

Clinical Policy: Caudal or Interlaminar Epidural Steroid Injections

Reference Number: CP.MP.164

Date of Last Revision: 07/22

[Coding Implications](#)

[Revision Log](#)

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

Description

Epidural steroid injections have been used for pain control in patients with radiculopathy, spinal stenosis, and nonspecific low back pain, despite inconsistent results as well as heterogeneous populations and interventions in randomized trials. Epidural injections are performed utilizing three approaches in the lumbar spine: caudal, interlaminar, and transforaminal. Generally, candidates for epidural steroid injection are individuals who have acute radicular symptoms or neurogenic claudication unresponsive to traditional analgesics and rest, with significant impairment in activities of daily living.

Note: For guidelines for transforaminal ESIs, reference *CP.MP.165 Selective Nerve Root Blocks and Transforaminal Epidural Steroid Injections*.

Policy/Criteria

It is the policy of health plans affiliated with Centene Corporation® that invasive pain management procedures performed by a physician are **medically necessary** when *the relevant criteria are met, only one procedure is performed per visit, with imaging guidance (except in rare instances, with documented justification), and the member/enrollee is not currently being treated with full anticoagulation therapy. If on warfarin, international normalized ratio (INR) should be ≤ 1.4 prior to the procedure.* Discontinuing anti-platelet therapy is a clinical decision balancing risks and benefits of the procedure on therapy, versus the underlying medical condition if not treated appropriately.²³

- I. It is the policy of health plans affiliated with Centene Corporation® that caudal or interlaminar epidural steroid injections (ESIs) are **medically necessary** for the following indications:
 - A. *One caudal or interlaminar ESI for acute pain management* (pain lasting < 3 months) when all of the following are met:
 1. There is severe radicular pain that interferes substantially with activities of daily living (ADLs);
 2. Severe pain persists after treatment with nonsteroidal anti-inflammatory drugs (NSAID) and/or opiates (both ≥ 3 days or contraindicated/not tolerated);
 3. The member/enrollee cannot tolerate chiropractic or physical therapy and the injection is intended as a bridge to therapy.
 - B. *Initial ESI for chronic pain*, all of the following:
 1. Request is for one caudal or interlaminar ESI at one level in the cervical, thoracic or lumbar region;
 2. Persistent radicular pain has been caused by spinal stenosis, disc herniation or degenerative changes in the vertebrae, as confirmed by physical exam and imaging;
 3. Pain interferes with ADLs and has lasted for at least 3 months;

Caudal or Interlaminar Epidural Steroid Injections

4. The member/enrollee has failed to respond to conservative therapy including all of the following:
 - a. ≥ 6 weeks chiropractic, physical therapy or prescribed home exercise program;
 - b. NSAID ≥ 3 weeks or NSAID contraindicated or not tolerated;
 - c. ≥ 6 weeks activity modification.
 - C. *Second caudal or interlaminar ESI for chronic pain that **did not** improve from the first ESI, all of the following:*
 1. Request is for an ESI at one level in the cervical, thoracic or lumbar region;
 2. At least 2 weeks have passed since the first ESI.
 - D. *Subsequent caudal or interlaminar ESI for recurrence of chronic pain that **had improved** from the first or second ESI, all of the following:*
 1. Initial injection(s) led to $\geq 50\%$ relief and functional improvement for at least 2 months;
 2. At least 2 months have passed since the last ESI;
 3. Less than 4 injections have been administered within 12 months;
 4. Less than 12 months have elapsed since the initial injection at the level requested.
- II. It is the policy of health plans affiliated with Centene Corporation that *a third or subsequent caudal or interlaminar ESI for chronic pain that **did not** improve from the first two ESIs is considered **not medically necessary** because effectiveness has not been established.*
- III. It is the policy of health plans affiliated with Centene Corporation that *continuation of injections beyond 12 months or more than 4 therapeutic injections is considered **not medically necessary** because effectiveness and safety have not been established. When more definitive therapies cannot be tolerated or provided, consideration will be made on a case by case basis.*
- IV. It is the policy of health plans affiliated with Centene Corporation that *caudal or interlaminar ESI for any other indication or location is considered **not medically necessary** because effectiveness has not been established.*

