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V1.0		HCMI
	Enrollment: Lung	
Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	
Completed By:	Completion Date (MM/DD/YYYY):	

Form Notes: An Enrollment Form should be completed for each HCMI case upon qualification notice from Leidos. All information provided on this form should include activity from the Date of Initial Pathologic Diagnosis to the most recent Date of Last Contact with the patient.

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
1	ID2		2003301	Provide the patient's ID2 (this ID will only be
				used by IMS for internal quality control).
2	ID3		5845012	Provide the HCMI-specific anonymized ID (ID3).
3	Index date	☐ Initial pathologic diagnosis ☐ Sample procurement ☐ First patient visit	6154722	Select the reference date used to calculate time intervals (e.g. days to treatment). Date of initial pathologic diagnosis is the HCMI standard and should be used unless it is unavailable. If an alternative index date is used, indicate it here and use it for all interval calculations.
Patient Inf			T	T =
4	Gender	☐ Male ☐ Female ☐ Unspecified	2200604	Provide the patient's gender using the defined categories. Identification of gender is based upon self-report and may come from a form, questionnaire, interview, etc.
5	Height		649	Provide the patient's height, in centimeters.
6	Weight		651	Provide the patient's weight, in kilograms.
7	Body mass index (BMI)		2006410	If the patient's height and weight are not collected, provide the patient's body mass index (BMI).
8	Race	☐ American Indian or Alaska Native ☐ Asian ☐ Black or African American ☐ Native Hawaiian or other Pacific Islander ☐ White ☐ Unknown ☐ Not reported	2192199	Provide the patient's race using the defined categories. American Indian or Alaska Native: A person having origins in any of the original peoples of North and South America (including Central America), and who maintains tribal affiliation or community attachment. Asian: A person having origins in any of the peoples of the Far East, Southeast Asia, or in the Indian subcontinent including, for example, Cambodia, China, India, Japan, Korea, Malaysia, Pakistan, the Philippine Islands, Thailand, and Vietnam. Black or African American: A person having origins in any of the black racial groups of Africa. Native Hawaiian or other Pacific Islander: A person having origins on any of the original peoples of Hawaii, Guam, Samoa, or other Pacific Island. White: A person having origins in any of the original peoples of Europe, the Middle East, or North Africa.
9	Ethnicity	☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown ☐ Not reported	2192217	Provide the patient's ethnicity using the defined categories. Hispanic or Latino: A person of Mexican, Puerto Rican, Cuban, Central or South American or other Spanish culture or origin, regardless of race. Not Hispanic or Latino: A person not meeting the definition of Hispanic or Latino.
10	Number of days from index date to date of last contact		3008273	Provide the number of days from the index date to the date of last contact.

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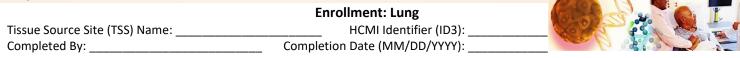
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Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	CAS SA	3
Completed By:	Completion Date (MM/DD/YYYY):	1000	-

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
10	Number of days		3008273	Provide the number of days from the index
	from index date to			date to the date of last contact.
	date of last contact			
11	Patient age on		6379572	Provide the age (in days) of the patient on
	index date			the index date. Note: If the patient's age is greater than 32,872
				days (90 years), please enter 32,872.
12	Year of birth		2896954	Provide the year of the patient's birth. If the
				patient was born prior to 1928, insert the
				date 1928.
13	Family history of	☐ Same	5832923	Has a first-degree relative of the patient
	cancer	☐ Different		been diagnosed with a cancer of the same
		□ None		or a different type?
		☐ Unknown		
14	Smoking history	☐ Lifelong non-smoker (<100 cigarettes	2181650	Indicate the patient's history of tobacco
14	Silloking history	smoked in a lifetime)	2181030	smoking as well as their current smoking
		☐ Current smoker (includes daily and non-		status using the defined categories.
		daily smokers)		
		☐ Current reformed smoker (duration not		
		specified)		
		☐ Current reformed smoker for >15 years		
		☐ Current reformed smoker for ≤15 years	225525	
15	Number pack years		2955385	Indicate the number of pack years smoked.
	smoked			Note: Numeric computed value to represent lifetime tobacco exposure defined as number of
				cigarettes smoked per day x number of years
				smoked divided by 20.
16	Tobacco smoking		2228610	Indicate the year in which the patient quit
	quit year			smoking.
17	Asbestos exposure?	☐ Yes	1253	Indicate whether the patient has been
	Assested exposure.	□ No	1233	exposed to asbestos.
		☐ Unknown		'
18	Radon exposure?	☐ Yes	2816352	Indicate whether the patient has been
		□ No		exposed to radon.
		☐ Unknown		
19	Metastasis at		3438571	Indicate whether there was evidence of
	diagnosis	☐ Metastatic		metastasis at the time of diagnosis of the
	assessment status	☐ Non-metastatic (confirmed) ☐ Non-metastatic (unconfirmed)		primary tumor. Note: If metastatic at diagnosis, proceed to Question 20, otherwise,
		Non metastatic (uncommica)		skip to Question 21.
20	Metastatic site(s) at	☐ Adrenal gland	3108271	Indicate the site(s) of metastasis at the time
	diagnosis	☐ Bone		of diagnosis of the primary tumor.
		☐ Brain		Note: If the anatomic site of metastasis is not
		Liver		listed, proceed to Question 20a, otherwise, skip to Question 21.
		Lung		to question 21.
		Lymph node(s)- non-regional		
20a	Specify metastatic	Other (specify)	3128033	If the site of metastasis is not included on
200	site(s)		3120033	the provided list, specify the site of
	2,00(0)			metastasis.
Biospecimo	en Information			
21	Tissue sample	☐ Normal tissue	2006911	Please select all the tissue sample types
	type(s) collected for	☐ Primary tumor		submitted for HCMI with this case.
	HCMI for this case	☐ Metastatic		
		☐ Recurrent		
		☐ Other tissue		
			1	



Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):
Completed By:	Completion Date (MM/DD/YYYY):

