cohere h e A L T H

Cohere Medical Policy - Lumbar or Thoracic Spinal Fusion

Clinical Guidelines for Medical Necessity Review

Version:3Effective Date:December 19, 2024

Important Notices

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Guideline Information:

Specialty Area: Disorders of the Musculoskeletal System **Guideline Name:** Cohere Medical Policy - Lumbar Spinal Fusion

Date of last literature review: 12/16/2024 **Document last updated:** 12/17/2024 **Type:** [X] Adult (18+ yo) | [X] Pediatric (0-17 yo)

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Medical Necessity Criteria

Service: Lumbar or Thoracic Spinal Fusion

Recommended Clinical Approach

Lumbar or thoracic spinal fusion (arthrodesis) may be necessary for conditions such as fracture or dislocation, spinal deformities (e.g., scoliosis or kyphosis), spinal stenosis, infection, tumors, or degenerative changes. This may be accomplished by an anterior, posterior, or posterolateral approach. Surgery may provide more rapid relief than non-surgical treatment options, as well as prevent further spinal cord dysfunction and neurological deficits.¹ Advanced imaging is recommended prior to surgical intervention.²⁻⁴

Medical Necessity Criteria

Indications

- → Lumbar or thoracic spinal fusion is considered appropriate if ANY of the following is TRUE:
 - A lumbar spinal fusion is considered appropriate if ALL of the following are TRUE:
 - No nicotine product use for 6 weeks with a negative lab test within 30 days (unless surgery is urgently required for progressive neurologic deficit); AND
 - **ANY** of the following is **TRUE**:
 - The procedure is **lumbar fusion with or without decompression,** and **ALL** of the following are **TRUE**:
 - **ANY** of the following:
 - Radiographic evidence of kyphosis or scoliosis greater than 40 degrees; **OR**
 - Failure of conservative management for greater than 6 weeks, including AT LEAST TWO of the following:
 - Anti-inflammatory medications, analgesics, or prescription medications (e.g., oral steroids,

narcotics, neuropathic pain medications) if not contraindicated; **OR**

- Physical therapy or physician-directed home exercise program; OR
- Injections, when medically appropriate, including **ANY** of the following:
 - ♦ Facet injections; OR
 - Medial branch blocks (MBB);
 OR
- The procedure is lumbar fusion with or without decompression with radiographic evidence of instability or iatrogenic instability caused by the decompression at all levels planned to be fused, and ANY of the following is TRUE⁵⁻⁶:
 - The patient has signs or symptoms of a potential cauda equina syndrome and ALL of the following⁷:
 - Magnetic resonance imaging (MRI)
 reveals compressive pathology; AND
 - **ANY** of the following symptoms:
 - Bowel, bladder, and erectile dysfunction; **OR**
 - Diffuse motor weakness; OR
 - Saddle-distribution anesthesia; OR
 - The patient has lumbar stenosis, and ALL of the following are TRUE:
 - **ANY** of the following symptoms⁵:
 - Lower extremity pain, weakness, fatigue, paresthesias, and sensory changes; OR
 - Gluteal and low back pain (LBP); **OR**
 - Bilateral or unilateral symptoms; **OR**
 - Symptoms that present only with activity; OR

- Exacerbating factors include standing, walking, and other upright exercises; OR
- Pain that is relieved in a sitting or supine position or with forward flexion at the waist; OR
- Lower extremity pain that is made worse by walking; AND
- **ANY** of the following physical examination findings^{5, 8-9}:
 - Focal motor weakness or sensory deficit; OR
 - Decreased or absent lower extremity reflexes; OR
 - Wide-based gait; OR
 - Positive Romberg's test (poor standing balance with eyes closed); OR
 - Positive straight leg raise (SLR; reproduction of lower extremity pain upon extension at the knee);
 AND
- MRI reveals compressive pathology and ANY of the following¹⁰:
 - Failure of conservative management for greater than 6 weeks, including AT LEAST TWO of the following:
 - Anti-inflammatory medications, analgesics or prescription medications (e.g., oral steroids, narcotics, neuropathic pain medications) if not contraindicated; OR
 - Physical therapy, including a physician-directed home exercise program; OR

