

Lung Project Data Elements			
DEMOGRAPHICS			
Data Elements	Options		
 Provider Date of initial Radiation/Oncology consult Date of Birth Gender Race 6. Medical Insurance (Check all that apply)	 First/Last mm/dd/yyyy Male/Female American Indian/Alaska Native Asian Native Hawaiian or Other Pacific Islander Black or African American White Arab/Middle Eastern Unknown or not reported Other (free text) No insurance/self-pay Medicare (all) Medicare Advantage-BCN Medicaid-Straight Medicaid -HMO Other Payer (government) Other Payer (Michigan and outstate) BCBSM-Michigan BCN- Michigan Commercial-HMO 		
7. Current Marital Status	 Married/ Domestic Partner Divorced Never Married 		
8. Cancer Type	 Separated Widowed Living with someone Single Breast Cancer Lung Cancer 		



LUNG PROJECT

L1: Patient: Pre-Treatment Lung Cancer Questionnaire

Time points: Pre RT Evaluation

Data Elements

Options

I have lack of energy	
I have nausea	
Because of my physical condition, I have trouble meeting the needs of my family	
I have pain	
I am bothered by side effects of treatment	
I feel ill	
I am forced to spend time in bed	
I am able to work	Not at all
My work (include work at home) is fulfilling	A little bit
	Somewhat
I am able to enjoy life	Quite a bit
I have accepted my illness	Very much
I am sleeping well	
I am enjoying the things I usually do for fun	
I am content with the quality of my life right now	
I have been short of breath	
I am losing weight	
My thinking is clear	
I have been coughing	



I am bothered by hair loss	
I have a good appetite	
I feel tightness in my chest	
Breathing is easy for me	
Have you ever smoked	No Yes
If Yes: I regret my smoking	Not at all A little bit Somewhat Quite a bit Very Much
Are you currently vaping?	• YES/NO
a) If yes, please select how often you change cartridges	 Cartridge refill lasts 2 days or longer 1 Cartridge refill per day 2 or more cartridge refills per day
b) Do you fill your own cartridge's or purchase prepackaged cartridges?	Fill own cartridgesPurchase prepackaged cartridges
c) Do you use flavored cartridges?	• YES/NO
d) Do you use any of the following while vaping?	 Marijuana Tobacco THC oil CBD with vaping



Have you ever, even once, used Cannabis? Think specifically about the past 30 days up to and including today. What is your best estimate of the number of days you used Cannabis during the past 30 days?	 Prefer not to answer – Proceed to question 28 No – Proceed to question 28 Yes 0 days 1 or 2 days 3 to 5 days 6 to 9 days 10 to 19 days 20 to 29 days
During the past 30 days, which one of the following ways did you use cannabis most often? Did you usually:	 All 30 days Smoke it (for example, in a joint, bong, pipe, or blunt) Eat it (for example, in brownies, cakes, cookies, or candy) Drink it (for example, in tea, cola, or alcohol) Vaporize it (for example, in an e-cigarette-like vaporizer or another vaporizing device) Dab it (for example, using waxes or concentrates) Apply to skin (for example, using lotions or oils) Administer rectally (for example, using suppositories). Use it some other way (please specify:
What is the major active ingredient in the cannabis product that you use the most? (This information can often be found on the package label.)	 THC –also called tetrahydrocannabinol CBD –also called cannabidiol Balanced levels of THC and CBD I don't know Other:
Please select the one response that best describes your swallowing ability over the course of the past week:	 No problems with swallowing this week Mild soreness only Can swallow solids with some difficulty Cannot swallow solids Cannot swallow liquids
Which best describes your race/ethnicity (mark all that apply)?	 American Indian/Alaska Native Arab/Middle Eastern Asian Black or African American Native Hawaiian or Pacific Islander White Other. Specify:



Are you of Hispanic/Latino origin?	• YES/NO	
What is the highest level of education you have completed?	 Some High School High School Graduate or GED Some College or Technical School Associate's Degree College Graduate (Bachelor's Degree) Graduate Degree 	
For the following additional symptoms or problems, please circle or mark the number that best applies to you during the past week.		
How much did you cough?		
Did you cough up blood?	Not at all	
Were you short of breath when you rested?	A little bit	
Were you short of breath when you walked?	Quite a bit	
Were you short of breath when you climbed stairs?	Very much	
Did you take any medicine for pain?	No Yes	
If yes, how much did it help?	Not at all A little bit Quite a bit Very much	
What is your height?	ftin	
What is your current weight?	pounds.	
L2: Patient: Weekly Lu	ung Cancer Swallowing Assessment	
	Time points: Weekly On-Treatment Visits	
Data Elements	Options	
Please select the one response that best describes your swallowing ability over the course of the past week:	 No problems with swallowing this week Mild soreness only Can swallow solids with some difficulty Cannot swallow solids 	