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
22	Number of		6584256	Please provide the number of normal tissue
	NORMAL tissues			specimens obtained for HCMI for this case.
	biospecimens			Note: This number is expected to be 1.
	collected for HCMI			
	for this case			
23	Number of		6584257	Please provide the number of primary
	PRIMARY cancer			tumor specimens obtained for HCMI for this
	tissue biospecimens			case. Note: A single primary tumor biospecimen
	collected for HCMI			obtained that is portioned for both sequencing
	model			and model generation counts as 1 single primary
	development for			tumor specimen. This number is expected to be 1.
	this case			1.
24	Number of		6584258	Please provide the number of metastatic
	METASTATIC/			and/or recurrent cancer biospecimens
	RECURRENT cancer			collected for HCMI for this case. <i>Note: A</i>
	tissue biospecimens			biospecimen obtained from a single site at a single
	collected for HCMI			timepoint in progression that is portioned for both sequencing and model generation counts as 1 single
	model			tumor specimen. A biospecimen obtained from
	development for			another site or at a later timepoint in progression
	this case			that is portioned for both sequencing and model
				generation counts as a second single tumor
25	Number of OTHER		6584259	specimen. Please provide the number of pre-
23	tissue biospecimens		0304233	malignant, non-malignant, or dysplastic
	collected for HCMI			tissue biospecimens collected for HCMI for
	model			this case. Note: A biospecimen obtained from a
	development for			single site at a single timepoint in progression
	this case			that is portioned for both sequencing and model
	tilis case			generation counts as 1 single tumor specimen. A
				biospecimen obtained from another site or at a
				later timepoint in progression that is portioned
				for both sequencing and model generation
26	Total number of		CE04271	counts as a second single tumor specimen.
26			6584271	Please provide the total number of tissue biospecimens collected for HCMI for this
	tissue biospecimens collected for HCMI			·
	for this case			case. Note: This number should be the sum of the normal, primary tumor, metastatic/recurrent
	TOT LITTS CASE			tumor, and other biospecimen counts above.
Normal Co	ntrol Information			
27	Normal tissue		6584264	Please provide a number to identify
	biospecimen			which biospecimen this is in the sequence.
	ordinal			Note: The first biospecimen should be number
				"1," the second should be number "2," etc.
28	CMDC sample ID		6586035	Please provide the CMDC sample ID for this
				biospecimen as it will appear on tubes and
				the Sample Submission Form transmitted to
				the BPC.
29	BPC submitter ID (if		6584919	Please provide the BPC-generated ID for this
	available)			sample as it will appear on the Sample
				Submission Form transmitted to the BPC.
30	Type of normal	☐ Whole blood	3081936	Indicate the type of normal control
	control	☐ Buccal cells		submitted for this case.
		☐ Buffy coat		
		☐ Lymphocytes		
		☐ Extracted DNA from blood		
		☐ Extracted DNA from saliva		
		☐ Extracted DNA from buccal cells		
		☐ Extracted DNA from normal tissue		
		☐ FFPE non-neoplastic tissue		
		☐ Non-neoplastic tissue		



normal tissue	ssue was submitted as , select the anatomic site e. Note: If the anatomic site
normal tissue	, select the anatomic site e. Note: If the anatomic site
☐ Left upper lobe lung ☐ Right lower lobe lung ☐ Right middle lobe lung ☐ Right middle lobe lung ☐ Question 31a, otherw	e. Note: If the anatomic site
☐ Right lower lobe lung	-
☐ Right middle lobe lung Question 31a, otherw	
I I RIGHT LINNER LONG LILING I I I I I I I I I I I I I I I I I I	vise, skip to Question 32.
Other (specify)	
□ Not applicable	
	ssue, adjacent tissue, or
	another anatomic site
was submitted as the	,
	nic site of the normal
tissue.	
,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	ce from the site of normal
	the primary tumor. <i>Note:</i>
	not submitted, select 'Not
not sloca) — Not applicable	
I I I I I I I I I I I I I I I I I I I	d used to preserve the
sample D FEDE	
preservation	rization.
method	
Primary Tumor Biospecimen Information	anda familia a satur
	code for the primary
	enerate the ID3 for this
	ICD-10 code is not listed,
	rwise, skip to Question 35.
	for the tumor used to
	el submitted to HCMI is
	e provided list, specify the
ICD-10 code.	
2 0032/3	pathology/laboratory
G 0070/3 G 0253/3	ICD-O-3 histology code
1 00/3/3 0 0200/3	or. Note: If the ICD-O-3
□ 8083/3 □ 8490/3 proceed to Question	primary tumor is not listed, 35a, otherwise, skip to
8084/3	33u, otherwise, skip to
□ 8230/3 □ Other (specify)	
35a Specify other 3226275 If the ICD-O-3 histo	logy code describing the
	patient's primary tumor
	the previous list, provide
the ICD-O-3 histolo	
	pathology/laboratory
origin report select the p	
Lung disease Mater Mater	primary site of the disease
1 I I Uner (Specify)	to Question 36a, otherwise
skip to Question 37.	
36a Other tissue or 5946219 If the primary site of	of the disease is not
organ of origin included on the pre	evious list, provide the
primary site of the	disease.
37 Histological Type 3081932 Select the surgical page 3081932	pathology text
	nistological tumor type.
description of the h	cal tumor tuno is not listed
☐ Lung cancer description of the h	cui cumor cype is not listea,
☐ Lung cancer ☐ Other (specify) ☐ Ung cancer ☐ Other (specify) ☐ Description of the Proceed to Question of Question of the Proceed to Question of Questi	37a, otherwise, skip to
☐ Lung cancer ☐ Other (specify) ☐ Other (specif	37a, otherwise, skip to
Ung cancer ☐ Other (specify) Other histological Dung cancer ☐ Other (specify) Other (specify) description of the histological Note: If the histological proceed to Question 38. 3294805 If the traditional su	37a, otherwise, skip to
Ung cancer ☐ Other (specify) Cher (specify) Description of the home in the	argical pathology text histological tumor type is
Ung cancer ☐ Other (specify) Cher (specify)	argical pathology text nistological tumor type is e previous list, please
Lung cancer Other (specify) Other (specify) Other histological type Lung cancer Note: If the histologic proceed to Question 38. 3294805 If the traditional su description of the histological type	argical pathology text nistological tumor type is e previous list, please

	Enrollment: Lung	TO THE	A CT	200	Contract of the second	7,
Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	1000	5	30		1
Completed By:	Completion Date (MM/DD/YYYY):	160	6 8	2000		-

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
38	Histological subtype	Acinar adenocarcinoma Adenocarcinoma in situ; mucinous Adenocarcinoma in situ; nonmucinous Adenosquamous carcinoma Basaloid squamous cell carcinoma Colloid adenocarcinoma Enteric adenocarcinoma Fetal adenocarcinoma Keratinizing squamous cell carcinoma Lepidic adenocarcinoma Micropapillary adenocarcinoma Minimally invasive adenocarcinoma; mucinous Minimally invasive adenocarcinoma; nonmucinous Nonkeratinizing squamous cell carcinoma Papillary adenocarcinoma Signet ring adenocarcinoma Solid adenocarcinoma Adenocarcinoma Colid adenocarcinoma Colloid ade	3081934	Using the patient's pathology/laboratory report, select the histological subtype of the primary tumor. Note: If the histological subtype is not listed, proceed to Question 38a, otherwise, skip to Question 39.
38a	Other histological subtype		3124492	If the histological subtype for the primary tumor is not included in the provided list, specify the histological subtype.
39	Prior malignancy (of the same cancer type)	☐ Yes ☐ No ☐ Unknown	5832924	Indicate whether the patient has a history of prior malignancy of the same cancer type.
40	Prior malignancy (other cancer type)	☐ Yes ☐ No ☐ Unknown	5878828	Indicate whether the patient has a history of prior malignancy of a different cancer type.
41	AJCC cancer staging edition	☐ 1 st ☐ 4 th ☐ 7 th ☐ 8 th ☐ 8 th	2722309	Select the AJCC staging handbook edition used to stage the patient's primary tumor.
42	Clinical stage group	□ Stage 0 □ Stage IIB □ Stage IA □ Stage III □ Stage IA1 □ Stage IIIA □ Stage IA2 □ Stage IIIB □ Stage IA3 □ Stage IIIC □ Stage IB □ Stage IV □ Stage II □ Stage IVA □ Stage IIA □ Stage IVB	3440332	Using the patient's pathology/laboratory report, select the clinical stage group of the primary tumor as defined by the American Joint Committee on Cancer (AJCC).
43	AJCC pathologic spread: Primary tumor (pT)	☐ T0 ☐ Tis ☐ Tis ☐ Tis (SCIS) ☐ Tis (AIS) ☐ T3 ☐ T1mi ☐ T1a ☐ T1b ☐ TX	3045435	Using the patient's pathology/laboratory report, select the code for the pathologic T (primary tumor) as defined by the American Joint Committee on Cancer (AJCC).
44	AJCC pathologic spread: Lymph nodes (pN)	□ N0 □ N3 □ N3 □ NX □ N2	3203106	Using the patient's pathology/laboratory report, select the code for the pathologic N (nodal) as defined by the American Joint Committee on Cancer (AJCC).
45	AJCC pathologic spread: Distant metastases (pM)	□ M0 □ M1b □ M1 □ M1c □ M1a □ MX	3045439	Using the patient's pathology/laboratory report, select the code for the pathologic M (metastasis) as defined by the American Joint Committee on Cancer (AJCC).

