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Enrollment: Endometrium

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Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):
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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. This form should be used for Endometrial Cancers and Uterine Sarcomas.

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	ormation		•	
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	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.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
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.
11	Patient age on index date		6379572	Provide the age (in days) of the patient on 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 cancer	☐ Same ☐ Different ☐ None ☐ Unknown	5832923	Has a first-degree relative of the patient been diagnosed with a cancer of the same or a different type?
14	Smoking history	 □ Lifelong non-smoker (<100 cigarettes smoked in a lifetime) □ Current smoker (includes daily and non-daily smokers) □ Current reformed smoker (duration not specified) □ Current reformed smoker for >15 years □ Current reformed smoker for ≤15 years 	2181650	Indicate the patient's history of tobacco smoking as well as their current smoking status using the defined categories.
15	Metastasis at diagnosis assessment status	☐ Metastatic☐ Non-metastatic (confirmed)☐ Non-metastatic (unconfirmed)	3438571	Indicate whether there was evidence of metastasis at the time of diagnosis of the primary tumor. Note: If metastatic at diagnosis, proceed to Question 16, otherwise, skip to Question 17.
16	Metastatic site(s) at diagnosis	□ Extra-abdominal lymph node(s) □ Soft tissue □ Pelvic lymph node(s) □ Right ovary □ Para-aortic lymph node(s) □ Right fallopian tube □ Peritoneum □ Left fallopian tube □ Lung □ Right tubo-ovarian □ Liver □ Left tubo-ovarian □ Adrenal gland □ Omentum □ Brain □ Other (specify) □ Bone	3029815	Indicate the site(s) of metastasis at the time of diagnosis of the primary tumor. Note: If the anatomic site of tumor tissue is not listed, proceed to Question 16a, otherwise, skip to Question 17.
16a	Specify metastatic site(s)		3128033	If the site of metastasis is not included on the provided list, specify the site of metastasis.
17	Clinical history (select all that apply)	□ Lynch syndrome □ Breast cancer □ Other cancer (specify) □ Polycystic ovarian syndrome (PCOS) □ Endometriosis □ Metabolic syndrome □ Diabetes □ Other (specify)	6690684	Select all relevant prior diseases/disorders in the patient's clinical history. Note: If 'Other cancer' is selected, proceed to Question 17a, otherwise, skip to Question 18. If the clinical history is not listed, proceed to Question 17b, otherwise, skip to Question 18.
17a	Other cancer clinical history		7028280	Specify the other cancer type in the patient's clinical history.
17b	Other clinical history		6690685	If not included in the previous list, specify other relevant prior diseases/disorders in the patient's clinical history.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
18	Has the participant ever received hormone replacement therapy?	☐ Yes ☐ No ☐ Unknown	5206889	Indicate whether the patient has ever used hormone replacement therapy. Note: If the patient has received hormone replacement therapy, proceed to Question 19, otherwise, skip to Question 20.
19	What type of hormone therapy was administered?	☐ Estrogen only☐ Progesterone only☐ Progesterone and Estrogen☐ Unknown	3690502	Indicate the type of hormones the patient mainly used during that time.
20	Has the participant ever used oral contraceptives?	☐ Yes ☐ No ☐ Unknown	7028283	Indicate whether the patient has ever used oral contraceptives. Note: If the patient has used oral contraceptives, proceed to Question 21, otherwise, skip to Question 22.
21	What type of oral contraceptives were used?	☐ Progestin only ☐ Progestin and estrogen ☐ Unknown	7028284	Indicate the type of oral contraceptive agent used.
Biospecim	en Information			
22	Tissue sample type(s) collected for HCMI for this case	 □ Normal tissue □ Primary tumor □ Metastatic □ Recurrent □ Other tissue 	2006911	Please select all the tissue sample types submitted for HCMI with this case.
23	Number of NORMAL tissues biospecimens collected for HCMI for this case		6584256	Please provide the number of normal tissue specimens obtained for HCMI for this case. <i>Note: This number is expected to be 1.</i>
24	Number of PRIMARY cancer tissue biospecimens collected for HCMI model development for this case		6584257	Please provide the number of primary tumor specimens obtained for HCMI for this case. Note: A single primary tumor biospecimen obtained that is portioned for both sequencing and model generation counts as 1 single primary tumor specimen. This number is expected to be 1.
25	Number of METASTATIC/RECURRENT cancer tissue biospecimens collected for HCMI model development for this case		6584258	Please provide the number of metastatic and/or recurrent cancer biospecimens collected for HCMI for this case. Note: 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.
26	Number of OTHER tissue biospecimens collected for HCMI model development for this case		6584259	Please provide the number of pre- malignant, non-malignant, or dysplastic tissue biospecimens collected for HCMI for this case. Note: 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.