Cannot swallow liquids



L3: Patient Lung End of Treatment and Follow-up		
	<i>Time points: Last week of treatment & Follow-up visits 1-3-6 months</i>	
Data Elements	Options	
I have lack of energy		
I have nausea		
Because of my physical condition, I have trouble meeting the needs of my family		
I have pain	Not at all	
I am bothered by side effects of treatment	A little bit	
I feel ill	Somewhat	
I am forced to spend time in bed	Quite a bit	
I am able to work	Very much	
My work (include work at home) is fulfilling		
I am able to enjoy life		
I have accepted my illness		
I am sleeping well		
I am enjoying the things I usually do for fun		
I am content with the quality of my life right now		
I have been short of breath	Not at all	
I am losing weight	A little bit	
My thinking is clear	Somewhat	
I have been coughing	Quite a bit	
I am bothered by hair loss	Very much	
I have a good appetite		
I feel tightness in my chest		



Breathing is easy for me	
Have you ever smoked	No Yes
If Yes: I regret my smoking	Not at all A little bit Somewhat Quite a bit Very Much
Are you currently vaping?	• YES/NO
a) If yes, please select how often you change cartridges	 Cartridge refill lasts 2 days or longer 1 Cartridge refill per day 2 or more cartridge refills per day
b) Do you fill your own cartridge's or purchase prepackaged cartridges?	 Fill own cartridges Purchase prepackaged cartridges
c) Do you use flavored cartridges?	• YES/NO
d) Do you use any of the following while vaping?	 Marijuana Tobacco THC oil CBD with vaping
Have any of your radiation oncology providers ever asked if you use cannabis? If the only mention of cannabis that you recall is from a survey form, then the answer is "No".	• YES/NO
Have you been satisfied with your radiation oncology providers' ability to answer questions about cannabis?	 I have not asked any questions about cannabis No Yes
Have you used cannabis, even once, since the first treatment in your course of radiation?	 Prefer not to answer No Yes



If you use cannabis, please check ALL the reason(s) that you have chosen to do so. Please select the one response that best describes your swallowing ability over the course of the past week (Select only one)	 For pain For nausea For anxiety For depression For poor appetite For trouble sleeping For the high (recreational) To fight Cancer Other: No problems with swallowing this week Mild soreness only Can swallow solids with some difficulty Cannot swallow solids Cannot swallow liquids
during the past week.	, , , , , , , , , , , , , , , , , , ,
How much did you cough?	
Did you cough up blood?	Not at all
Were you short of breath when you rested?	A little bit
Were you short of breath when you walked?	Quite a bit
Were you short of breath when you climbed stairs?	Very much
Did you take any medicine for pain?	No Yes
If yes, how much did it help?	Not at all A little bit Quite a bit Very much



L4 CDA: Lung Cancer Baseline Clinical Data		
	Time points: Pre RT Evaluation	
Data Elements	Options	
Weight (specify lbs. or kg):	lb/kg.	
BMI or Height (specify inches or cm):	BMI and inches/cm.	
Tumor Stage:	T N M	
Histology:	 NSCLC – Squamous cell carcinoma NSCLC Adenocarcinoma SCLC – Iimited extensive Other (specify): No biopsy 	
Tumor location: (Check all that apply)	 Right upper lobe Right middle lobe Right lower lobe Left upper lobe Left lower lobe Superior sulcus 	
Surgical resection prior to radiation?	YES/NO Date:	
Margin status	 Positive Negative N/A 	



Г

Has there ever been any previous lung surgery?	 YES/ NO Date: Lobectomy Pneumonectomy Wedge resection Other (specify):
Was Is the patient on O2 prior to radiation? When is O2 used?	 NO/YES How many liters? Always Daytime Nighttime When short of breath
Pulmonary function prior to RT:	 PFT not done FEV1Liters,% predicted DLCO not done DLCO%/mm,% predicted
Current smoker?	 YES NO Unknown
Former smoker? (quit at least one month prior to diagnosis)	 YES NO Unknown