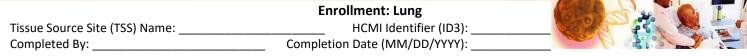
V1.0		HCMI
	Enrollment: Lung	
Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	
Completed By:	Completion Date (MM/DD/YYYY):	

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
46	Tumor stage (pathological)	□ Stage 0 □ Stage IA □ Stage IA □ Stage IA1 □ Stage IB □ Stage IB □ Stage IB □ Stage III □ Stage III □ Stage III □ Stage IV □ Stage III □ Stage IV □ Stage III □ Stage IV	3065862	Using the patient's pathology/laboratory report, in conjunction with the patient's medical record, select the stage as defined by the American Joint Committee on Cancer (AJCC).
47	Tumor grade	☐ G1-Well differentiated ☐ G2-Moderately differentiated ☐ G3-Poorly differentiated ☐ G4-Undifferentiated ☐ GB-Borderline histologic grade ☐ GX-Unknown	2785839	Using the patient's pathology/laboratory report, select the grade of the primary tumor.
Prognostic		eatures for Primary Tumor Prognosis or Responsi		eatment
48	Was KRAS mutation analysis performed?	☐ Yes☐ No☐ Unknown	6060079	Indicate whether KRAS mutation analysis was performed. Note: If KRAS mutation analysis was not performed, skip to Question 51.
49	Was a mutation in KRAS identified?	☐ Yes ☐ No	6060081	Indicate whether a mutation in KRAS was identified through mutation analysis. <i>Note: If KRAS mutation was not identified, skip to Question 51.</i>
50	If KRAS mutation identified, which one?	□ G12A □ G12V □ G13V □ Q61H □ G12C □ G13A □ Q61L □ G12D □ G13C □ A146T □ G12S □ G13R □ Other (specify)	6060083	Indicate the specific KRAS mutation identified. Note: If the KRAS mutation is not listed, proceed to Question 50a, otherwise, skip to Question 51.
50a	Other KRAS mutation(s)		6101691	If the KRAS mutation identified is not provided in the previous list, specify the KRAS mutation.
51	Was EGFR mutation analysis performed?	☐ Yes ☐ No ☐ Unknown	6063955	Indicate whether EGFR mutation analysis was performed. Note: If EGFR mutation analysis was not performed, skip to Question 54.
52	Was a mutation in EGFR identified?	☐ Yes ☐ No	6063530	Indicate whether a mutation in EGFR was identified through mutation analysis. <i>Note: If EGFR mutation was not identified, skip to Question 54.</i>
53	If EGFR mutation identified, which one?	□ G719X □ L861Q □ T790M □ Exon 19 deletion □ C797S □ Exon 20 insertion □ L858R □ Other (specify)	3147627	Indicate the specific EGFR mutation identified. Note: If the EGFR mutation is not listed, proceed to Question 53a, otherwise, skip to Question 54.
53a	Other EGFR mutation(s)		4173882	If the EGFR mutation identified is not provided in the previous list, specify the EGFR mutation.
54	Was BRAF mutation analysis performed?	☐ Yes☐ No☐ Unknown	6061813	Indicate whether BRAF mutation analysis was performed. Note: If BRAF mutation analysis was not performed, skip to Question 57.
55	Was a mutation in BRAF identified?	☐ Yes ☐ No	6061809	Indicate whether a mutation in BRAF was identified through mutation analysis. <i>Note: If BRAF mutation was not identified, skip to Question 57.</i>
56	If BRAF mutation identified, which one?	□ V600D □ V600R □ V600E □ K601E □ V600K □ Other (specify)	6061810	Indicate the specific BRAF mutation identified. Note: If the BRAF mutation is not listed, proceed to Question 56a, otherwise, skip to Question 57.
56a	Other BRAF mutation		6101687	If the BRAF mutation identified is not provided in the previous list, specify the BRAF mutation.

Enrollment: Lung

Tissue Source Site (TSS) Name: _____ HCMI Identifier (ID3): _____
Completion Date (MM/DD/YYYY): ____

Question	Question Text	Data Entry Options		CDE ID	Instruction Text
57	Was MET mutation analysis performed?	☐ Yes ☐ No ☐ Unknown		6078874	Indicate whether MET mutation analysis was performed. Note: If MET mutation analysis was not performed, skip to Question 60.
58	Was a mutation in MET identified?	☐ Yes ☐ No		6078886	Indicate whether a mutation in MET was identified through mutation analysis. Note: If MET mutation was not identified, skip to Question 60.
59	If MET mutation identified, which one?	☐ D963 splice mutation ☐ D1010N ☐ D1010 splice mutation ☐ Intron 13 mutation	☐ Intron 14 mutation ☐ Exon 14 mutation ☐ Exon 14 deletion ☐ Other (specify)	6078901	Indicate the specific MET mutation identified. Note: If the MET mutation is not listed, proceed to Question 59a, otherwise, skip to Question 60.
59a	Other MET mutation		-	6230525	If the BRAF mutation is not included in the list provided, specify the MET mutation identified.
60	Was EML4-ALK translocation analysis performed?	☐ Yes ☐ No ☐ Unknown		6063976	Indicate whether EML4-ALK translocation analysis was performed. Note: If EML4-ALK translocation analysis was not performed, skip to Question 64.
61	Was an EML4-ALK translocation identified?	☐ Yes ☐ No		6063959	Indicate whether an EML4-ALK translocation was identified. Note: If EML4-ALK translocation was not identified, skip to Question 64.
62	If EML4-ALK translocation found, which variant?	☐ Variant 1 ☐ Variant 2 ☐ Variant 3	☐ Variant 4 ☐ Variant 5	3139445	Indicate the EML4-ALK translocation variant that was identified.
63	What method was used to identify the EML4-ALK translocation	☐ Cytogenetics☐ FISH☐ NGS	☐ IHC (nuclear staining) ☐ RT-PCR ☐ Other	3139449	Specify the method used to identify the EML4-ALK translocation variant.
64	Was ROS1 FISH performed?	☐ Yes ☐ No ☐ Unknown		6078871	Indicate whether fluorescence in situ hybridization (FISH) analysis of the ROS1 gene was performed. Note: If ROS FISH was not performed, skip to Question 66.
65	Was ROS1 rearrangement detected by FISH?	☐ Yes ☐ No		6078881	Indicate whether a ROS1 rearrangement was identified by fluorescence in situ hybridization (FISH).
66	Was RET gene rearrangement analysis performed?	☐ Yes ☐ No ☐ Unknown		6078872	Indicate whether RET gene rearrangement analysis was performed. Note: If RET rearrangement analysis was not performed, skip to Question 68.
67	Was a RET rearrangement identified?	☐ Yes ☐ No		6078882	Indicate whether a RET rearrangement was identified.
68	Was PD-L1 IHC performed?	☐ Yes ☐ No ☐ Unknown		6078873	Indicate whether PD-L1 expression was assessed by immunohistochemistry (IHC). Note: If PD-L1 IHC was not performed, skip to Question 70.
69	PD-L1 expression by IHC	☐ Positive☐ Negative☐ Equivocal		4798631	Indicate the status of PD-L1 protein expression as determined by immunohistochemistry (IHC).
70	Was HER2 IHC performed?	☐ Yes ☐ No ☐ Unknown		6063454	Indicate whether HER2 expression was assessed by immunohistochemistry (IHC). Note: If HER2 IHC was not performed, skip to Question 72.