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Tissue Source Site (TSS) Name: _	HCMI Identifier (ID3):
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
27	Total number of tissue biospecimens collected for HCMI for this case		6584271	Please provide the total number of tissue biospecimens collected for HCMI for this case. Note: This number should be the sum of the normal, primary tumor, metastatic/recurrent tumor, and other biospecimen counts above.
	ntrol Information		_	
28	Normal tissue biospecimen ordinal		6584264	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.
29	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.
30	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.
31	Type of normal control	 □ Whole blood □ Buccal cells □ 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 	3081936	Indicate the type of normal control submitted for this case.
32	Anatomic site of normal tissue	☐ Right ovary ☐ Left ovary ☐ Right fallopian tube ☐ Left fallopian tube ☐ Right tubo- ovarian ☐ Left tubo-ovarian ☐ Lower uterine segment ☐ Skin ☐ Other (specify) ☐ Not applicable	4132152	If non-neoplastic tissue was submitted as the normal control, select the anatomic site of the normal tissue. Note: If the anatomic site of normal tissue is not listed, proceed to Question 32a, otherwise, skip to Question 33.
32a	Other anatomic site of normal tissue		3288189	If non-neoplastic tissue, adjacent tissue, or normal tissue from another anatomic site was submitted as the normal control, provide the anatomic site of the normal tissue.
33	Distance from tumor to normal control tissue (if not blood)	 □ Adjacent (< or = 2cm) □ Distal (>2cm) □ Unknown □ Not applicable 	3088708	Indicate the distance from the site of normal tumor collection to the primary tumor. Note: If normal tissue was not submitted, select 'Not applicable'.
34	Normal tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the normal tissue sample collected for molecular characterization.
Primary Tu	ımor Biospecimen Informatio			
35	ICD-10 code for primary tumor	☐ C54.0 ☐ C54.1 ☐ C54.2 ☐ C54.3 ☐ C54.8 ☐ C54.9 ☐ Other (specify)	3226287	Provide the ICD-10 code for the primary tumor as used to generate the ID3 for this subject. Note: If the ICD-10 code is not listed, proceed to 35a, otherwise, skip to Question 36.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
35a	Other ICD-10 code for primary tumor		3226287	If the ICD-10 code for the tumor used to generate the model submitted to HCMI is not included on the provided list, specify the ICD-10 code.
36	Tumor morphology	□ 8263/3 □ 8933/3 □ 8441/3 □ 8380/3 □ 8890/3 □ 8480/3 □ 8382/3 □ 8891/3 □ 8013/3 □ 8383/3 □ 8896/3 □ 8013/3 □ 8570/3 □ 8900/3 □ 8041/3 □ 8930/3 □ 8714/3 □ 8020/3 □ 8931/3 □ 8805/3 □ Other (specify)	3226275	Using the patient's pathology/laboratory report, provide the ICD-O-3 histology code of the primary tumor. Note: If the ICD-O-3 histology code of the primary tumor is not listed, proceed to Question 36a, otherwise, skip to Question 37.
36a	Specify other morphology		3226275	If the ICD-O-3 histology code describing the morphology of the patient's primary tumor is not included on the previous list, provide the ICD-O-3 histology code.
37	Tissue or organ of origin	☐ Endometrium☐ Uterus☐ Other (specify)	3427536	Using the patient's pathology/laboratory report, select the primary site of the disease. Note: If the primary site of the disease is not listed, proceed to Question 37a, otherwise skip to Question 38.
37a	Other tissue or organ of origin		5946219	If the primary site of the disease is not included on the previous list, provide the primary site of the disease.
38	Histological type	☐ Endometrial cancer ☐ Uterine cancer ☐ Other (specify)	3081932	Select the surgical pathology text description of the histological tumor type. Note: If the histological tumor type is not listed, proceed to Question 38a, otherwise, skip to Question 39.
38a	Other histological type		3294805	If the traditional surgical pathology text description of the histological tumor type is not included on the previous list, please specify the histological type.
39	Histological subtype: endometrial carcinomas	 □ Endometrioid carcinoma, NOS □ Endometrioid carcinoma with squamous differentiation □ Endometrioid carcinoma, villoglandular variant □ Endometrioid carcinoma with secretory differentiation □ Endometrioid carcinoma, other variant (specify) □ Serous endometrial intraepithelial carcinoma □ Carcinosarcoma (malignant mixed Müllerian tumor) □ Mucinous carcinoma □ Clear cell carcinoma □ Small cell neuroendocrine carcinoma □ Large cell neuroendocrine carcinoma □ Mixed cell carcinoma (specify types and percentages) □ Undifferentiated carcinoma 	3081934	Using the patient's pathology/laboratory report, select the histological subtype of the primary tumor. Note: If "Endometrioid carcinoma, other variant" is selected, proceed to Question 39a. If "Mixed cell carcinoma" is selected, proceed to Question 39b-c. If the histological subtype is not listed, proceed to Question 39d, otherwise, skip to Question 40.

