



Radiation Therapy Small Cell Lung Cancer Request

For NON-URGENT requests, please complete this document for authorization along with any relevant clinical documentation requested within this document (i.e. radiation therapy consultation, comparison plan, etc.) before submitting the case by web, phone, or fax. Failure to provide all relevant information may delay the determination. Phone and fax numbers can be found on evicore.com under the Guidelines and Fax Forms section. You may also log into the provider portal located on the site to submit an authorization request. **URGENT (same day) requests must be submitted by phone.**

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

Clinical Information	ICD-10 Code(s):	
	What is the radiation therapy treatment start date (mm/dd/yyyy)?	
	<i>eviCore is utilizing a clinical decision support submission model for this diagnosis. Please note that only some of the following example questions will need to be answered during the submission of your prior authorization request. For best results, the answers to these questions should be submitted online.</i>	
	Does the patient have a history of distant metastases (stage M1) (i.e. to brain, lung, liver, bone)? <input type="checkbox"/> Yes <input type="checkbox"/> No	
	What is the treatment intent?	
	<input type="checkbox"/> Curative, No surgery planned or performed <input type="checkbox"/> Curative, Post-operative (adjuvant) <input type="checkbox"/> Curative, Pre-operative (neo-adjuvant) <input type="checkbox"/> Curative, Treatment of the primary in an oligometastatic setting <input type="checkbox"/> Palliative (non-curative, to alleviate symptoms) <input type="checkbox"/> Prophylactic cranial irradiation (PCI)	
	What is the stage of the lung cancer at the time of original diagnosis?	
	<input type="checkbox"/> IA <input type="checkbox"/> IIIB <input type="checkbox"/> IB <input type="checkbox"/> IIIC <input type="checkbox"/> IIA <input type="checkbox"/> IV or Extensive stage <input type="checkbox"/> IIB <input type="checkbox"/> Loco-regional Recurrence <input type="checkbox"/> IIIA	
	If chemotherapy has been delivered, what has been the response?	
	<input type="checkbox"/> Complete response (CR) <input type="checkbox"/> Partial response (PR) <input type="checkbox"/> No response or stable disease <input type="checkbox"/> Progressive disease <input type="checkbox"/> Not applicable (N/A)	

How many fractions will be used for each phase?			
Phase 1	Phase 2	Phase 3	Treatment Technique
			Conventional isodose planning, complex
			3D conformal
			Intensity Modulated Radiation Therapy (IMRT)
			Tomotherapy (IMRT)
			Rotational Arc Therapy
			Proton Beam Therapy
			Stereotactic Body Radiation Therapy (SBRT)
			Biology-guided Radiation Therapy (BgRT)
			N/A
Will image guided radiation therapy (IGRT) be used for treatment?			<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Will the patient be receiving concurrent chemotherapy?			<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Will treatment be delivered twice daily (i.e. BID)?			<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
<i>Please be prepared to submit consult note, results of imaging from the past 60 days and radiation prescription or clinical treatment plan in order to expedite the review process. Failure to provide all relevant information may result in a delay.</i>			
Additional Comments/Information:			

