



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



PAIN MANAGEMENT FACET JOINT INJECTIONS 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 contact email:
Provider name:
Provider TIN:
Provider NPI:
Practice/group name:
Provider physical address:
Provider mailing address (if different):

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:

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
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 medial branch block (answer a – j)	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. Does the pain interfere with daily activities and get worse with bending or twisting?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	c. Are any radiculopathy or claudication symptoms (burning, tingling, cramping) present?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	d. Is the injection being done for diagnostic purposes only?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	e. Will more than the recommended amount of anesthetic be used? (Total amount less than 0.3cc for cervical spine and 0.5cc for lumbar)	<input type="checkbox"/> Yes <input type="checkbox"/> No
	f. Are more than 2 levels (either unilateral or bilateral) planned for the procedure?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	g. Has conservative treatment been attempted for at least 4 weeks/1 month?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	h. Has medication been attempted as part of conservative treatment?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	i. 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
	j. 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
<input type="checkbox"/> Second medial branch block (i.e. facet joint(s) that received one block prior) (answer a – d)	a. Were criteria met for initial block?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	b. Does the medical record show at least 80% reduction in pain and improvement in function with initial block?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	c. Have 4 or more medial branch block sessions been done in this same spine region in the past 12 months (cervical/thoracic or lumbar)?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	d. Have 8 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"/> Third or greater medial branch block (i.e. facet joint(s) that received 2 or more blocks prior)	No questions related to this procedure; proceed to next section.	
<input type="checkbox"/> Initial therapeutic joint injection for treatment of facet cyst (answer a – b)	a. Does imaging (CT, MRI) confirm facet cyst causing nerve root compression or displacement?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	b. Does imaging correlate with symptoms and rule other possible causes out?	<input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Second therapeutic joint injection for treatment of facet cyst (answer a – b)	a. Did the original symptoms return?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	b. Does the medical record confirm at least 50% reduction in pain and improvement in function after the first procedure?	<input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Third or greater joint injection for treatment of facet cyst	No questions related to this procedure; proceed to next section.	
<input type="checkbox"/> Initial therapeutic injection for chronic facet-related pain (answer a – d)	a. Were two medial branch blocks performed at the same location as the planned therapeutic intervention?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	b. 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
	c. Are more than 2 levels (either unilateral or bilateral) planned for the procedure?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	d. Does the medical record document why a radiofrequency ablation is not possible?	<input type="checkbox"/> Yes <input type="checkbox"/> No
<input type="checkbox"/> Second or greater therapeutic injection for chronic facet-related pain (i.e. facet joint(s) that received at least one therapeutic injection prior) (answer a – e)	a. Were criteria met for initial therapeutic injection?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	b. Are more than 2 levels (either unilateral or bilateral) planned for the procedure?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	c. Does the medical record confirm at least 50% reduction in pain and improvement in function for 3 months after the prior procedure?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	d. Have 4 or more therapeutic facet joint injection sessions been done in this same spine region in the past 12 months (cervical/thoracic or lumbar)?	<input type="checkbox"/> Yes <input type="checkbox"/> No
	e. Have 8 or more total therapeutic facet joint injection sessions for the entire spine been done in the past 12 months?	<input type="checkbox"/> Yes <input type="checkbox"/> No

Do any of the following apply? (Answer a – f)	
a. Injection with steroid planned with uncontrolled diabetes, uncontrolled hypertension, or congestive heart failure present	<input type="checkbox"/> Yes <input type="checkbox"/> No
b. Systemic or localized infection at planned injection site	<input type="checkbox"/> Yes <input type="checkbox"/> No
c. Facet joint intervention is planned at a fused spine level	<input type="checkbox"/> Yes <input type="checkbox"/> No
d. Facet joint intervention is planned at the site of a previously successful radiofrequency ablation	<input type="checkbox"/> Yes <input type="checkbox"/> No
e. 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
f. 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

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: