

# Radiation Therapy, Intensity Modulated (IMRT) 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, refer to the clinical policy document MC/L009 Radiation Therapy, Intensity Modulated (IMRT) 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
Date of Service	ICD 10 Diagnosis	Procedure Code(s)	Patient Date of Birth
			Number of Fractions
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
<b>Request for IMRT for any of the following conditions: <i>check all that apply</i></b>			
<input type="checkbox"/> Anus or anal canal cancer <input type="checkbox"/> Breast cancer – any of the following: <input type="checkbox"/> Treatment of left-sided internal mammary nodes <input type="checkbox"/> Partial breast irradiation of up to 5 fractions <input type="checkbox"/> Central nervous system (CNS) tumors (primary or benign) including the brain, brain stem, and spinal cord <input type="checkbox"/> Cervical cancer <input type="checkbox"/> Endometrial cancer <input type="checkbox"/> Esophageal cancer <input type="checkbox"/> Gastroesophageal junction (Siewert III tumors) <input type="checkbox"/> Head and neck cancer, including lymphoma and solitary plasmacytomas - Treatment includes the following areas (check all that apply): <input type="checkbox"/> Pharynx (nasopharynx, oropharynx, hypopharynx) <input type="checkbox"/> Larynx cancer (stage III or IV glottic cancer) <input type="checkbox"/> Salivary glands <input type="checkbox"/> Oral cavity (includes tongue) <input type="checkbox"/> Nasal cavity <input type="checkbox"/> Paranasal sinuses <input type="checkbox"/> Mediastinal tumors (eg, lymphomas, thymomas, including tracheal cancer) <input type="checkbox"/> Pancreatic cancer <input type="checkbox"/> Prostate cancer			
<b>Request for IMRT for a condition not listed above – must meet any of the following:</b>			
<i>Check all that apply; Include treatment plan comparison documentation of IMRT and non-IMRT technique</i>			
<input type="checkbox"/> A non-IMRT technique would increase the probability of clinically meaningful normal tissue toxicity (e.g., as specified by the Radiation Therapy Oncology Group (RTOG) or QUANTEC guidelines) and demonstrated on a comparison of treatment plans for the IMRT and non-IMRT technique (e.g., three-dimensional conformal treatment plan).			
<input type="checkbox"/> The same or immediately adjacent area has been previously irradiated, and the dose distribution within the individual must be sculpted to avoid exceeding the cumulative tolerance dose of nearby tissue.			