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		☐ De-differentiated carcinoma ☐ Other (specify)		
39a	Specify the other endometrioid carcinoma variant		7029865	If not provided in the previous list, specify the endometrioid adenocarcinoma variant histologic subtypes present.
39b	Specify mixed cell carcinoma types		7029863	Specify the endometrial mixed cell adenocarcinoma histologic subtypes present.
39c	Specify mixed cell carcinoma percentages		7029864	Specify the percentages present of the endometrial mixed cell adenocarcinoma histologic subtypes.
39d	Other histological subtype		3124492	If the histological subtype for the primary tumor is not included in the provided list, specify the histological subtype.
40	Histological subtype: uterine sarcomas	 □ Leiomyosarcoma □ Leiomyosarcoma, epithelioid type □ Leiomyosarcoma, myxoid type □ Endometrial stromal sarcoma □ Endometrial stromal sarcoma with smooth muscle differentiation □ Endometrial stromal sarcoma with sex cord elements □ Endometrial stromal sarcoma with glandular elements □ Undifferentiated uterine/endometrial sarcoma □ Adenosarcoma □ Adenosarcoma with rhabdomyoblastic differentiation □ Adenosarcoma with cartilaginous differentiation □ Adenosarcoma with other heterologous element (specify) □ Adenosarcoma with sarcomatous overgrowth □ Endometrial rhabdomyosarcoma □ Malignant perivascular epithelioid cell tumor □ Other (specify) 	3081934	Provide the traditional surgical pathology text description of the histological tumor subtype. Note: If "Adenosarcoma with other heterologous element" is selected, proceed to Question 40a. If the histological subtype is not listed, proceed to Question 40b, otherwise, skip to Question 41.
40a	Specify the other heterologous element		7029866	Specify the other heterologous elements present in the endometrioid adenocarcinoma histologic subtype.
40b	Other histological subtype: uterine sarcomas		3124492	If not provided in the previous list, specify the traditional surgical pathology text description of the histological tumor subtype.
41	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.
42	Prior malignancy (other cancer type)	☐ Yes ☐ No ☐ Unknown	5878828	Indicate whether the patient has a history of prior malignancy of a different cancer type.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
43	AJCC cancer staging edition	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	2722309	Select the AJCC staging handbook edition used to stage the patient's primary tumor.
44	AJCC pathologic spread: Primary tumor (pT)	□ TX □ T1b □ T2a □ T3a □ T0 □ T1c □ T2b □ T3b □ T1 □ T2 □ T3 □ T4	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).
45	AJCC pathologic spread: Lymph nodes (pN)	□ NX □ N1 □ N2 □ N0 □ N1mi □ N2mi □ N0(i+) □ N1a □ N2a	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).
46	AJCC pathologic spread: Distant metastases (pM)	□ MX □ M1	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).
47	Tumor stage (pathological)	□ Stage I □ Stage IIIB □ Stage IA □ Stage IIIC □ Stage IB □ Stage IIIC1 □ Stage II □ Stage IIIC2 □ Stage III □ Stage IVA □ Stage IIIA □ Stage IVB	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).
48	FIGO Stage	□ Stage I □ Stage IIIB □ Stage IA □ Stage IIIC □ Stage IB □ Stage IIIC1 □ Stage IIA □ Stage IIIC2 □ Stage IIB □ Stage IVA □ Stage III □ Stage IVB	5326898	Using the patient's pathology /laboratory report, in conjunction with the patient's medical record, select the stage as defined by the Federation of Gynecology and Obstetrics (FIGO), 2014 edition.
49	FIGO grade (applies to endometrioid and mucinous carcinomas only)	☐ Grade 1 ☐ Other (specify) ☐ Grade 2 ☐ Cannot be assessed ☐ Grade 3 ☐ Not applicable	7028262	Select the grade representing the extent of endometrial cancer in the body, based on staging criteria from the Federation of Gynecology and Obstetrics (FIGO). Note: If the FIGO grade is not listed, proceed to Question 49a, otherwise, skip to Question 50.
49a	Specify other FIGO grade		7028262	If not included in the previous list, provide the grade representing the extent of endometrial cancer in the body, based on staging criteria from the Federation of Gynecology and Obstetrics (FIGO).
50	Histologic grade (applies to sarcomas only)	☐ Low grade ☐ Cannot be ☐ assessed ☐ With sarcomatous overgrowth ☐ Not applicable	7028298	Select the grade representing the extent of endometrioid stroma sarcoma in the body.