Was this patient counseled by a doctor or other healthcare worker about quitting cigarettes? (applies to current smokers only)

YES ٠ NO

•

Comorbidities:					
Does the patient have:	YES	NO	Does the patient have:	YES	NO
Hypertension?			Hemiplegia?	۵	
Diabetes mellitus?			Leukemia?		
Scleroderma?			Malignant lymphoma?		
Rheumatoid arthritis?			Myocardial infarction?		
Lupus?			Peripheral vascular disease?		
Cerebrovascular disease?			Ulcer disease?		
Chronic pulmonary disease?			Liver disease?		
Congestive heart failure?			Renal disease?		
Connective tissue disease?			Malignant solid tumor (other than lung)?		
Confusion?			Arrhythmia?		



	Time points: Last Week of RT
Data Elements	Options
Did the patient receive systemic therapy?	YESNO
If yes, check all that apply:	
Before radiation therapy	
Prior to Surgery	Agents (check all that apply): Cisplatin/Platinol-AQ/Platinol Etoposide/Eposin/VePesid/VP-16 Paclitaxel/Taxol/TAX Carboplatin/Paraplatin/Paraplatin-AQ Pemetrexed/Alimta Docetaxel/Taxotere/TXT Gemcitabine/Gemzar Vinorelbine/Navelbine Other (specify)
During radiation therapy	
 Prior to Surgery After surgery Without planned surgery 	Agents (check all that apply): Cisplatin/Platinol-AQ/Platinol Etoposide/Eposin/VePesid/VP-16 Paclitaxel/Taxol/TAX Carboplatin/Paraplatin/Paraplatin-AQ Pemetrexed/Alimta Docetaxel/Taxotere/TXT Gemcitabine/Gemzar Vinorelbine/Navelbine Other (specify)
Immunotherapy before radiation therapy	
 Prior to Surgery 	Agents (check all that apply): Atezolizumab Durvalumab Nivolumab Pembrolizumab Other (specify)



Immunotherapy during radiation therapy Prior to Surgery Agents (check all that apply): After surgery Atezolizumab Without planned surgery Durvalumab Nivolumab Pembrolizumab Other (specify) _____ L6 Physician: Lung Baseline-1st Week Time points: 1st Week of RT **Data Elements** Options What was the date of the first fraction? (mm/dd/yyyy) completed by the CDA Baseline Toxicity Scoring (CTCAE v 4.0) Please circle one number in each row. Adverse Event 4 5 2 **Gastrointestinal disorders** Asymptomatic; clinical or Symptomatic; altered Severely altered Life-threatening Death Esophagitis diagnostic observations only; eating/swallowing; none eating/swallowing; oral consequences; urgent intervention not indicated supplements indicated tube feeding, TPN or intervention indicated hospitalization indicated **Esophageal Pain** Mild pain Moderate pain; limiting Severe pain; limiting none instrumental ADL self-care ADL **General disorders** Fatigue none Fatigue relieved by rest Fatigue not relieved by Fatigue not relieved by rest; limiting rest, limiting self-care instrumental ADL ADL Respiratory, thoracic and mediastinal disorders Cough none Mild symptoms; Moderate symptoms, Severe symptoms; medical intervention limiting self-care ADL nonprescription intervention indicated; limiting indicated instrumental ADL Shortness of breath at Dyspnea none Shortness of breath with Shortness of breath with Life-threatening Death moderate exertion minimal exertion; rest; limiting self-care consequences; urgent limiting instrumental ADL ADL intervention indicated Pleuritic pain Mild pain Moderate pain; limiting Severe pain; limiting none instrumental ADL self-care ADL Pneumonitis Asymptomatic; clinical or Severe symptoms; none Symptomatic; medical Life-threatening Death diagnostic observations only; intervention indicated; limiting self-care ADL; respiratory compromise; intervention not indicated limiting instrumental ADL oxygen indicated urgent intervention indicated (e.g., tracheotomy or intubation) ECOG Performance Status'