Question	Question Text	Data Entry Options	CDE ID	Instruction Text
71	HER2 expression by	☐ Positive	2957563	Indicate the expression of HER2 as assessed
	IHC	☐ Negative		by immunohistochemistry (IHC).
		☐ Equivocal		
Primary Tu	mor Sample Informati	on	7	
72	Are you submitting a primary tumor tissue sample for this case?	☐ Yes ☐ No		If yes, proceed to question 73. If submitting a metastatic/recurrent tumor sample, proceed to Question 99.
73	Primary tumor biospecimen ordinal		6584265	Please provide a number to identify which biospecimen this is in the sequence. <i>Note:</i> This number should be "1".
74	CMDC sample ID		6586035	Please provide the CMDC sample ID for this biospecimen as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
75	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
76	Sample represents primary diagnosis?	☐ Yes ☐ No	6584730	Does this primary tumor specimen represent the PRIMARY DIAGNOSIS for this Case ID3? Note: If no, continue to Question 77, otherwise, skip to Question 78.
77	Specify the ICD-10 code		3226287	Provide the ICD-10 code for the primary tumor used to generate the model submitted to HCMI.
78	Tumor tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the tumor tissue sample collected for molecular characterization.
79	Anatomic Site of tumor from which Model was Derived	□ Left lower lobe lung □ Left upper lobe lung □ Right lower lobe lung □ Right middle lobe lung □ Right upper lobe lung □ Adrenal gland □ Bone □ Brain □ Bronchus □ Liver □ Lymph node □ Other (specify)	4214629	Select the anatomic site of the tumor tissue sample used to generate the model for HCMI. Note: If the tissue or organ of origin is not listed, proceed to Question 79a. Otherwise, skip to Question 80.
79a	Other anatomic site from which the tumor was obtained		5946219	If not provided in the previous list, provide the anatomic site of the tumor tissue sample used to generate the model for HCMI.
80	Method of cancer sample procurement	 □ Cytology □ Fine needle aspiration biopsy □ Incisional biopsy □ Excisional biopsy □ Tumor resection □ Other (specify) 	3103514	Provide the procedure performed to obtain the primary tumor tissue. Note: If the method of procurement is not listed, proceed to Question 80a, otherwise, skip to Question 81.
80a	Specify the other method of tumor sample procurement		2006730	Specify the procedure performed to obtain the primary tumor tissue, if not included in the previous list.



Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	1000	K
Completed By:	Completion Date (MM/DD/YYYY):	100	

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
81	Number of days from index date to date of tumor sample procurement		3288495	Provide the number of days from the index date to the date of the procedure that produced the tumor tissue submitted for HCMI.
82	Tumor tissue type	□ Primary□ Additional Primary□ NOS	3288124	Provide the primary tumor tissue type for this sample.
	ımor Model Informatio	n		
83	Primary model biospecimen ordinal		6594596	Please provide a number to identify which biospecimen this is in the sequence. Note: This number is expected to be "1".
84	CMDC model ID		6586036	Please provide the CMDC model ID for this sample as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
85	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
86	Model represents primary diagnosis?	☐ Yes ☐ No	6584730	Does this MODEL represent the PRIMARY DIAGNOSIS for this Case ID3?
87	Model's primary tumor tissue CMDC sample ID		6586035	Enter the CMDC Sample ID of the PRIMARY TUMOR TISSUE from which this model is derived.
88	Model's primary tumor biospecimen ordinal		6584265	Enter the biospecimen ordinal of the PRIMARY TUMOR TISSUE from which this model is derived.
-	Information			
89	History of neoadjuvant treatment	 □ No □ Yes; radiation prior to resection □ Yes; pharmaceutical treatment prior to resection □ Yes; both radiation and pharmaceutical treatment prior to resection □ Unknown 	3382737	Indicate whether the patient received neoadjuvant radiation or pharmaceutical treatment. Note: Pharmaceutical therapy is addressed in Questions 90-96. Radiation therapy is addressed in Questions 97-98.
90	Neoadjuvant chemotherapy type	 □ Cytotoxic chemotherapy □ Hormonal □ Immunotherapy (cellular and immune checkpoint) □ Targeted therapy (small molecule inhibitors and targeted antibodies) □ Not applicable 	5832928	Select all neoadjuvant chemotherapy types that were administered to the patient. Note: Cytotoxic chemotherapy is addressed in Questions 90-92. Immunotherapy is addressed in Questions 93-94. Targeted therapy is addressed in Questions 95-96.
91	Neoadjuvant chemotherapeutic regimen	□ Albumin-bound paclitaxel □ Paclitaxel □ Pemetrexed □ Vinblastine □ Vinorelbine □ Docetaxel □ Other (specify) □ Chemotherapy not given	2853313	Select all chemotherapeutics used for neoadjuvant therapy. Note: If neoadjuvant chemotherapy was not given, skip to Question 93. If the neoadjuvant chemotherapeutic regimen is not listed, proceed to Question 91a, otherwise, skip to Question 92.
91a	Other neoadjuvant chemotherapeutic regimen		62694	If the neoadjuvant therapy is not included in the provided list, specify neoadjuvant therapies administered.
92	Days to neoadjuvant chemotherapy treatment from index date		5102411	Provide the number of days from index date to the date of treatment with neoadjuvant chemotherapy.

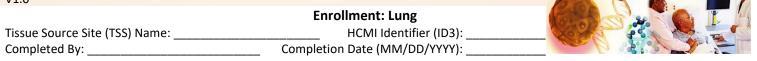
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ned, skip to Question 95. If the not listed, proceed to rwise, skip to Question 94. of the immunotherapy over of days from the index of treatment with the day therapy administered to If the targeted therapy d, proceed to Question 95a,
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herapy was not administered n 99. If the radiation therap
ed to Question 97a, otherwise
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erapy type is not included
st, specify the type.
per of days from the index
of treatment with radiation
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en was collected for this
es, proceed to Question 100
HER tissue sample, proceed t
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number to identify which
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en should be number "1", the
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
103	Metastatic/ recurrent tumor tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the metastatic/recurrent tumor tissue sample collected for molecular characterization.
104	Number of days from index date to date of diagnosis of metastasis/ recurrence		6132218	Provide the number of days from the index date to the date of diagnosis of metastatic/recurrent disease.
105	Method of metastatic/ recurrent cancer sample procurement	☐ Cytology ☐ Fine needle aspiration biopsy ☐ Incisional biopsy ☐ Excisional biopsy ☐ Tumor resection ☐ Other (specify)	6587389	Indicate the procedure performed to obtain the metastatic/recurrent tumor tissue. Note: If the method of procurement is not listed, proceed to Question 105a, otherwise, skip to Question 106.
105a	Other method of cancer sample procurement		6587390	If the procedure performed to obtain the tumor tissue is not included in the provided list, specify the procedure.
106	Number of days from index date to date of metastatic/ recurrent sample procurement		3288495	Provide the number of days from the index date to the date of the procedure that produced the metastatic/recurrent tumor tissue submitted for HCMI.
107	Metastatic/ recurrent site	☐ Left lower lobe lung	6587394	Select the site from which the metastatic/ recurrent tissue used to develop the model was derived. Note: If the metastatic/recurrent site is not listed, proceed to Question 107a, otherwise, skip to Question 108.
107a	Other metastatic/ recurrent site		6587395	If not included in the previous list, specify the site from which the metastatic/recurrent tissue used to develop the model was derived.
108	Site of relapse	☐ Local ☐ Regional ☐ Distant ☐ Not applicable	2002506	If the primary tumor relapsed, provide the site of relapse.
109	ICD-10 code		3226287	Provide the ICD-10 code for the metastatic/recurrent tumor used to generate the model submitted to HCMI.
110	ICD-O-3 histology code		3226275	Provide the ICD-O-3 histology code describing the morphology of the metastatic/recurrent tumor used to generate the model submitted to HCMI.
111	Maintenance and/or consolidation therapy administered prior to collection of metastatic/ recurrent tissue		6119066	Provide the name(s) of the maintenance and/or consolidation therapy administered to the patient prior to the collection of the metastatic/recurrent tissue used to develop the model.