- Injections, when medically appropriate, including ANY of the following:
 - Facet injections; **OR**
 - Medial branch blocks (MBB); OR
- The patient has severe pain or disability affecting their quality of life and limiting their daily life (including working and inability to provide self-care); OR
- The patient has progressive neurological motor deficits; OR
- Unstable fracture noted on imaging with neuro deficit or myelopathy with **ANY** of the following;
 - ♦ Chance fracture; OR
 - Burst fracture with neuro deficit;
 - Fracture-dislocation; OR
- The patient has lumbar radiculopathy and ALL of the following are TRUE:
 - **ANY** of the following symptoms⁸:
 - Lower extremity pain, paresthesia, weakness, or numbness in a myotomal or dermatome distribution; **OR**
 - Radicular pain with coughing, sneezing, or straining; OR
 - LBP; AND
 - **ANY** of the following physical examination findings⁸⁻⁹:
 - Sensory disturbance (e.g., loss of sensation or decreased sensory response) or weakness in a dermatomal or myotomal distribution; **OR**

- Absent or decreased lower extremity reflex; OR
- Reduced spinal mobility; **OR**
- ANY of the following positive specialty tests (unless medically contraindicated):
 - Straight leg raise; OR
 - Crossed Lasègue's (or crossed straight leg raise);
 OR
 - Femoral nerve stretch; OR
 - ♦ Slump; AND
- MRI reveals compressive pathology and the patient has **ANY** of the following¹⁰:
 - Failure to show significant improvement in pain or disability level due to symptoms, despite receiving non-surgical management for more than 6 weeks, including AT LEAST TWO of the following (unless medically contraindicated):
 - Anti-inflammatory medications, analgesics or prescription medications (e.g., oral steroids, narcotics, neuropathic pain medications) if not contraindicated; OR
 - Physical therapy, including a physician-directed home exercise program; OR
 - Injections, when medically appropriate, including ANY of the following:
 - Facet injections; **OR**
 - Medial branch blocks (MBB); OR

- Severe pain or disability affecting their quality of life and limiting their daily life (including working and being unable to provide self-care);
 OR
- Progressive neurological motor deficits; OR
- A thoracic spinal fusion is considered appropriate if ALL of the following are TRUE:
 - No nicotine product use for 6 weeks with a negative lab test within 30 days (unless surgery is urgently required for progressive neurologic deficit); **AND**
 - **ANY** of the following is **TRUE**:
 - The procedure is thoracic fusion with or without decompression with **ALL** of the following:
 - Radiographic evidence of **ANY** of the following:
 - Lumbar curve greater than 40 degrees requiring extension into the thoracic spine; OR
 - Positive sagittal balance or sagittal vertebral axis (SVA) greater than 5 cm;
 OR
 - Pelvic incidence lumbar lordosis (PI-LL) mismatch greater than 10 degrees; **AND**
 - Failure to show significant improvement in pain or disability level due to symptoms, despite receiving non-surgical management for more than 6 weeks, including AT LEAST TWO of the following (unless medically contraindicated):
 - Anti-inflammatory medications, analgesics or prescription medications (e.g., oral steroids, narcotics, neuropathic pain medications) if not contraindicated; OR
 - Physical therapy, including a physician-directed home exercise program; OR

- Injections, when medically appropriate, including **ANY** of the following:
 - Facet injections; OR
 - Medial branch blocks (MBB); **OR**
- Junctional fusion for adjacent segment disease (e.g., stenosis, kyphosis, listhesis) is appropriate when ANY of the following is TRUE:
 - Adjacent cervical fusion is medically appropriate and approvable; **OR**
 - Adjacent lumbar fusion is medically appropriate and approvable; **OR**
 - Adjacent thoracic fusion is medically appropriate and approvable; **OR**
- Revision or repeat spinal fusion (e.g., due to prior unhealed fusion attempt) when at least 12 months have elapsed since the original surgery and imaging studies confirm the absence of healing in the preceding 3 months unless failure has occurred (e.g., pseudoarthrosis)¹.

Non-Indications

- → A thoracic or lumbar spinal fusion is not considered appropriate if ANY of the following is TRUE:
 - Current laboratory-confirmed nicotine use (unless surgery is urgently required for progressive neurologic deficit); OR
 - **ANY** of the following^z:
 - In disc herniation, adjunct to primary excision of a central or posterolateral disc herniation at any level in the absence of instability or spondylolisthesis; OR
 - In lumbar stenosis when greater than 50% bilateral facet resection is not required to achieve neurologic decompression; OR
 - Discogenic LBP that does not meet the criteria listed in the Indications section above.