	Fully active	Restricted in physically strenuous activity but ambulatory and able to carry out work of a light or sedentary nature,e.g., light housework, office work	of all s to car activit	selfcare ry out a ties. Up than 50	and capable but unable any work and about 0% of waking	Capable of only limited self-care, confined to bed or chair more than 50% of waking hours	Completely disabled. Cannot carry on any self- care. Totally confined to bed or chair	Dead
What % weight loss did the patient experience over the 6 months prior to the initiation of lung cancer treatment: (Enter "0" if patient's weight remained static, or if patient gained weight.)			_		%			
Is this patient enrolled on any lung cancer treatment clinical trial, study, or protocol (do not include MROQC)?				•	YES NO			
Does this study (these studies) influence your radiation dose / treatment plan for this patient			-	• •	YES NO Not appl	icable/not on study		
Does this study (these studies) influence your choice or sequencing of systemic therapy (immunotherapy and/or chemotherapy)?			-	•	YES NO Not appli	cable/not on study		
Is this a hypofractionated course of treatment (≤20 fractions inclusive of SBRT)?				•	YES NO			
<i>If yes, answer questions:</i> Was this a pathologically confirmed cancer or clinical diagnosis?			r	•	patholog clinical d			
Was nodal staging performed?				•	YES NO			



a) If yes, how was it determined (check all that apply)?	 Clinically by PET Pathologically by EBUS/bronchoscopy Pathologically by Mediastinoscopy Other (specify):
Is this a peripheral, central or ultra-central lesion?	 Peripheral Central Ultra-central

L7 Physician: Lung Weekly

Time points: Weekly On-Treatment Visits

Data Elements			Options			
Toxicity Scoring	(CTCAE v 4.0)	Please circle one	number in each ro	w.		
Adverse Event	0	1	2	3	4	5
Gastrointestinal	disorders					
Esophagitis	None	Asymptomatic; clinical or diagnostic observations only; intervention not indicated	Symptomatic; altered eating/ swallowing; oral supplements indicated	Severely altered eating/swallowing; tube feeding, TPN or hospitalization indicated	Life- threatening consequences; urgent intervention indicated	Death
Esophageal Pain	None	Mild pain	Moderate pain; limiting instrumental ADL	Severe pain; limiting self-care ADL		
General Disorders					·	
Fatigue	None	Fatigue relieved by rest	Fatigue not relieved by rest; limiting instrumental ADL	Fatigue not relieved by rest; limiting self-care ADL		
Respiratory disord	lers					
Cough	None	Mild symptoms; nonprescription intervention indicated	Moderate symptoms, medical intervention indicated; limiting instrumental ADL	Severe symptoms; limiting self-care ADL		
ECOG Performanc	e Status*					



	Fully active	Restricted in physically strenuous activit but ambulatory and able to carry out work of a light or sedentar nature, e.g., light housework, offic work	 unable to carry out any work activities. Up and about more than 50% of waking 	Capable of only limited self-care, confined to bed or chair more than 50% of waking hours	Completely disabled. Cannot carry on any self- care. Totally confined to bed or chair	Dead
	L8 Phys	ician: Lung C	ancer Clinical Ou			
			• · · ·	Tim	e points: Last weel	k of RT
Data Elements			Options			
What was the dat	What was the date of the last fraction?			yy) completed by the	e CDA	
Weight	Weight			bs. or kg)		
Did any break in treatment occur?			• YES/ NO			
If Yes, was it due to toxicity?			• YES/NO			
Was the toxicity-related treatment break >5 days?			• YES/NO			
Rate current esophageal pain on a scale of 0-10:			(0-10)			



Please select what you have recommended to the patient in the past month to manage esophagitis:			medie Miles • Narco (e.g. v oxyco • Other	narcotic prescription cations (e.g. Magic m mixture, xylocaine) otic prescription mec Vicodin, Percocet, m odone, fentanyl) r intervention ify):	lications orphine,		
Has the patient been admitted for a cardiac event? a) If YES, date of admission:			• YES/	NO ate)			
b) Select the cardiac event related to admission			CongPeric	ythmia gestive Heart failure cardial Effusion cardial Infarction			
Toxicity Sco	ring (CTC	AE v 4.0). Please ci	rcle one r	umber in each r	ow.		
Adverse Event	0	1		2	3	4	5
Gastrointestina	l disorders						
Esophagitis	none	Asymptomatic; clinical or diagnostic observations only; intervention not indicated	Symptomatic; altered eating/swallowing; oral supplements indicated		Severely altered eating/swallowing; tube feeding, TPN or hospitalization indicated	Life-threatening consequences; urgent intervention indicated	Death
Esophageal	none	Mild pain	Moderate	pain; limiting	Severe pain; limiting		
Pain			instrumer		self-care ADL		
General Disorde	ers						
Fatigue	none	Fatigue relieved by rest	-	ot relieved by rest; strumental ADL	Fatigue not relieved by rest, limiting self-care ADL		
Respiratory, the	oracic and r	nediastinal disorders					
Cough	none	Mild symptoms; nonprescription intervention indicated	medical ir indicated; instrumer	ital ADL	Severe symptoms; limiting self-care ADL		
Dyspnea	none	Shortness of breath with moderate exertion		of breath with xertion; limiting tal ADL	Shortness of breath at rest; limiting self-care ADL	Life-threatening consequences; urgent intervention indicated	Death



Pleuritic pain	none	Mild pain	Moderate pain; limiting instrumental ADL	Severe pain; limiting self-care ADL		
Pneumonitis	none	Asymptomatic; clinical or diagnostic observations only; intervention not indicated	Symptomatic; medical intervention indicated; limiting instrumental ADL	Severe symptoms; limiting self-care ADL; oxygen indicated	Life-threatening respiratory compromise; urgent intervention indicated (e.g., tracheotomy or intubation)	Death
ECOG Performan	ce Status*					
	Fully active	Restricted in physically strenuous activity but ambulatory and able to carry out work of a light or sedentary nature	Ambulatory and capable of all self-care but unable to carry out any work activities. Up and about more than 50% of waking hours	Capable of only limited self-care, confined to bed or chair more than 50% of waking hours	Completely disabled. Cannot carry on self-care. Totally confined to bed or chair	Dead

L9 Physician: Lung Clinical Outcomes Follow up					
Time points: Follow-up visits 1-3-6 month					
Data Elements	Options				
Weight	(specify lbs. or kg)				
Rate current esophageal pain on a scale of 0-10:	(0-10)				
Please select what you have recommended to the patient in the past month to manage esophagitis:	 None Non-narcotic prescription medications (e.g. Magic mouthwash, Miles mixture, xylocaine) Narcotic prescription medications (e.g. Vicodin, Percocet, morphine, oxycodone, fentanyl) Other intervention (specify): 				



MROQC Lung Data Elements Guide

-		ion of the primar diotherapy?	y tumor	• YES/î	NO		
a) If YES,	date per	formed:		(dat	e)		
b) If YES, what was the surgical margin status?			PositNega				
Disease status: (check one			 No ev disea Local Dista 	vidence of disease vidence of progressic se /Regional progressio nt progression (Local & Distant prog	n		
Has the patient been admitted for a cardiac event? a) If YES, date of admission: b) Select the cardiac event related to admission				CongPeric			
Toxicity Sco	oring (CTC/	AE v 4.0). Please ci	rcle one n	umber in each ro	w.		
Adverse Event	0	1		2	3	4	5
Gastrointestin Esophagitis	al disorders none	Asymptomatic; clinical or diagnostic observations only;	Symptomatic; altered eating/swallowing; oral supplements indicated		Severely altered eating/swallowing; tube feeding, TPN or	Life-threatening consequences; urgent intervention indicated	Death