Question	Question Text	Data Entry Options	CDE ID	Instruction Text
112	Days to start of maintenance and/or consolidation therapy from index date		5102411	Provide the number of days from the index date to the date maintenance and/or consolidation therapy started.
113	Days to last known administration date of maintenance and/or consolidation therapy from index date		5102431	Provide the number of days from the index date to the last known date of maintenance and/or consolidation therapy.
114	Is the patient still receiving treatment?	☐ Yes ☐ No ☐ Unknown	6379568	Indicate whether the patient is still undergoing maintenance and/or consolidation therapy.
115	Disease status	□ No evidence of disease□ Progressive disease□ Stable disease□ Unknown	2188290	Provide the disease status following maintenance and/or consolidation therapy.
		eatures for Metastatic/Recurrent 1		
116	Was KRAS mutation analysis performed?	☐ Yes ☐ No ☐ Unknown	6060079	Indicate whether KRAS mutation analysis was performed. Note: If KRAS mutation analysis was not performed, skip to Question 119.
117	Was a mutation in KRAS identified?	☐ Yes ☐ No	6060081	Indicate whether a mutation in KRAS was identified through mutation analysis. <i>Note: If KRAS mutation was not identified, skip to Question 119.</i>
118	If KRAS mutation identified, which one?	☐ G12A ☐ G12V ☐ G12C ☐ G13A ☐ G12D ☐ G13C ☐ G12R ☐ G13D	□ G13V 6060083 □ Q61H □ Q61L □ A146T □ Other (specify)	Indicate the specific KRAS mutation identified. Note: If the KRAS mutation is not listed, proceed to Question 118a, otherwise, skip to Question 119.
118a	Other KRAS mutation(s)		6101691	If the KRAS mutation identified is not provided in the previous list, specify the KRAS mutation.
119	Was EGFR mutation analysis performed?	☐ Yes ☐ No ☐ Unknown	6063955	Indicate whether EGFR mutation analysis was performed. Note: If EGFR mutation analysis was not performed, skip to Question 122.
120	Was a mutation in EGFR identified?	☐ Yes ☐ No	6063530	Indicate whether a mutation in EGFR was identified through mutation analysis. Note: If EGFR mutation was not identified, skip to Question 122.
121	If EGFR mutation identified, which one?	☐ C797S ☐ Exor	Q 3147627 1 19 deletion 20 insertion er (specify)	Indicate the specific EGFR mutation identified. Note: If the EGFR mutation is not listed, proceed to Question 121a, otherwise, skip to Question 122.
121a	Other EGFR mutation(s)		4173882	If the EGFR mutation identified is not provided in the previous list, specify the EGFR mutation.
122	Was BRAF mutation analysis performed?	☐ Yes ☐ No ☐ Unknown	6061813	Indicate whether BRAF mutation analysis was performed. Note: If BRAF mutation analysis was not performed, skip to Question 125.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
123	Was a mutation in BRAF identified?	☐ Yes ☐ No	6061809	Indicate whether a mutation in BRAF was identified through mutation analysis. <i>Note: If BRAF mutation was not identified, skip to Question 125.</i>
124	If BRAF mutation identified, which one?	□ V600D □ V600R □ V600E □ K601E □ V600K □ Other (s	6061810 pecify)	Indicate the specific BRAF mutation identified. Note: If the BRAF mutation is not listed, proceed to Question 124a, otherwise, skip to Question 125.
124a	Other BRAF mutation		6101687	If the BRAF mutation identified is not provided in the previous list, specify the BRAF mutation.
125	Was MET mutation analysis performed?	☐ Yes ☐ No ☐ Unknown	6078874	Indicate whether MET mutation analysis was performed. Note: If MET mutation analysis was not performed, skip to Question 128.
126	Was a mutation in MET identified?	☐ Yes ☐ No	6078886	Indicate whether a mutation in MET was identified through mutation analysis. Note: If MET mutation was not identified, skip to Question 128.
127	If MET mutation identified, which one?	□ D963 splice mutation □ Intron 1 □ D1010N mutatio □ D1010 splice □ Exon 14 mutation □ Exon 14 □ Intron 13 □ Other (s	n mutation deletion pecify)	Indicate the specific MET mutation identified. Note: If the MET mutation is not listed, proceed to Question 127a, otherwise, skip to Question 128.
127a	Other MET mutation		6230525	If the MET mutation is not included in the list provided, specify the MET mutation identified.
128	Was EML4-ALK translocation analysis performed?	☐ Yes ☐ No ☐ Unknown	6063976	Indicate whether EML4-ALK translocation analysis was performed. Note: If EML4-ALK translocation analysis was not performed, skip to Question 132.
129	Was an EML4-ALK translocation identified?	☐ Yes ☐ No	6063959	Indicate whether an EML4-ALK translocation was identified. Note: If EML4-ALK translocation was not identified, skip to Question 132.
130	If EML4-ALK translocation found, which variant?	☐ Variant 1 ☐ Variant 2 ☐ Variant 3 ☐ Variant 4 ☐ Variant 5	3139445	Indicate the EML4-ALK translocation variant that was identified.
131	What method was used to identify the EML4-ALK translocation	Cytogenetics IHC (nuc staining FISH RT-PCR NGS Other		Specify the method used to identify the EML4-ALK translocation variant.
132	Was ROS1 FISH performed?	☐ Yes ☐ No ☐ Unknown	6078871	Indicate whether fluorescence in situ hybridization (FISH) analysis of the ROS1 gene was performed. Note: If ROS FISH was not performed, skip to Question 134.
133	Was ROS1 rearrangement detected by FISH?	☐ Yes ☐ No	6078881	Indicate whether a ROS1 rearrangement was identified by fluorescence in situ hybridization (FISH).
134	Was RET gene rearrangement analysis performed?	☐ Yes ☐ No ☐ Unknown	6078872	Indicate whether RET gene rearrangement analysis was performed. Note: If RET rearrangement analysis was not performed, skip to Question 136.
135	Was a RET rearrangement identified?	☐ Yes ☐ No	6078882	Indicate whether a RET rearrangement was identified.



