

**Prior Authorization for Therapeutic Radiation Procedures Including Intensity Modulation Radiation Treatment (IMRT), Stereotactic Radio Surgery (SRS) and Stereotactic Body Radiation Therapy (SBRT)**

This worksheet is a tool to help you identify and collect the clinical information required to support a prior authorization request for UnitedHealthcare Medicare Advantage members. After the clinician completes the clinical information, please go to [UHCprovider.com](http://UHCprovider.com) > Prior Authorization and Notification > Oncology > Medicare Advantage Therapeutic Radiation to submit the prior authorization request. Submitting the clinical information from this worksheet alone won't automatically result in prior authorization approval; additional clinical information may be requested during the clinical review process. If you have questions about your submission or need to request an expedited review, please call 866-889-8054.

<b>Patient/ Member</b>	First Name:	Middle Initial:	Last Name:
	DOB (mm/dd/yyyy):		Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female
	Health Plan:		Member ID:

<b>Clinical Information</b>	ICD-10 Code(s):
	What is the radiation therapy treatment start date (mm/dd/yyyy)?
	<b><i>Please have the answers to below questions prepared as these questions may be asked.</i></b>
	Is radiation being delivered as:
	<input type="checkbox"/> Initial treatment for a newly diagnosed prostate cancer without distant metastatic disease <input type="checkbox"/> Post-prostatectomy adjuvant therapy due to adverse pathology without distant metastatic disease <input type="checkbox"/> Post-prostatectomy salvage therapy due to recurrence without distant metastatic disease <input type="checkbox"/> Palliative therapy (i.e. non-curative therapy to alleviate obstructive symptoms or bleeding) <input type="checkbox"/> Other (e.g. Recurrent prostate cancer, Definitive treatment of prostate in the metastatic setting)
	What is/was the patient's risk group (as defined by NCCN®)?
	<input type="checkbox"/> Very Low-risk (T1c and Gleason <= 6 and PSA under 10 ng/mL and 1-2 Positive Cores with <=50% involvement in each core and PSA density < 0.15 ng/mL/g) <input type="checkbox"/> Low-risk (T1-T2a and Gleason <= 6 and PSA under 10 ng/mL) <input type="checkbox"/> Favorable Intermediate-risk (T2b-T2c or PSA 10-20 ng/mL; Gleason (3+4) and <50% of cores are positive) <input type="checkbox"/> Unfavorable Intermediate-risk (T2b-T2c and/or PSA 10-20 ng/mL; and Gleason (4+3)) <input type="checkbox"/> High-risk (T3a or Gleason 8-10 or PSA > 20) <input type="checkbox"/> Very High-risk (T3b-T4 or > 4 Cores of Gleason 8-10 or Primary Gleason 5) <input type="checkbox"/> Regional (any T, N1, M0) <input type="checkbox"/> Distant metastases (i.e. spread to bone)
	If high or very high risk, will the pelvic lymph nodes be treated? <span style="float: right;"><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A</span>
	What treatment technique will be used?
	<input type="checkbox"/> Hypofractionated Intensity Modulated Radiation Therapy (IMRT) <input type="checkbox"/> Conventionally fractionated Intensity Modulated Radiation Therapy (IMRT) <input type="checkbox"/> Stereotactic Body Radiation Therapy (SBRT)

<b>Clinical Information</b>	Additional Comments/Information: