

Prior Authorization Request Form for Intraosseous Basivertebral Nerve Ablation (Intracept® System) #486

Medical Policy #485 Intraosseous Basivertebral Nerve Ablation (Intracept® system)

Please use this form to assist in identifying members who meet Blue Cross Blue Shield of Massachusetts' (BCBSMA's) medical necessity criteria for Intraosseous Basivertebral Nerve Ablation (Intracept® system). For members who do not meet the criteria, submit a letter of medical necessity with a request for Clinical Exception (Individual Consideration).

Once completed, please fax to: This form must be completed and faxed to: Medical and Surgical: 1-888-282-0780; Medicare Advantage: 1-800-447-2994.

CLINICAL DOCUMENTATION

Copies of clinical documentation that supports the medical necessity criteria for Intracept® system must be submitted with this form. If the patient does not meet all the criteria listed below, please submit a letter of medical necessity explaining why an exception is justified.

ratient information		
Patient Name:	Today's Date:	
BCBSMA ID#:	Date of Treatment:	
Date of Birth:	Place of Service: Outpatient ☐ Inpatient ☐	
	Distributor:	
Physician Information	Facility Information	
Name:	Name:	
Address:	Address:	
Phone #:	Phone #:	
Fax#:	Fax#:	
NPI#:	NPI#:	
Please check off that the patient meets <u>ALL</u> the following	criteria:	
Chronic lower back pain >6 months		
Refractory to optimal nonsurgical medical management incl		
chiropractic therapy, epidural or facet injection therapy, luml		
home use of heat/cold therapies, pharmacotherapy, cognitive		
Modic type I or II changes on MRI, endplate hypointensity (
plus hyperintensity on T2 images (Type 1) involving in the e		
inflammation, edema, disruption, and fissuring of the endpla		
marrow, and changes to the vertebral body marrow including	g replacement of normal bone marrow by fat.	

No previous history of BVN ablation at the planned level of treatment.	
No more than one to two (1-2) vertebral bodies treated during a single session.	
Treatment of no more than 4 vertebral bodies per patient lifetime.	

CONTRAINDICATIONS

Please check off that the patient <u>DOES NOT HAVE ANY</u> of the following contraindications:	
Individual does not have any of the following:	
 Evidence on imaging (MRI, flexion/extension radiographs, etc.) indicating that pain may be due to another condition including but not limited to lumbar stenosis, spondylolisthesis, segmental instability, disc herniation, degenerative scoliosis, or facet arthropathy or effusion with clinically suspected facet joint pain, or 	
 Metabolic bone disease (eg, osteoporosis), treatment of spine fragility fracture, trauma/compression fracture, or 	
 History of or active spinal cancer, or 	
Spine infection or active systemic infection, or	
 Bleeding diathesis, or 	
 Neurogenic claudication, lumbar radiculopathy or radicular pain due to neurocompression (eg, HNP, stenosis), as primary symptoms, or 	
Radiographic evidence of:	
 Lumbar/lumbosacral disc extrusion or protrusion >5mm at levels L3-S1; Lumbar/lumbosacral spondylolisthesis > Grade 2 at any level; Lumbar/lumbosacral spondylolysis at levels L3-S1; 	
 Lumbar/lumbosacral facet arthrosis/effusion correlated with facet-mediated pain at levels L3- S1 	
 Patients with severe cardiac or pulmonary compromise, or 	
 Patients with implantable pulse generators (eg, pacemakers, defibrillators) or other electronic implants, or 	
Pregnancy, or	
• BMI >40.	

Providers should enter the <u>relevant diagnosis code(s)</u> below:

Code	Description	
64628	Thermal destruction of intraosseous basivertebral nerve, first 2 vertebral bodies	
64629	Thermal destruction of intraosseous basivertebral nerve, each additional vertebral body	

Providers should enter other relevant code(s) below:

Code	Description	