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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
136	Was PD-L1 IHC performed?	☐ Yes ☐ No ☐ Unknown	6078873	Indicate whether PD-L1 expression was assessed by immunohistochemistry (IHC). Note: If PD-L1 IHC was not performed, skip to
137	PD-L1 expression by IHC	☐ Positive ☐ Negative ☐ Equivocal	4798631	Question 138. Indicate the status of PD-L1 protein expression as determined by immunohistochemistry (IHC).
138	Was HER2 IHC performed?	☐ Yes ☐ No ☐ Unknown	6063454	Indicate whether HER2 expression was assessed by immunohistochemistry (IHC). Note: If HER2 IHC was not performed, skip to Question 140.
139	HER2 expression by IHC	□ Positive □ Negative □ Equivocal	2957563	Indicate the expression of HER2 as assessed by immunohistochemistry (IHC).
Additional	 Metastatic/Recurrent	Tumor Biospecimen Information (if applicable)	1	
140	Are you submitting an additional metastatic/ recurrent tumor tissue sample?	☐ Yes ☐ No		A biospecimen obtained from a single site at a single timepoint in progression that is portioned for both sequencing and model generation counts as 1 single tumor specimen. A biospecimen obtained from another site or at a later timepoint in progression that is portioned for both sequencing and model generation counts as a second single tumor specimen. Note: If yes, proceed to Question 141, otherwise, skip to Question 181.
141	Metastatic/ recurrent tissue biospecimen ordinal		6584266	Please provide a number to identify which biospecimen this is in the sequence. The first biospecimen should be number "1," the second should be number "2," etc.
142	CMDC tissue ID		6586035	Please provide the CMDC sample ID for this biospecimen as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
143	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
144	Metastatic/ recurrent tumor tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the metastatic/recurrent tumor tissue sample collected for molecular characterization.
145	Number of days from index date to date of diagnosis of additional metastasis/ recurrence		6132218	Provide the number of days from the index date to the date of diagnosis of additional metastatic/recurrent disease.
146	Method of metastatic/ recurrent cancer sample procurement	☐ Cytology ☐ Fine needle aspiration biopsy ☐ Incisional biopsy ☐ Excisional biopsy ☐ Tumor resection ☐ Other (specify)	6587389	Indicate the procedure performed to obtain the metastatic/recurrent tumor tissue. Note: If the method of procurement is not listed, proceed to Question 146a, otherwise, skip to Question 147.
146a	Other method of cancer sample procurement		6587390	If the procedure performed to obtain the tumor tissue is not included in the provided list, specify the procedure.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
147	Number of days		3288495	Provide the number of days from the index
	from index date to			date to the date of the procedure that
	date of metastatic/			produced the metastatic/recurrent tumor
	recurrent sample			tissue submitted for HCMI.
	procurement			
148	Metastatic/	☐ Left lower lobe lung	6587394	Select the site from which the
2.0	recurrent site	☐ Left upper lobe lung		metastatic/recurrent tissue used to develop
		☐ Right lower lobe lung		the model was derived. Note: If the
		☐ Right middle lobe lung		metastatic/recurrent site is not listed, proceed to
		☐ Right upper lobe lung		Question 148a, otherwise, skip to Question 149.
		☐ Adrenal gland		
		☐ Bone		
		☐ Brain		
		Liver		
		Lung		
		☐ Lymph node(s)- non-regional		
		☐ Other (specify)		
		D other (specify)		
148a	Other metastatic/		6587395	If not included in the previous list, specify
	recurrent site			the site from which the metastatic/
				recurrent tissue used to develop the model
				was derived.
149	Site of relapse	Local	2002506	If the primary tumor relapsed, provide the
		☐ Regional		site of relapse.
		Distant		
150	ICD-10 code	☐ Not applicable	3226287	Provide the ICD-10 code for the
	100 20 0000		0220207	metastatic/recurrent tumor used to
				generate the model submitted to HCMI.
151	ICD-O-3 histology		3226275	Provide the ICD-O-3 histology code
	code			describing the morphology of the
				metastatic/recurrent tumor used to
				generate the model submitted to HCMI.
152	Maintenance		6119066	Provide the name(s) of the maintenance
	and/or			and/or consolidation therapy administered
	consolidation			to the patient prior to the collection of the
	therapy			metastatic/recurrent tissue used to develop
	administered prior			the model.
	to collection of			
	metastatic/ recurrent tissue			
153	Days to start of		5102411	Provide the number of days from the index
133	maintenance		3102411	date to the date maintenance and/or
	and/or			consolidation therapy started.
	consolidation			consolidation therapy started.
	therapy from index			
	date			
154	Days to last known		5102431	Provide the number of days from the index
	administration date			date to the last known date of maintenance
	of maintenance			and/or consolidation therapy.
	and/or			· ·
	consolidation			
	therapy from index			
	date			
155	Is the patient still	☐ Yes	6379568	Indicate whether the patient is still
	receiving	□ No		undergoing maintenance and/or
	treatment?	☐ Unknown		consolidation therapy.



Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
156	Disease status	☐ No evidence of disease	2188290	Provide the disease status following
		☐ Progressive disease		maintenance and/or consolidation therapy.
		☐ Stable disease		
	/p !: .: /::: . ! =	Unknown		
157	Was KRAS mutation	eatures for Additional Metastatic/Recurrent Tun	6060079	Indicate whether KRAS mutation analysis
137	analysis	Yes	0000079	was performed. <i>Note: If KRAS mutation</i>
	performed?	□ No		analysis was not performed, skip to Question
	periorinea.	☐ Unknown		160.
158	Was a mutation in		6060081	Indicate whether a mutation in KRAS was
	KRAS identified?	□ Yes		identified through mutation analysis. Note: If
		□ No		KRAS mutation was not identified, skip to Question 160.
				question 100.
159	If KRAS mutation	□ G13V	6060083	Indicate the specific KRAS mutation
	identified, which	☐ G12A ☐ G12V ☐ Q61H ☐ G12C ☐ G13A ☐ Q61H		identified. Note: If the KRAS mutation is not
	one?	☐ G12C ☐ G13A ☐ Q61L ☐ G13C ☐ T 14.4.5		listed, proceed to Question 159a, otherwise, skip
		П G12В П G13D □ А1461		to Question 160.
		□ G12S □ G13R □ Other		
159a	Other KRAS	(specify)	6101691	If the KRAS mutation identified is not
139a	mutation(s)		0101031	provided in the previous list, specify the
	matation(3)			KRAS mutation.
160	Was EGFR mutation	П У	6063955	Indicate whether EGFR mutation analysis
	analysis	☐ Yes ☐ No		was performed. Note: If EGFR mutation
	performed?	□ Unknown		analysis was not performed, skip to Question
1.54		2 chillown	6060500	163.
161	Was a mutation in EGFR identified?	☐ Yes	6063530	Indicate whether a mutation in EGFR was
	EGFR identified?	□ No		identified through mutation analysis. Note: If EGFR mutation was not identified, skip to
				Question 163.
162	If EGFR mutation	☐ G719X ☐ L861Q	3147627	Indicate the specific EGFR mutation
	identified, which	☐ T790M ☐ Exon 19 deletion		identified. Note: If the EGFR mutation is not
	one?	☐ C797S ☐ Exon 20 insertion		listed, proceed to Question 162a, otherwise, skip to Question 163.
162a	Other EGFR	☐ L858R ☐ Other (specify)	4173882	If the EGFR mutation identified is not
1024	mutation(s)		41/3002	provided in the previous list, specify the
	matation(s)			EGFR mutation.
163	Was BRAF mutation	☐ Yes	6061813	Indicate whether BRAF mutation analysis
	analysis	□ No		was performed. Note: If BRAF mutation
	performed?	☐ Unknown		analysis was not performed, skip to Question
164	Was a mutation in		6061809	166. Indicate whether a mutation in BRAF was
104	BRAF identified?	☐ Yes	0001003	identified through mutation analysis. Note: If
	Divid identified.	□ No		BRAF mutation was not identified, skip to
				Question 166.
165	If BRAF mutation	□ V600D □ V600R	6061810	Indicate the specific BRAF mutation
	identified, which	□ V600E □ K601E		identified. Note: If the BRAF mutation is not
	one?	☐ V600K ☐ Other (specify)		listed, proceed to Question 165a, otherwise, skip to Question 166.
165a	Other BRAF		6101687	If the BRAF mutation identified is not
	mutation			provided in the previous list, specify the
				BRAF mutation.
166	Was MET mutation	☐ Yes	6078874	Indicate whether MET mutation analysis
	analysis	□ No		was performed. Note: If MET mutation analysis
	performed?	□ Unknown		was not performed, skip to Question 169.
167	Was a mutation in	□ Vos	6078886	Indicate whether a mutation in MET was
	MET identified?	│ □ Yes │ □ No		identified through mutation analysis. Note: If MET mutation was not identified, skip to
				Question 169.
		ı	ı	