Level of Care Criteria Inpatient or Outpatient

Procedure Codes (CPT/HCPCS)

| CPT/HCPCS Code | Code Description | | |
|----------------|---|--|--|
| 20999 | Unlisted procedure, musculoskeletal system, general | | |
| 22532 | Arthrodesis, lateral extra cavitary technique, including minimal discectomy to prepare interspace (other than for decompression); thoracic | | |
| 22533 | Arthrodesis, lateral, lumbar | | |
| 22534 | Each additional, thoracic or lumbar, (add-on code) | | |
| 22556 | Arthrodesis, anterior interbody technique, including minimal discectomy to prepare interspace (other than for decompression); thoracic | | |
| 22558 | Arthrodesis, anterior interbody technique, including minimal discectomy to prepare interspace (other than for decompression); lumbar | | |
| 22585 | Arthrodesis, anterior interbody technique, including minimal discectomy to prepare interspace (other than for decompression); each additional interspace (List separately in addition to code for primary procedure) | | |
| 22586 | Arthrodesis, pre-sacral interbody technique, including disc space preparation, discectomy, with posterior instrumentation, with image guidance, includes bone graft when performed, L5-S1 interspace | | |
| 22610 | Arthrodesis, posterior or posterolateral technique, single interspace; thoracic (with lateral transverse technique, when performed) | | |
| 22612 | Arthrodesis, posterior or posterolateral technique, single interspace; lumbar (with lateral transverse technique, when performed) | | |
| 22614 | Arthrodesis, posterior or posterolateral technique, single | | |

| | interspace; each additional interspace (List separately in addition to code for primary procedure) | | |
|-------|--|--|--|
| 22630 | Arthrodesis, posterior interbody technique, including laminectomy and/or discectomy to prepare interspace (other than for decompression), single interspace, lumbar | | |
| 22632 | Arthrodesis, posterior interbody technique, including laminectomy and/or discectomy to prepare interspace (other than for decompression), single interspace, lumbar; each additional interspace (List separately in addition to code for primary procedure) | | |
| 22633 | Arthrodesis, combined posterior or posterolateral technique with posterior interbody technique including laminectomy and/or discectomy sufficient to prepare interspace (other than for decompression), single interspace, lumbar | | |
| 22634 | Arthrodesis, combined posterior or posterolateral technique with posterior interbody technique including laminectomy and/or discectomy sufficient to prepare interspace (other than for decompression), single interspace, lumbar; each additional interspace (List separately in addition to code for primary procedure) | | |
| 22800 | Arthrodesis, posterior, for spinal deformity, with or without cast; up to 6 vertebral segments | | |
| 22802 | Arthrodesis, posterior, for spinal deformity, with or without cast; 7 to 12 vertebral segments | | |
| 22804 | Arthrodesis, posterior, for spinal deformity, with or without cast; 13 or more vertebral segments | | |
| 22808 | Arthrodesis, anterior, for spinal deformity, with or without cast; 2 to 3 vertebral segments | | |
| 22810 | Arthrodesis, anterior, for spinal deformity, with or without cast; 4 to 7 vertebral segments | | |

| 22812 | Arthrodesis, anterior, for spinal deformity, with or without cast; 8 or more vertebral segments | | |
|-------|---|--|--|
| 22830 | Exploration of spinal fusion | | |
| 22840 | Posterior non-segmental instrumentation (eg, Harrington rod technique, pedicle fixation across 1 interspace, atlantoaxial transarticular screw fixation, sublaminar wiring at C1, facet screw fixation) (List separately in addition to code for primary procedure) | | |
| 22841 | Internal spinal fixation by wiring of spinous processes (List separately in addition to code for primary procedure) | | |
| 22842 | Posterior segmental instrumentation (eg, pedicle fixation, dual rods with multiple hooks and sublaminar wires); 3 to 6 vertebral segments (List separately in addition to code for primary procedure) | | |
| 22843 | Posterior segmental instrumentation (eg, pedicle fixation, dual rods with multiple hooks and sublaminar wires); 7 to 12 vertebral segments (List separately in addition to code for primary procedure) | | |
| 22844 | Posterior segmental instrumentation (eg, pedicle fixation, dual rods with multiple hooks and sublaminar wires); 13 or more vertebral segments (List separately in addition to code for primary procedure) | | |
| 22845 | Anterior instrumentation; 2 to 3 vertebral segments (List separately in addition to code for primary procedure) | | |
| 22846 | Anterior instrumentation; 4 to 7 vertebral segments (List separately in addition to code for primary procedure) | | |
| 22847 | Anterior instrumentation; 8 or more vertebral segments (List separately in addition to code for primary procedure) | | |
| 22848 | Pelvic fixation (attachment of caudal end of | | |

