior lumbar ESI provided no relief, a second ESI is allowed following reassessment of the patient, injection technique and/or medication used.
- No more than 3 lumbar ESIs and/or DSNRBs may be performed in a 6-month period of time.
- No more than 6 lumbar ESIs and/or DSNRBs may be performed in a 12-month period of time regardless of the number of levels involved.
- Films that adequately document final needle position and injectate flow must be retained and made available upon request.
- No more than 2 transforaminal injections may be performed at a single setting (e.g. single level bilaterally or two levels)
- For caudal or lumbar interlaminar injections, only one per session may be performed and NOT in conjunction with a transforaminal injection.
- For each session, no more than 80mg of triamcinolone, 80 mg of methylprednisolone, 12 mg of betamethasone, 15 mg of dexamethasone or equivalent corticosteroid dosing should be used.
- Given the recent RCT evidence (Kennedy et al, Pain Medicine, 2014; El-Yahchouchi et al, Pain Medicine, 2014) for the therapeutic equivalency of dexamethasone to particulate steroid, particulate-free steroid, such as dexamethasone, should be used as the first line drug in all transforaminal ESIs. Particulate steroid should be used only after failure of particulate-free steroid and with appropriate patient counseling and safeguards, such as digital subtraction imaging.
- Local anesthesia is usually sufficient for a majority of lumbar ESIs though on occasion minimal to moderate conscious sedation is an appropriate option
- If monitored anesthesia care is utilized, the need for such sedation should be clearly documented in the medical records.

Rationale

Lumbar epidural steroid injections are one the most commonly performed injection procedures in the treatment of spine-related pain. The proposed Coverage Policy (also known as the "Policy") put forth by the North American Spine Society utilizes an evidence-based approach to spinal care when possible. In the absence of strict evidence-based criteria, the Policy utilizes the multidisciplinary and non-conflicted experience and expertise of the task force in order to reflect reasonable standard practice indications in the United States.

For lumbar radicular pain, the rationale for coverage is based on high-level evidence and what most practitioners would consider to be accepted practice patterns. Lumbar radicular pain may be caused by

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a myriad of pathologic conditions including, but not limited to lumbar disc herniation, lumbar stenosis (central or foraminal), lumbar spondylolisthesis, post-operative perineural fibrosis, or failed low back surgery syndrome. Multiple randomized-controlled trials have demonstrated that ESIs are effective in the treatment lumbar radiculitis caused by disc herniation¹⁻⁸. There is sufficient literature to suggest that at least a trial of ESI's for radicular pain caused by conditions other than disc herniation is appropriate ⁹⁻¹⁷ prior to considering surgical intervention.

For neurogenic claudication, the rationale for coverage is based on what most practitioners would consider to be accepted practice patterns. Neurogenic claudication is caused by spinal stenosis, either degenerative or isthmic. There is literature to suggest that ESIs are effective in reducing pain in this patient population^{10,18,19} though this treatment seems to be less effective in this group than in patients with herniated discs^{20,21}. In addition, there is data that shows that the injection of epidural steroid is equivalent to epidural local anesthetic^{15, 22-26}. It should be noted that epidural injection of local anesthetic has been clearly demonstrated to be more effective than a placebo²⁷. Based on these data, it is felt that a trial of epidural injections is reasonable prior to the consideration of surgical intervention for neurogenic claudication associated with lumbar spinal stenosis.

For selected cases of LBP, the rationale for coverage is based on what most practitioners would consider to be accepted practice patterns. While epidural injections are not typically considered an effective treatment for isolated, non-specific low back pain, they can be helpful in certain circumstances as described above. It is acknowledged that there is a paucity of data on this topic. In the absence of quality data, this coverage recommendation is guided by what appears to be reasonable and accepted practice patterns.

The rationale for the procedural **requirements, utilization, and restrictions** is based on what most practitioners would consider to be accepted practice patterns. In addition, there are a number of reports of complications associated with epidural injections²⁸⁻³³ that have occurred primarily as a result of intravascular injection. The use of live, contrast-enhanced fluoroscopy, digital subtraction, and the use of non-particulate steroids minimizes these risks.

As the potential risks with ESIs are both local from the procedure itself and systemic from the medications injected (specifically steroids), it is reasonable to place limits on the number of injections that should be administered in a given time. Currently, there are no data to support performing a predetermined "series" of injections. The determination to perform more than one injection should be based on the patient's response to the prior injection, the approach/location it was administered, the patient's symptoms, the medications used, and the imaging findings. This evaluation needs to be done via a face-to-face encounter and the reasons for repeating the injection clearly documented.

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Author Disclosure

Scott Kreiner, MD: Nothing to Disclose.

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