Background

There is much debate on the efficacy and medical necessity of multiple interventions for managing spinal pain. Epidural glucocorticoid injections have been used for pain control in patients with radiculopathy, spinal stenosis, and nonspecific low back pain despite inconsistent results as well as heterogeneous populations and interventions in randomized controlled trials (RCTs). Epidural injections are performed utilizing 3 approaches in the lumbar spine: caudal, interlaminar, and transforaminal.² Generally, candidates for epidural steroid injection are individuals who have acute radicular symptoms or neurogenic claudication unresponsive to traditional analgesics and rest, with significant impairment in activities of daily living. Epidural steroid injections have been used in the treatment of spinal stenosis for many years, and no validated long-term outcomes have been reported to substantiate their use. However, significant improvement in pain scores have been reported at 3 months after injection.

CLINICAL POLICY

Caudal or Interlaminar Epidural Steroid Injections

Zhai et al conducted a meta-analysis to assess the effects of various surgical and nonsurgical modalities, including epidural injections, used to treat lumbar disc herniation (LDH) or radiculitis. A systemic literature review identified RCTs that compared the use of local anesthetic with and without steroids. The outcomes included pain relief, functional improvement, opioid intake, and therapeutic procedural characteristics. The reviewers concluded the meta-analysis confirms that epidural injections of local anesthetic with or without steroids have beneficial but similar effects in the treatment of patients with chronic low back and lower extremity pain.¹

Results of a 2-year follow-up of 3 randomized, double-blind, controlled trials, with a total of 360 patients with chronic persistent pain of disc herniation receiving either caudal, lumbar interlaminar or transforaminal epidural injections, showed similar efficacy of the 3 techniques with local anesthetic alone or local anesthetic with steroid.² Caudal and interlaminar trials used in the assessment showed some superiority of steroids over local anesthetic at 3 and 6 month follow-up. Interlaminar with steroids were superior to transforaminal at 12-months.²

Coding Implications

This clinical policy references Current Procedural Terminology (CPT®). CPT® is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2020, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

| CPT® Codes | Description |
|------------|--|
| 62320 | Injection(s), of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, including needle or catheter placement, interlaminar epidural or subarachnoid, cervical or thoracic; without imaging guidance |
| 62321 | Injection(s), of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, including needle or catheter placement, interlaminar epidural or subarachnoid, cervical or thoracic; with imaging guidance (i.e., fluoroscopy or CT) |
| 62322 | Injection(s), of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, including needle or catheter placement, interlaminar epidural or subarachnoid, lumbar or sacral (caudal); without imaging guidance |
| 62323 | Injection(s), of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, including needle or catheter placement, interlaminar epidural or subarachnoid, lumbar or sacral (caudal); with imaging guidance (i.e., fluoroscopy or CT) |
| 62324 | Injection(s), including indwelling catheter placement, continuous infusion or intermittent bolus, of diagnostic or therapeutic substance(s) (eg, anesthetic, |

Caudal or Interlaminar Epidural Steroid Injections

| CPT [®] Codes | Description |
|---------------------------|--|
| | antispasmodic, opioid, steroid, other solution), not including neurolytic substances, interlaminar epidural or subarachnoid, cervical or thoracic; without imaging guidance |
| 62325 | Injection(s), including indwelling catheter placement, continuous infusion or intermittent bolus, of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, interlaminar epidural or subarachnoid, cervical or thoracic; with imaging guidance (i.e., fluoroscopy or CT) |
| 62326 | Injection(s), including indwelling catheter placement, continuous infusion or intermittent bolus, of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, interlaminar epidural or subarachnoid, lumbar or sacral (caudal); without imaging guidance |
| 62327 | Injection(s), including indwelling catheter placement, continuous infusion or intermittent bolus, of diagnostic or therapeutic substance(s) (eg, anesthetic, antispasmodic, opioid, steroid, other solution), not including neurolytic substances, interlaminar epidural or subarachnoid, lumbar or sacral (caudal); with imaging guidance (i.e., fluoroscopy or CT) |