		intervention not			hospitalization		
		indicated			indicated		
		maleatea			maleated		
Esophageal	none	Mild pain	Moderate pai		Severe pain; limiting		
Pain			instrumental	ADL	self-care ADL		
General Disor		1					1
Fatigue	none	Fatigue relieved by	-	elieved by rest;	Fatigue not relieved by		
		rest	limiting instru	imental ADL	rest, limiting self-care		
	<u> </u>				ADL		
		mediastinal disorders					
Cough	none	Mild symptoms;	,	nptoms, medic	, , ,		
		nonprescription intervention	intervention i limiting instru		limiting self-care ADL		
		indicated	initia institu				
Dyspnea	none	Shortness of	Shortness of	breath with	Shortness of breath at	Life-threatening	Death
,		breath with	minimal exer		rest; limiting self-care	consequences; urgent	
		moderate exertion	instrumental		ADL	intervention indicated	
Pleuritic	none	Mild pain	Moderate pai	in; limiting	Severe pain; limiting		
pain			instrumental ADL		self-care ADL		
Pneumonitis	none	Asymptomatic;	Symptomatic; medical		Severe symptoms;	Life-threatening respiratory	Death
		clinical or	intervention i	,	limiting self-care ADL;	compromise; urgent intervention indicated (e.g.,	
		diagnostic	limiting instru	imental ADL	oxygen indicated	tracheotomy or intubation)	
		observations only; intervention not				the leadening of intrabution,	
		indicated					
ECOG Performa	nce Status*	indicated					
	Fully	Restricted in	Ambulatory a	ind capable of a	Capable of only limited	Completely disabled. Cannot	Dead
	active	physically	self-care but	unable to carry	self-care, confined to	carry on self-care. Totally	
		strenuous activity	out any work	activities. Up	bed or chair more than	confined to bed or chair	
		but ambulatory	and about mo	ore than 50% o	50% of waking hours		
		and able to carry	waking hours				
		out work of a light					
		or sedentary					
		nature, e.g., light					
		housework, office					
		work					
		L10 Phys	sician: Lu	ng Clinic	al Outcomes Fol	low up	
					Time noir	nts: Follow-up visits 1-3-6	months
				0	rinic poir		
Data Elem	ents			Options			
Did the patient receive systemic <u>therapy after</u> receiving radiation therapy?			• Y	ES			
				0			
receiving radiation therapy?			- N	0			
If yes, che	ck all tha	t apply:					
		- ~phil.					
				1			



Chemotherapy	Agents (check all that apply): Cisplatin/Platinol-AQ/Platinol Etoposide/Eposin/VePesid/VP-16 Paclitaxel/Taxol/TAX Carboplatin/Paraplatin/Paraplatin-AQ Pemetrexed/Alimta Docetaxel/Taxotere/TXT Gemcitabine/Gemzar Vinorelbine/Navelbine Other (specify)
Immunotherapy	Agents (check all that apply): • Atezolizumab • Durvalumab • Nivolumab • Pembrolizumab • Other (specify)
L11 Lung:	Annual Clinical Outcomes
	Time points: Annually
Data Elements	Options
Since the end of treatment, has the patient been admitted for a cardiac event?	• YES/NO
If YES, date of admission 1	(date)
Select the cardiac event related to admission 1:	 Arrhythmia Congestive Heart failure Pericardial Effusion Myocardial Infarction None of the above



Since the end of treatment, has the patient been admitted for a cardiac event?	• YES/NO
If YES, date of admission 2	(date)
Select the cardiac event related to admission 2:	 Arrhythmia Congestive Heart failure Pericardial Effusion Myocardial Infarction None of the above
Since the end of treatment, has the patient been admitted for a lung event?	• YES/NO
If YES, date of admission 1	(date)
Select the lung event related to admission 1	 Chronic obstructive pulmonary disease-exacerbation Pneumonia Pneumonitis None of the above



Since the end of treatment, has the patient been admitted for a lung event?	• YES/NO
If YES, date of admission 2	(date)
Select the lung event related to admission 2	 Chronic obstructive pulmonary disease-exacerbation Pneumonia Pneumonitis None of the above
Disease status: (check one	 No evidence of disease No evidence of progression of disease Local/Regional progression Distant progression Both (Local & Distant progression)
Has the patient received any thoracic RT after their initial treatment?	• YES/NO



Has the patient tested positive for COVID- 19?	• YES/NO
If YES, date of positive diagnosis:	 (date) Date not available
If YES, was the patient:	SymptomaticAsymptomaticNot documented

SE2 CDA: Early Termination of MROQC Patient Participation Form

Data Elements	Options
Date of Early Termination:	If the patient died this would be the date of death otherwise it is the last eval. date the patient had
Reason patient is no longer being followed /participating in MROQC:	 Moved To continue treatment elsewhere Deceased Hospice Metastatic disease Patient chose to stop treatment Patient not returning to RT department for follow up Medical issues (i.e. CVA, MI) prevent further participation Lung only- annual documentation not available