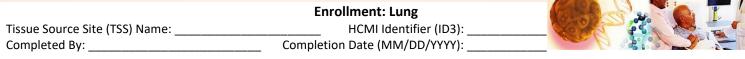
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
168	If MET mutation identified, which one?	☐ D963 splice mutation ☐ D1010N ☐ D1010 splice mutation ☐ Intron 13 mutation ☐ Intron 14 mutation ☐ Exon 14 mutation ☐ Exon 14 deletion	6078901	Indicate the specific MET mutation identified. Note: If the MET mutation is not listed, proceed to Question 168a, otherwise, skip to Question 169.
		☐ Other (specify)		
168a	Other MET mutation		6230525	If the BRAF mutation is not included in the list provided, specify the MET mutation identified.
169	Was EML4-ALK translocation analysis performed?	☐ Yes ☐ No ☐ Unknown	6063976	Indicate whether EML4-ALK translocation analysis was performed. Note: If EML4-ALK translocation analysis was not performed, skip to Question 173.
170	Was an EML4-ALK translocation identified?	☐ Yes ☐ No	6063959	Indicate whether an EML4-ALK translocation was identified. Note: If EML4-ALK translocation was not identified, skip to Question 173.
171	If EML4-ALK translocation found, which variant?	☐ Variant 1 ☐ Variant 2 ☐ Variant 3 ☐ Variant 4 ☐ Variant 5	3139445	Indicate the EML4-ALK translocation variant that was identified.
172	What method was used to identify the EML4-ALK translocation	☐ Cytogenetics ☐ FISH ☐ NGS ☐ IHC (nuclear staining) ☐ RT-PCR ☐ Other	3139449	Specify the method used to identify the EML4-ALK translocation variant.
173	Was ROS1 FISH performed?	☐ Yes ☐ No ☐ Unknown	6078871	Indicate whether fluorescence in situ hybridization (FISH) analysis of the ROS1 gene was performed. Note: If ROS FISH was not performed, skip to Question 175.
174	Was ROS1 rearrangement detected by FISH?	☐ Yes ☐ No	6078881	Indicate whether a ROS1 rearrangement was identified by fluorescence in situ hybridization (FISH).
175	Was RET gene rearrangement analysis performed?	☐ Yes ☐ No ☐ Unknown	6078872	Indicate whether RET gene rearrangement analysis was performed. Note: If RET rearrangement analysis was not performed, skip to Question 177.
176	Was a RET rearrangement identified?	☐ Yes ☐ No	6078882	Indicate whether a RET rearrangement was identified.
177	Was PD-L1 IHC performed?	☐ Yes ☐ No ☐ Unknown	6078873	Indicate whether PD-L1 expression was assessed by immunohistochemistry (IHC). Note: If PD-L1 IHC was not performed, skip to Question 179.
178	PD-L1 expression by IHC	☐ Positive ☐ Negative ☐ Equivocal	4798631	Indicate the status of PD-L1 protein expression as determined by immunohistochemistry (IHC).
179	Was HER2 IHC performed?	☐ Yes ☐ No ☐ Unknown	6063454	Indicate whether HER2 expression was assessed by immunohistochemistry (IHC). Note: If HER2 IHC was not performed, skip to Question 181.
180	HER2 expression by IHC	☐ Positive ☐ Negative ☐ Equivocal	2957563	Indicate the expression of HER2 as assessed by immunohistochemistry (IHC).



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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
Metastatio	/Recurrent Tumor Mod	del Information		
181	METASTATIC/ RECURRENT model biospecimen ordinal		6594587	Please provide a number to identify which biospecimen this is in the sequence. Note: The first biospecimen should be number "1," the second should be number "2," etc.
182	CMDC model ID		6586036	Please provide the CMDC model ID for this sample as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
183	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
184	Model's METASTATIC/ RECURRENT tumor tissue CMDC sample ID		6586035	Enter the CMDC Sample ID of the METASTATIC/RECURRENT tissue from which this model is derived.
185	Model's METASTATIC/ RECURRENT tumor tissue biospecimen ordinal		6584266	Enter the biospecimen ordinal of the METASTATIC/RECURRENT tissue from which this model is derived.
Additional	Metastatic/Recurrent	Biospecimen Tumor Model Information (if applie	cable)	
186	METASTATIC/ RECURRENT model biospecimen ordinal		6594587	Please provide a number to identify which biospecimen this is in the sequence. Note: The first biospecimen should be number "1," the second should be number "2," etc.
187	CMDC model ID		6586036	Please provide the CMDC model ID for this sample as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
188	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
189	Model's METASTATIC/ RECURRENT tumor tissue CMDC sample ID		6586035	Enter the CMDC Sample ID of the METASTATIC/RECURRENT tissue from which this model is derived.
190	Model's METASTATIC/ RECURRENT tumor tissue biospecimen ordinal		6584266	Enter the biospecimen ordinal of the METASTATIC/RECURRENT tissue from which this model is derived.
Other Bios	pecimen Information		1	
191	Are you submitting an OTHER tissue sample?	☐ Yes ☐ No		Indicate whether an OTHER tissue sample (e.g. pre-malignant, non-malignant, or dysplastic tissue, etc.) was collected for HCMI for this case. <i>Note: If yes, proceed to Question 192.</i>
192	OTHER tissue biospecimen ordinal		6584267	Please provide a number to identify which biospecimen this is in the sequence. Note: The first biospecimen should be number "1," the second should be number "2," etc.

