

# Radiofrequency Ablation (Initial and Repeat) Prior Authorization Form



This form must be completed by a person with thorough clinical knowledge of the member's current clinical presentation and his/her clinical evaluation history. **Clinical documentation supporting the medical necessity of this request is required.** For more information, please refer to the medical policy document Radiofrequency Ablation (Neurotomy, Denervation, Rhizotomy) Cervical, Thoracic, Lumbosacral, Sacroiliac or Knee Pain (MC/F024) located at <https://www.aspirushealthplan.com>. Please contact Customer Service at 866.631.5404 if there are questions.

**Return completed form and clinical documentation to:** Aspirus Health Plan, Attn: Integrated Health Services, PO Box 1062, Minneapolis, MN 55440 or Fax to 763.847.4014.

PATIENT INFORMATION			
Patient Last Name	Patient First Name	Member ID	Patient Date of Birth
Date of Service	ICD 10 Diagnosis	Procedure Code(s)	

ORDERING PROVIDER INFORMATION			
Ordering Provider Name	Ordering Provider Signature		NPI
Clinic Name	Clinic Phone Number	Clinic Fax Number	NPI
Clinic Address	City	State	Zip Code

SERVICING PROVIDER INFORMATION			
Servicing Provider Name			NPI
Facility Name	Facility Phone Number	Facility Fax Number	NPI
Facility Address	City	State	Zip Code

THE PROCEDURE WILL BE DONE IN ONE OF THE FOLLOWING SITES OF CARE
<input type="checkbox"/> Office or Ambulatory Surgery Center <input type="checkbox"/> Outpatient Hospital. <i>Nearest office or ambulatory surgery center capable of providing service is 60 miles or more from member's home.</i> <input type="checkbox"/> Outpatient Hospital. <i>Documentation supports that the member is considered at high risk for complications that require a hospital setting, such as, but not limited to, the member's physical status is classified as ASA III-VI, per the American Society of Anesthesiologists Physical Status Classification System.</i> <input type="checkbox"/> Outpatient Hospital. <i>Documentation supports that the servicing provider does not hold privileges at an office or ambulatory surgery center within 60 miles driving distance from the member's home.</i>

INITIAL REQUEST FOR NON-PULSED RADIOFREQUENCY ABLATION FOR FACET-MEDIATED CERVICAL, THORACIC, LUMBOSACRAL, OR SACROILIAC JOINT PAIN
Level of Procedure: _____ <input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral
<input type="checkbox"/> The member has chronic (at least 6 months) cervical, thoracic, or lumbosacral pain suggestive of facet or sacroiliac joint origin. Must be documented in the medical record on history and physical exam.
<input type="checkbox"/> The member has failed at least 3 months of conservative therapy. Please indicate below the type of conservative therapy ( <i>check all that apply</i> ). Clinical documentation should indicate dates of services. <input type="checkbox"/> Chronic Back Program <input type="checkbox"/> Physical Therapy <input type="checkbox"/> Steroid Injections <input type="checkbox"/> Pharmacotherapies <input type="checkbox"/> Structured Home Exercise Program <input type="checkbox"/> Weight Loss, if indicated <input type="checkbox"/> Activity Modification <input type="checkbox"/> Spinal Manipulation
<input type="checkbox"/> The member has not had prior fusion surgery at the level where treatment is being considered.
<input type="checkbox"/> The member has undergone at least one anesthetic block of the involved facet, medial, primary dorsal-rami, or sacral lateral branch nerves. Anesthetic block date: _____ % Pain Reduction: _____

**INITIAL REQUEST FOR INTRAOSSEOUS ABLATION (e.g., INTRACEPT®) OF THE BASIVERTBRAL NERVE (BVN) FOR LOW BACK PAIN**

- The member has chronic (at least 6 months) isolated low back pain suggestive of skeletal endplate inflammation. Must be documented in the medical record on history and physical exam.
- MRI shows Type 1 or Type 2 Modic changes of the vertebral endplates at 3 or less contiguous levels, L3-S1.
- The member has failed at least 3 months of conservative therapy. Please indicate below the type of conservative therapy (*check all that apply*). Clinical documentation should indicate dates of services.
  - Chronic Back Program     Physical Therapy     Steroid Injections     Pharmacotherapies     Structured Home Exercise Program
  - Weight Loss, if indicated     Activity Modification     Spinal Manipulation

**INITIAL REQUEST FOR NON-PULSED RADIOFREQUENCY ABLATION OF THE GENICULAR NERVE (ARTICULAR NERVE BRANCHES) FOR KNEE PAIN**

- The member has chronic (at least 6 months) knee pain
- The member has failed at least 3 months of conservative therapy. Please indicate below the type of conservative therapy (*check all that apply*). Clinical documentation should indicate dates of services.
  - Pharmacotherapies     Physical Therapy     Activity Modification     Structured Home Exercise Program
  - Weight Loss, if indicated
- The member has undergone at least 1 anesthetic block of the genicular nerve with 50% pain reduction  
Anesthetic block date: \_\_\_\_\_ % Pain Reduction: \_\_\_\_\_
- The member has no history of total knee arthroplasty (TKA) – knee pain is due to knee osteoarthritis (OA) and member is not a good surgical candidate for TKA due to medical comorbidities and/or a high body mass index (BMI).  
**OR**
- The member has a history of TKA for osteoarthritis and is equal to or greater than 6 months post-op.

**REPEAT REQUEST FOR RFA ON THE SAME NERVE (SITE)**

- |   |  |
|---|--|
| Level of Procedure: _____ <input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral | Date of Prior Radiofrequency Treatment of the Same Nerve |
|---|--|
- A minimum of six (6) months has elapsed since prior ablative treatment of the same nerve.
  - Prior ablative treatment resulted in at least a 50% reduction in pain for a minimum of 10 weeks following the previous treatment.