

Cohere Medical Policy -Epidural Steroid Injections (ESI) Clinical Guidelines for Medical Necessity Review

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Guideline Information:

Specialty Area: Disorders of the Musculoskeletal System

Guideline Name: Cohere Medical Policy - Epidural Steroid Injection (ESI)

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Type: $[\underline{\mathbf{X}}]$ Adult (18+ yo) | $[\underline{\mathbf{X}}]$ Pediatric (0-17yo)

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Medical Necessity Criteria

Service: Epidural Steroid Injections (ESI)

Recommended Clinical Approach

Epidural steroid injections (ESIs) are used to treat radicular back and neck pain when conservative and noninvasive treatments fail. ESI involves the administration of an anesthetic and/or steroid in the epidural space of the spinal cord to treat inflammation that affects nerve roots. Three approaches for ESI include (1) Interlaminar or translaminar, (2) caudal, and (3) transforminal. ESI should be performed using radiographic image guidance. This guideline does not apply to epidural, spinal injections performed for perioperative pain management, or intrathecal, epidural injections done to assess candidacy for intrathecal pump

Medical Necessity Criteria

Indications

- → An **epidural steroid injection (ESI)** is considered appropriate if **ANY** of the following is **TRUE**^{2-16,19}:
 - The injection is a diagnostic, transforaminal ESI or diagnostic selective nerve root block (SNRB) to identify the cause of pain for surgical planning, and ALL of the following are met:
 - The pain is causing a functional or vocational disability, or pain levels of greater than or equal to 4 on a scale of 0 to 10;
 AND
 - Documentation of a preoperative evaluation and plan for surgery; OR
 - ◆ ALL of the following are TRUE:
 - The patient has ANY of the following:
 - o Acute herpes zoster; OR
 - Postherpetic neuralgia that has failed or had adverse side effects with trial of neuropathic medications;
 AND

- The patient has received no more than 6 ESIs over 12 consecutive months¹⁶; OR
- The patient has radicular pain or neurogenic claudication and ALL of the following are TRUE:
 - Findings are consistent with radicular symptoms or neurogenic claudication (e.g., history, signs and symptoms, or physical exam); AND
 - The patient has undergone advanced imaging (CT scan or MRI) that demonstrates findings consistent with ANY of the following:
 - Radicular symptoms that correlate to the side requested (e.g., spinal stenosis, foraminal stenosis, neural impingement, disc herniation, advanced degenerative changes, postlaminectomy changes etc.); OR
 - Neurogenic claudication; AND
 - **ALL** of the following are **TRUE**:
 - The patient's pain level is greater than or equal to 4 out of 10 on a scale of 0 to 10; AND
 - The pain is causing a functional or vocational disability; AND
 - ANY of the following:
 - ALL of the following:
 - The injection is for acute radicular pain secondary to acute disc herniation; AND
 - Failure of conservative management for greater than 2 weeks, including anti-inflammatory medications, analgesics, or prescription medications (e.g., oral steroids, narcotics, neuropathic pain medications); OR
 - The injection is an initial ESI injection with failure of conservative management, defined by ALL of the following:
 - Greater than 6 weeks duration; AND
 - Anti-inflammatory medications, analgesics, or prescription medications (e.g., oral steroids, narcotics, neuropathic pain medications) if not contraindicated; AND

- Physical therapy; OR
- A second ESI when ALL of the following are TRUE:
 - The initial injection has failed; AND
 - The injection is planned with a different approach, level, or medication type; OR
- The injection is a repeat ESI, and ALL of the following are TRUE:
 - ◆ The most recent ESI injection resulted in ALL of the following:
 - 50% or greater pain or symptom relief;
 AND
 - The pain or symptom relief lasted greater than or equal to 2 months duration; AND
 - The patient is participating in ongoing rehabilitative approaches (e.g., physical therapy, chiropractic care, or physician-guided home exercise program); AND
 - No more than 4 injections are given per rolling
 12-month period per spinal region.

Non-Indications

- → Epidural steroid injection (ESI) is not considered appropriate if ANY of the following is TRUE¹⁵⁻¹⁶:
 - ◆ Transforaminal epidural steroid injections [TFESIs] at more than 2 nerve roots (more than 1 spinal level when bilateral or more than 2 spinal levels when unilaterally)¹⁶; **OR**
 - ◆ Presence of ANY of the following¹⁶:
 - Non-specific low back pain (LBP); OR
 - Myelopathy; OR
 - Cauda equina syndrome; OR
 - Rapidly progressing neurological deficit; OR
 - Epidural abscess; OR
 - ESIs are performed at multiple anatomical regions during the same date of service; OR
 - ◆ ESIs performed with biologics (e.g., platelet-rich plasma, stem cells, amniotic fluid, etc.)¹⁶; **OR**

- ◆ ESI is performed with ultrasound image guidance (except for injections performed during pregnancy) where it would be reasonable to avoid radiographs¹⁶; OR
- ◆ ESI injection is performed without fluoroscopy or computed tomography (CT) guidance²⁰; **OR**
- ◆ Allergy or sensitivity to local anesthetics or other medications; **OR**
- When other types of spine injections (e.g., facet, sacroiliac joint injections, ESIs to other spinal regions etc.) are performed on the same date of service, unless medical necessity to perfrom multiple injections is documented in chart (e.g., risk of holding anticoagulant multiple times, presence of facet cyst causing radicular symptoms for which ESI and facet injection can be allowed at the same time, etc.); OR
- ESIs performed for perioperative pain management; OR
- Intrathecal ESIs done to assess candidacy for intrathecal pump;
 OR
- ◆ A series of ESIs are included in a single request.