| HCPCS Codes | Description |
|----------------|-------------|
| N/A | |

ICD-10-CM Diagnosis Codes that Support Coverage Criteria

+ Indicates a code requiring an additional character

| Code | Description |
|-----------------------|--|
| M47.22 | Other spondylosis with radiculopathy, cervical region |
| M47.23 | Other spondylosis with radiculopathy, cervicothoracic region |
| M47.24 | Other spondylosis with radiculopathy, thoracic region |
| M47.25 | Other spondylosis with radiculopathy, thoracolumbar region |
| M47.26 | Other spondylosis with radiculopathy, lumbar region |
| M47.27 | Other spondylosis with radiculopathy, lumbosacral region |
| M48.00 through M48.08 | Spinal Stenosis |
| M50.10 through M50.13 | Cervical disc disorder with radiculopathy |
| M51.14 through M51.17 | Thoracic, thoracolumbar and lumbosacral intervertebral disc disorders with radiculopathy |
| M54.12 | Radiculopathy, cervical region |
| M54.13 | Radiculopathy, cervicothoracic region |
| M54.14 | Radiculopathy, thoracic region |
| M54.15 | Radiculopathy, thoracolumbar region |
| M54.16 | Radiculopathy, lumbar region |
| M54.17 | Radiculopathy, lumbosacral region |

Caudal or Interlaminar Epidural Steroid Injections

| Code | Description |
|-------|--|
| M54.5 | Low back pain |
| M54.6 | Pain in thoracic spine |
| M96.1 | Postlaminectomy syndrome, not elsewhere classified |

| Reviews, Revisions, and Approvals | Revision Date | Approval Date |
|--|---------------|---------------|
| Caudal and interlaminar ESI criteria reviewed in CP.MP.118 | 04/18 | 04/18 |
| Split from CP.MP.118 Injections for Pain Management. No criteria changes. | 08/18 | |
| In section D regarding second or subsequent ESI for chronic pain that improved from the diagnostic injections, changed requirement for 3 months having passed from the previous injection to 2 months. Anticoagulation indication moved to policy/criteria section as it is applicable to all injections in this policy. | 08/19 | 08/19 |
| References reviewed and updated | 06/20 | 07/20 |
| In policy statement, changed “with or without radiographic guidance” to “with imaging, (except in rare instances, with documented justification).” Added, “Request is not for cervical interlaminar ESI above C7” to B.5, C.3 and D.5. Changed “review date” in the header to “date of last revision” and “date” in the revision log header to “revision date.” References reviewed and updated. Replaced “member” with “member/enrollee” in all instances. Specialist review. | 07/21 | 07/21 |
| Removed “Request is not for cervical interlaminar ESI above C7” from B.5, C.3 and D.5. | 09/21 | 09/21 |
| Annual review. Note added regarding guidelines for transforaminal ESIs. Background updated with no impact on criteria. References reviewed and updated. | 07/22 | 07/22 |

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Caudal or Interlaminar Epidural Steroid Injections

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Caudal or Interlaminar Epidural Steroid Injections

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Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

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CLINICAL POLICY

Caudal or Interlaminar Epidural Steroid Injections



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Note: For Medicaid member/enrollees, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

Note: For Medicare member/enrollees, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs, LCDs, and Medicare Coverage Articles should be reviewed prior to applying the criteria set forth in this clinical policy. Refer to the CMS website at <http://www.cms.gov> for additional information.

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