LUNG Radiotherapy Technical Details Form	
Simulation Which lung has the primary tumor?	RightLeft
Was intravenous contrast used for the patient's treatment planning simulation?	YesNo
Select the primary method used to assess the motion of the tumor and organs-at-risk during simulation.	 DCT Fluoroscopy Slow CT Motion not assessed Scans at multiple breath hold states Other. Please specify:
Targets Which modalities were used for target delineation? Only choose datasets which were registered and fused to the treatment planning scan. Check all that apply.	 CT Diagnostic CT PET MRI
How was motion accounted for during the treatment of this patient?	 ITV approach: no motion control technique was applied, but the target volumes were designed to account for breathing motion (using 4DCT, scans at multiple breath hold states, slow CT, etc.) Voluntary breath hold without a device Breath hold with a device (ABC, SDX, etc.) Gating of radiotherapy (RPM, AlignRT, etc.) Abdominal compression Motion was not taken into account while designing volumes or by a motion management technique Other. Please specify:



Г

What was the reason for not considering motion in accordance with the MROQC target delineation guidelines?	 DCT is not available at treating institution Use of slow CT was not feasible, due to time constraints or experience with technique Other. Please specify:
Was patient specific reproducibility testing performed to ensure the breath hold position was reliable? [If " Voluntary breath hold without a device" or "Breath hold with a device (ABC, SDX, etc.)" or "Abdominal compression"]	YesNo
Was motion considered in the delineation of target volumes? [If "Other. Please specify:"]	YesNo
Was a motion encompassing GTV (IGTV) structure contoured?	YesNo
Select the name of the GTV structure:	 Drop-down menu: GTV, GTVp, IGTV, Other. Please specify:
Was a CTV or ICTV structure contoured?	YesNo
Enter the volume of the IGTV (GTV) in cc:	cc
Enter the reason(s):	 Institutional practice Physician preference Ambiguity in imaging/could not define Other. Please specify:
Was a motion encompassing CTV or ICTV structure defined/contoured?	YesNo



Select the name of the CTV structure:	 Drop-down menu: CTV, CTVp, CTV_High, ICTV, Other. Please specify:
Enter the volume of the ICTV (CTV) in cc	cc
What is the approximate margin between the IGTV (GTV)structure and ICTV (CTV) structure in cm?	cm
Enter the reason(s):	 Institutional practice Physician preference Ambiguity in imaging/could not define Other. Please specify:
Was a PTV structure defined?	YesNo
Select the name of the PTV structure:	Drop-down menu: PTV, PTVp, PTV_High, Other. Please specify:
Enter the volume of the PTV in cc	cc
What is the approximate margin between the CTV structure (or GTV structure if CTV structure was not defined) and PTV structure in cm?	cm
Enter the reason(s):	 Institutional practice Physician preference Ambiguity in imaging/could not define Other. Please specify:



Treatment Planning	
Do any of these structures overlap with a 2 cm expansion of the PTV? Check all that apply.	 Spinal cord Brachial plexus Heart Other structure of interest. Please specify: Esophagus No, the PTV is greater than 2 cm from all other structures
Select the number of plans treated	• drop-down menu: 1-10
For each plan, specify:	
 a. Planning type 	Forward planningInverse planning
b) Dose delivered with this plan (Gy	• between 1 and 90
 C) Number of fractions delivered with this plan 	• between 1 and 40
d. Was the patient treated BID?	YesNo
 e. Treatment region 	 Primary target Primary target & nodes Nodes



Reason for plan	Initial
	Planned Boost
	Planned Adaptation
	Unplanned Modification
f) If not initial, what was the reason?	• Minimize dose to critical structures (e.g. off-cord or off
	brachial plexus boost)
	• Patient anatomy change (e.g. lung inflation, pleural effusion
	change)
	 Change in motion management strategy
	Other. Please specify:
g. Was this plan considered SBRT?	Mar .
0	• Yes
	• No
 h) Did this plan include a concomitant 	
boost?	Yes
DOOSLE	• No
	Gy (between 1 and 90)
j) If no, enter planned dose:	/ \
k) If no, enter planned number of	Gy (between 1 and 40)
fractions:	
Treatment Delivery and Image Guidance	
ineatment Denvery and image Guiuance	
What tupo of imaging was used to verify this	a W//W/ portal
What type of imaging was used to verify this	kV/MV portal
patient's setup?	CT (CBCT or TomoTherapy CT)
	• Films
	Video-based system
	Other. Please specify:



Г

For each imaging type, specify how often the patient was imaged during treatment.	 Daily Less than daily but more than weekly Weekly Other. Please specify:
---	---