Level of Care Criteria

Outpatient

Procedure Codes (CPT/HCPCS)

CPT/HCPCS Code	Code 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 (ie, 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 (ie, fluoroscopy or CT)	
64479	Injection(s), anesthetic agent(s) and/or steroid; transforaminal epidural, with imaging guidance (fluoroscopy or CT), cervical or thoracic, single level	
64480	Injection(s), anesthetic agent(s) and/or steroid; transforaminal epidural, with imaging guidance (fluoroscopy or CT), cervical or thoracic, each additional level (List separately in addition to code for primary procedure)	
64483	Injection(s), anesthetic agent(s) and/or steroid; transforaminal epidural, with imaging guidance (fluoroscopy or CT), lumbar or sacral, single level	
64484	Injection(s), anesthetic agent(s) and/or steroid; transforaminal epidural, with imaging guidance (fluoroscopy or CT), lumbar or sacral, each additional level (List separately in addition to code for primary procedure)	
64999	Unlisted procedure, nervous system	

Medical Evidence

The Centers for Medicare and Medicaid Services (CMS) (2023) published Local Coverage Determination (LCD) Epidural Steroid Injections (ESIs) for Pain Management (L39036). The LCD includes several recommendations and non-indications for ESIs for radiculopathy and pain related to the spine, including Grade A recommendations for (i) contrast-enhanced fluoroscopy to guide ESIs for improved accuracy of delivery and (ii) transforaminal ESIs in patients with lumbar disc herniations with radiculopathy.¹⁶

Kreiner et al. (2014) developed a clinical guideline for the North American Spine Society (NASS) to diagnose and treat lumbar disc herniation. Twenty-nine questions and recommendations resulted from this effort, including a Grade A recommendation for use of contrast-enhanced fluoroscopy in the routine performance of ESIs; a Grade A recommendation for ESI in treatment of lumbar disc herniation with radiculopathy; no recommendation given for optimal frequency or quantity of injections; and insufficient evidence found to make a recommendation for or against the 12-month efficacy of transforaminal ESI in lumbar disc herniation with radiculopathy.¹⁶

Manchikanti et al. (2012) conducted a randomized, double-blind, active-control trial. Patients with a diagnosis of disc herniation, radiculitis, facet joint pain, or sacroiliac joint pain were excluded. The two groups of subjects received local anesthetic only or local anesthetic mixed with non-particulate betamethasone. Success was to be measured by a greater than 50% decrease in pain and disability, and the outcome was 77% in the local anesthetic-only group and 67% in the local anesthetic mixed with non-particulate betamethasone.¹⁰

In a systematic review, Kim et al. (2011) tested the hypothesis that dexamethasone phosphate (DP) and methylprednisolone acetate (MPA) would achieve similar outcomes when used in the treatment of lumbar radiculopathy by epidural injection. The study determined that, when compared with particulate methylprednisolone, nonparticulate dexamethasone appeared to be close in safety and effectiveness.⁸

Furman et al. (2010) conducted a prospective, single-arm, pilot, observational study. Inclusion criteria included patients with lumbar radicular pain, and exclusion criteria included the presence of irreversible psychological barriers or anatomical anomalies. The group concluded that subjects undergoing fluoroscopic-guided, contrast-enhanced lumbar interlaminar epidural steroids for primarily radicular pain versus axial pain experienced at least a 3-month improvement on the Numeric Pain Rating Scale (NPRS).²

Ahmed et al. (2023) conducted a systematic meta-analysis of randomized controlled trials to compare ultrasonography (USG) guidance and conventional fluoroscopy (FL) guidance for ESI in the treatment of radiculopathy. Of the 7 RCTs included in the meta-analysis, the authors found no statistically significant difference in pain reduction between USG and FL groups in lumbosacral ESIs at 1 and 3 months. Functional improvement after ESIs was comparable between the groups. The risk of inadvertent vascular puncture in USG-guided ESIs was lower compared to conventional FL-guided ESIs [odds ratio 0.21 (0.07, 0.64)], and the procedure time in the USG group was significantly lower compared to the FL group.¹⁷

Mahmoud et al. (2024) conducted a systematic review and meta-analysis comparing different ESI approaches with the aim of investigating the effectiveness of various ESI procedures in adults with lower back pain. The authors concluded that a transforaminal (TF) ESI approach was most effective in decreasing pain indices (visual analog score [VAS] and Oswestery disability index [ODI]) in the long term. The authors also found that a parasagittal intralaminar (PIL) approach was most effective in decreasing VAS in the short term (less than 6 months).¹⁸

Yang et al. (2020) performed a meta-analysis of 6 RCTs to compare the clinical effectiveness of ESI to conservative treatments for patients with lumbosacral radicular pain, including physical therapy, lifestyle modifications, exercise, and patient education. The meta-analysis included 490 adult patients; 249 patients received ESI, and 241 received conservative treatment. The patients had a confirmed diagnosis of lumbar disc herniation or spinal stenosis. Pooled analysis of pain scores at short-term and intermediate follow-ups for 3 RCTs demonstrated a significant reduction in pain for ESI as compared to conservative treatments. Long-term follow-up for 2 RCTs showed comparable effectiveness of ESI and conservative treatments. Pooled



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Clinical Guideline Revision History/Information

Original Date: April 21, 2021			
Review History			
Version 2	12/29/2023		
Version 3	9/20/2024	Updated language regarding conservative treatment.	
Version 4	12/19/2024	 Annual review Aligned to current policy guidelines. Updated medical evidence section. References updated. CPT codes and descriptions updated; temporary codes deleted Updated indications and non-indications for clarity and ease of conversion to rules. 	