



**Blue Cross
Blue Shield
Blue Care Network**
of Michigan

Nonprofit corporations and independent licensees
of the Blue Cross and Blue Shield Association



**PAIN MANAGEMENT
NEUROABLATION
AUTHORIZATION REQUEST FORM**

Utilization management toll-free phone: 1-833-217-9670
Utilization management local phone: 313-908-6040
Utilization management fax: 313-483-7323

| |
|---|
| Today's date (mm/dd/yyyy): ___ / ___ / ___ |
| Provider contact name: |
| Provider contact phone: |
| Provider contact fax: |
| Provider name: |
| Provider TIN: |
| Provider NPI: |
| Practice/group name: |
| Provider physical address: |
| Provider mailing address (if different): |

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|--|
| Member name: |
| Date of birth (mm/dd/yyyy): ___ / ___ / ___ |
| Member ID (including any alpha prefix): |
| Health plan: |
| Notification method preference: <input type="checkbox"/> Postal mail <input type="checkbox"/> Fax |
| Mailing address or fax number: |
| Notes: |

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|--|---|
| Where will the procedure take place? <input type="checkbox"/> Provider office <input type="checkbox"/> Outpatient facility <input type="checkbox"/> Inpatient hospital <input type="checkbox"/> Ambulatory surgical center | |
| Facility name: | Facility contact name: |
| Facility TIN: | Facility contact phone: |
| Facility NPI: | Facility contact fax: |
| Facility physical address: | Facility mailing address (if different): |

| Requested procedure code | Modifier: LT, RT or 50 (bilateral) | Quantity | Spine level |
|---|------------------------------------|--|-------------|
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| Diagnosis code(s): | | Anticipated date of service (mm/dd/yyyy): ___ / ___ / ___ | |
| Case urgency <input type="checkbox"/> Standard <input type="checkbox"/> Expedited In keeping with guidelines from the National Committee for Quality Assurance and Centers for Medicare & Medicaid Services, prior authorization requests qualify for expedited review when the standard review time frame could do one of the following: <ul style="list-style-type: none"> • Seriously jeopardize the life, health or safety of the member or others, due to the member's psychological state. • In the opinion of a practitioner with knowledge of the member's medical or behavioral health condition, subject the member to adverse health consequences without the care or treatment that is the subject of the request. | | | |
| Patient's height: | Patient's weight: | Patient's BMI: | |

| What type of procedure is planned? (Select one and answer all adjacent questions.) | | |
|---|--|--|
| <input type="checkbox"/> Initial cervical/thoracic or lumbar thermal radiofrequency ablation (answer a – i) | a. Has moderate to severe pain (rated at least 3 out of 10), primarily axial in nature, been present for 3 months? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | b. Are any radiculopathy or claudication symptoms (burning, tingling, cramping) present? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | c. Are more than 2 levels (either unilateral or bilateral) planned for the procedure? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | d. Has conservative treatment been attempted for at least 12 weeks/3 months? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | e. Has medication been attempted as part of conservative treatment? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | f. Has chiropractic care, physical therapy, and/or home exercise program been attempted as part of conservative treatment? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | g. Does imaging show any other possible causes of pain (such as stenosis, nerve impingement, fracture, or infection)? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | h. Were two medial branch blocks performed at the same location as planned radiofrequency ablation? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | i. Does the medical record show that BOTH medial branch blocks resulted in at least 80% reduction in pain and improvement in function? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| <input type="checkbox"/> Repeat cervical/thoracic or lumbar thermal radiofrequency ablation (answer a – d) | a. Were criteria met for initial ablation? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | b. Does the medical record confirm at least 50% reduction in pain and improvement in function for at least 6 months after the prior procedure? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | c. Have 2 or more radiofrequency ablation sessions been done in this same spine region in the past 12 months (cervical/thoracic or lumbar/sacral)? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | d. Have 4 or more total medial branch block sessions for the entire spine been done in the past 12 months? | <input type="checkbox"/> Yes <input type="checkbox"/> No |

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| <input type="checkbox"/> Initial sacroiliac thermal radiofrequency ablation (answer a – f) | a. Is there presence of moderate to severe pain (rated at least 3 out of 10) that interferes with daily activities? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | b. Has conservative treatment been attempted for at least 12 weeks/3 months? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | c. Has medication been attempted as part of conservative treatment? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | d. Has chiropractic care, physical therapy, and/or home exercise program been attempted as part of conservative treatment? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | e. Were 2 diagnostic injections performed on the L5 primary dorsal ramus and the 1 st – 3 rd sacral dorsal rami branches? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | f. Does the medical record show at least 75% reduction in pain and improvement in function with both diagnostic injections? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| <input type="checkbox"/> Repeat sacroiliac thermal radiofrequency ablation (answer a – c) | a. Does the medical record confirm at least 50% reduction in pain and improvement in function for at least 6 months after the prior procedure? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | b. Have 2 or more radiofrequency ablation sessions been done in this same spine region in the past 12 months (cervical/thoracic or lumbar/sacral)? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| | c. Have 4 or more total medial branch block sessions for the entire spine been done in the past 12 months? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| <input type="checkbox"/> Intraosseous radiofrequency ablation (basivertebral nerve) | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Pulsed radiofrequency ablation | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Cooled radiofrequency ablation | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Endoscopic radiofrequency ablation | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Chemical ablation | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Laser ablation | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Cryoablation, cryoanalgesia, or cryoneurolysis | No questions related to this procedure; proceed to next section. | |
| <input type="checkbox"/> Procedure planned for any area other than the spine | No questions related to this procedure; proceed to next section. | |

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| Do any of the following apply? (Answer a – d) | |
| a. Systemic or localized infection at planned injection site | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| b. Radiofrequency ablation is planned at a fused spine level | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| c. Other pain management interventions planned same day (i.e. epidural steroid injection, SI joint injection, trigger point injection, etc.) | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| d. Pain management procedures planned in multiple regions (i.e. cervical/thoracic AND lumbar or sacral) | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Will the procedure be performed with fluoroscopic guidance? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Is general anesthesia, conscious sedation, or monitored anesthesia care planned? | <input type="checkbox"/> Yes <input type="checkbox"/> No |



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|--|--|
| Is this request for lovera (cryoablation)? | <input type="checkbox"/> Yes <input type="checkbox"/> No |
| Include imaging reports, surgical plan and clinical documentation of all conservative therapies that have been attempted as well as the duration of each type of conservative treatment. | |
| Form completed by: | Date: |