| | instrumentation to pelvic bony structures) other than sacrum (List separately in addition to code for primary procedure) | | |
|-------|---|--|--|
| 22849 | Reinsertion of spinal fixation device | | |
| 22853 | Insertion of interbody biomechanical device(s) (eg, synthetic cage, mesh) with integral anterior instrumentation for device anchoring (eg, screws, flanges), when performed, to intervertebral disc space in conjunction with interbody arthrodesis, each interspace (List separately in addition to code for primary procedure) | | |
| 22854 | Insertion of intervertebral biomechanical device(s) (eg, synthetic cage, mesh) with integral anterior instrumentation for device anchoring (eg, screws, flanges), when performed, to vertebral corpectomy(ies) (vertebral body resection, partial or complete) defect, in conjunction with interbody arthrodesis, each contiguous defect (List separately in addition to code for primary procedure) | | |
| 22859 | Insertion of intervertebral biomechanical device(s) (eg, synthetic cage, mesh, methylmethacrylate) to intervertebral disc space or vertebral body defect without interbody arthrodesis, each contiguous defect (List separately in addition to code for primary procedure) | | |
| 22899 | Unlisted procedure, spine | | |

Medical Evidence

Kreiner et al. (2020) published a systematic review of guidelines for low back pain (LBP) diagnosis and treatment. Insufficient evidence was found to recommend for or against a particular fusion technique for the treatment of LBP. No literature evidence was found to adequately address differences in clinical outcomes or functional status for single-level vs. multilevel fusions. No studies were found to address the effectiveness of fusion over discectomy, discectomy with rhizotomy or decompression alone.¹³

The American College of Radiology (ACR) Expert Panel on Neurological Imaging has published appropriateness criteria related to myelopathic evaluation. Agarwal et al. (2021) updated the previous criteria for myelopathy with magnetic resonance imaging (MRI) recommended as initial imaging for acute onset myelopathy. MRI is also recommended for chronic or progressive myelopathy due to its superior resolution of soft tissue and ability to evaluate surrounding structures. Computed tomography (CT) may be appropriate, with CT myelography of possible use prior to surgical intervention.² Non-contrast MRI is usually appropriate for (LBP; radiography and CT may be appropriate for LBP with and without radiculopathy. This applies to surgical candidates with persistence or progression of symptoms having failed six weeks of medical management. MRI, CT, and CT myelography are recommended for suspected cauda equina syndrome. In cases of osteoporosis or chronic steroid use, radiography, non-contrast MRI, or CT is usually appropriate.³

Reid et al. (2019) completed a literature review regarding the current state of lumbar fusion indications and techniques for degenerative spine disease. The group states that few randomized trials exist that studied lumbar fusion for degenerative disease. Lumbar instrumentation technologies in degenerative diseases are discussed, including issues from Harrington rod implantation, such as sagittal imbalance and flatback syndrome. Pedicle screw fixation development is stated to significantly improve successful fusion rates.¹²

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Clinical Guideline Revision History/Information

| Original Date: September 29, 2023 | | | | |
|-----------------------------------|------------|--|--|--|
| Review History | | | | |
| Version 2 | 7/25/2024 | Updated language regarding conservative treatment and nicotine use. | | |
| Version 2A | 9/20/2024 | Added CPT code 22840. | | |
| Version 3 | 12/19/2024 | Annual review. Added thoracic criteria. Revised conservative care and nicotine use sections. Literature review- additions made Style updates made per Style Guide | | |