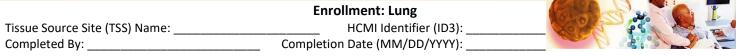


Question	Question Text	Data Entry Options	CDE ID	Instruction Text
193	CMDC sample ID		6586035	Please provide the CMDC sample ID for this specimen as it will appear on tubes and the Sample Submission Form transmitted to the BPC.
194	BPC submitter ID (if available)		6584919	Please provide the BPC-generated ID for this sample as it will appear on the Sample Submission Form transmitted to the BPC.
195	OTHER tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the OTHER tissue sample collected for molecular characterization.
196	Other method of cancer sample procurement	 □ Cytology □ Fine needle aspiration biopsy □ Incisional biopsy □ Excisional biopsy □ Tumor resection □ Other (specify) 	6587398	Provide the procedure performed to obtain the OTHER tissue. Note: If the method of procurement is not listed, proceed to Question 196a, otherwise, skip to Question 197.
196a	Specify method of OTHER tissue sample procurement		6587399	Specify the procedure performed to obtain the OTHER tissue.
197	Number of days from index date to date of OTHER sample procurement		3288495	Provide the number of days from the index date to the date of the procedure that produced the OTHER tissue submitted for HCMI.
198	Tissue type	☐ Pre-malignant ☐ Other (specify)	64784	Indicate the OTHER tissue type. Note: If the OTHER tissue type is not listed, proceed to Question 198a, otherwise, skip to Question 199.
198a	Specify tissue type		64785	Specify the OTHER tissue type if not in the provided list.
199	Anatomic site of OTHER tissue	□ Left lower lobe lung □ Left upper lobe lung □ Right lower lobe lung □ Right middle lobe lung □ Right upper lobe lung □ Adrenal gland □ Bone □ Brain □ Liver □ Lung □ Lymph node(s)- non-regional □ Other (specify)	6696813	Select the site from which the OTHER tissue used to develop the model was derived. Note: If the OTHER tissue site is not listed, proceed to Question 199a, otherwise, skip to Question 200.
199a	Specify anatomic site of OTHER tissue		6584916	Specify the site of OTHER tissue, if not in the previous list.
200	ICD-10 code		3226287	Provide the ICD-10 code for the OTHER tissue used to generate the model submitted to HCMI.
201	ICD-O-3 histology code		3226275	Provide the ICD-O-3 histology code describing the morphology of the OTHER tissue used to generate the model submitted to HCMI.
Additional	OTHER biospecimen In	formation (if applicable)		



		BUOM BUILDING
Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	16
Completed By:	Completion Date (MM/DD/YYYY):	160

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
202	Are you submitting			Indicate whether an additional OTHER tissue
	an additional			sample (pre-malignant, non-malignant, or
	OTHER tissue	☐ Yes		dysplastic tissue, etc.) is being submitted for
	sample?	□ No		HCMI for this case.
				Note: If yes, proceed to Question 203, otherwise, skip to Question 213.
203	OTHER tissue		6584267	Please provide a number to identify
	biospecimen			which biospecimen this is in the sequence.
	ordinal			Note: The first biospecimen should be number "1," the second should be number "2," etc.
204	CMDC sample ID		6586035	Please provide the CMDC sample ID
204	CIVIDE Sample 1D		0300033	for this specimen as it will appear on tubes
				and the Sample Submission Form
				transmitted to the BPC.
205	BPC submitter ID (if		6584919	Please provide the BPC-generated ID for this
	available)			sample as it will appear on the Sample
	,			Submission Form transmitted to the BPC.
206	OTHER tissue	☐ Cryopreserved	5432521	Provide the method used to preserve the
	sample	□ FFPE		OTHER tissue sample collected for molecular
	preservation	☐ Frozen		characterization.
	method	□ ост		
		☐ Snap frozen		
207	Other method of	·	6587398	Provide the procedure performed to obtain
	cancer sample	☐ Cytology☐ Fine needle aspiration biopsy		the OTHER tissue. Note: If the method of
	procurement	☐ Incisional biopsy		procurement is not listed, proceed to Question
		☐ Excisional biopsy		207a, otherwise, skip to Question 208.
		☐ Tumor resection		
		☐ Other (specify)		
207-	Constitution at least	_ c.i.e. (spee//	CE07200	Constitution and an automorphism
207a	Specify method of OTHER tissue		6587399	Specify the procedure performed to obtain the OTHER tissue.
	sample			the OTHER tissue.
	procurement			
208	Number of days		3288495	Provide the number of days from the index
200	from index date to		3200433	date to the date of the procedure that
	date of			produced the OTHER tissue submitted for
	OTHER sample			HCMI.
	procurement			
209	Tissue type		64784	Indicate the OTHER tissue type.
		☐ Pre-malignant		Note: If the OTHER tissue type is not listed,
		☐ Other (specify)		proceed to Question 209a, otherwise, skip to
		= other (speeny)		Question 210.
209a	Specify tissue type		64785	Specify the OTHER tissue type
2034	Specify tissue type		04765	if not in the provided list.
				c the provided hat
210	Anatomic site of	☐ Left lower lobe lung	6696813	Select the site from which the OTHER tissue
	OTHER tissue	☐ Left upper lobe lung		used to develop the model was derived.
		☐ Right lower lobe lung		Note: If the OTHER tissue site is not listed,
		☐ Right middle lobe lung		proceed to Question 210a, otherwise, skip to
		☐ Right upper lobe lung		Question 211.
		☐ Adrenal gland		
		☐ Bone		
		☐ Brain		
		☐ Liver		
		☐ Lung		
		☐ Lymph node(s)- non-regional		
		☐ Other (specify)		



Question	Question Text	Data Entry Options	CDE ID	Instruction Text
210a	Specify anatomic		6584916	Specify the site of OTHER tissue, if not in the
	site of OTHER tissue			previous list.
211	ICD-10 code		3226287	Provide the ICD-10 code for the OTHER
				tissue used to generate the model
				submitted to HCMI.
212	ICD-O-3 histology		3226275	Provide the ICD-O-3 histology code
	code			describing the morphology of the OTHER
				tissue used to generate the model
				submitted to HCMI.
Other Tissa	ue Model Information			
213	OTHER tissue		6594590	Please provide a number to identify
	model biospecimen			which biospecimen this is in the sequence.
	ordinal			Note: The first biospecimen should be number
24.4	CNADC d-LID		CE0C02C	"1," the second should be number "2," etc.
214	CMDC model ID		6586036	Please provide the CMDC model ID
				for this sample as it will appear on tubes and
				the Sample Submission Form transmitted to
215	BPC submitter ID (if		6584919	the BPC. Please provide the BPC-generated ID
213	available)		0364919	for this sample as it will appear on the
	available)			Sample Submission Form transmitted
				to the BPC.
216	Model's OTHER		6586035	Enter the CMDC Sample ID of the OTHER
210	tissue CMDC		0300033	tissue from which this model is derived.
	sample ID			dissue from which this model is derived.
217	Model's OTHER		6584267	Enter the biospecimen ordinal of the OTHER
	tissue biospecimen		000.207	tissue from which this model is derived.
	ordinal			
Additional	Other Tissue Model In	formation (if applicable)	l	
218	OTHER tissue	., ,,	6594590	Please provide a number to identify which
	model biospecimen			biospecimen this is in the sequence. <i>Note:</i>
	ordinal			The first biospecimen should be number "1," the
				second should be number "2," etc.
219	CMDC model ID		6586036	Please provide the CMDC model ID
				for this sample as it will appear on tubes and
				the Sample Submission Form transmitted to
25.5				the BPC.
220	BPC submitter ID (if		6584919	Please provide the BPC-generated ID
	available)			for this sample as it will appear on the
				Sample Submission Form transmitted
224	Modelle OTUED		6500005	to the BPC.
221	Model's OTHER tissue CMDC		6586035	Enter the CMDC Sample ID of the OTHER tissue from which this model is derived.
				ussue from which this model is derived.
222	sample ID Model's OTHER		6584267	Enter the biospecimen ordinal of the OTHER
222	tissue biospecimen		0304207	tissue from which this model is derived.
	ordinal			ussue from which this model is derived.
	Orumai			