Prognostic	/Predictive/Lifestyle Feature	s for Primary Tumor Prognosis or Responsiveness t	o Treatmen	
51	Specimen integrity	☐ Intact ☐ Opened ☐ Mocellated ☐ Other (specify)	2783662	Indicate the physical status of the specimen. Note: If the specimen integrity is not listed, proceed to Question 51a, otherwise, skip to Question 52.
51a	Specify specimen integrity		7028264	If not included in the previous list, specify the physical condition of the specimen.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
52	Is there myometrial invasion?	☐ Yes ☐ No ☐ Unknown	3440922	Indicate whether tumor cells have invaded the myometrium. Note: If myometrial invasion is present, proceed to Question 53, otherwise, skip to Question 56.
53	Myometrial invasion depth (mm)	mm	7028265	Indicate the depth (in mm) of invasion of the myometrium by cancer.
54	Myometrial thickness (mm)	mm	3440923	Provide the greatest depth of the myometrial invasion as a percentage of the thickness of the involved wall.
55	Percentage of myometrial invasion	<u>*************************************</u>	3440923	Provide the greatest depth of the myometrial invasion as a percentage of the thickness of the involved wall.
56	Tumor size, largest dimension	cm	64215	Provide the largest dimension of the primary tumor in cm, regardless of anatomical plane.
57	Adenomyosis	□ Not identified □ Present, uninvolved by carcinoma □ Present, involved by carcinoma □ Cannot be determined	7028267	Indicate whether or not the growth of endometrial tissue inside the muscular wall of the uterine corpus is involved with endometrial cancer.
58	Uterine serosa involvement	☐ Yes ☐ No ☐ Cannot be determined	7024271	Indicate whether or not the uterine serosa (perimetrium) is involved in the disease.
59	Lower uterine segment involvement	 □ Not identified □ Present, superficial (non-myoinvasive) □ Present, myoinvasive □ Cannot be determined 	7028268	Indicate whether the isthmus uteri has been involved with endometrial cancer.
60	Cervical stromal involvement	☐ Present ☐ Not identified ☐ Cannot be determined	7028269	Indicate whether or not the cervix uteri stromal cells are involved with endometrial cancer.
61	Other tissue/organ involvement	 Not applicable Not identified Right ovary Left ovary Ovary (side not specified) Right fallopian tube Left fallopian tube Fallopian tube (side not specified) Vagina Right parametrium Left parametrium Pelvic wall Bladder wall Bladder mucosa Rectal wall Bowel mucosa Omentum Pelvic lymph node(s) Other organ/tissue (specify) Cannot be determined 	2793699	Select all sites of other tissue/organ involvement. This does not include the primary tumor site. Note: If the other tissue/organ involved is not listed, proceed to Question 61a, otherwise, skip to Question 62.
61a	Specify other tissue/organ involvement		7029867	If not included in the previous list, specify the other site of other tissue/organ involvement. This does not include the primary tumor site.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
62	Peritoneal/Ascitic fluid	 □ Not submitted/unknown □ Negative for malignancy (normal/benign) □ Atypical and/or suspicious □ Malignant (positive for malignancy) □ Unsatisfactory/nondiagnostic 	6690681	Indicate the outcome of cytological analysis for peritoneal/ascitic fluid.
63	Lymphovascular invasion	□ Present□ Absent□ Cannot be determined	64727	Indicate whether large vessel (vascular) invasion or small, thinwalled (lymphatic) invasion was detected in the tumor specimen.
64	Lymph nodes	 □ No lymph nodes submitted or found □ All lymph nodes negative for tumor cells □ Positive lymph nodes 	7028287	Indicate whether the lymph nodes are involved with endometrial cancer as determined by pathologic assessment. Note: If lymph nodes are positive, proceed to Question 65, otherwise, skip to Question 75.
65	Number of pelvic nodes with micrometastases		7028289	Indicate the number of pelvic lymph nodes that have micrometastases present.
66	Number of pelvic nodes with macrometastases		7028288	Indicate the number of pelvic lymph nodes that have macrometastases present.
67	Number of pelvic nodes with isolated tumor cells		7028290	Indicate the number of pelvic lymph nodes that have isolated tumor cells present.
68	Total number of pelvic nodes examined		7028294	Indicate the total number of pelvic lymph nodes assessed for the presence of metastases and/or tumor cells.
69	Number of pelvic sentinel nodes examined		7028295	Indicate the total number of pelvic sentinel lymph nodes assessed for the presence of metastases and/or tumor cells.
70	Number of para-aortic nodes with micrometastases		7028292	Indicate the number of para-aortic lymph nodes that have micrometastases present.
71	Number of para-aortic nodes with macrometastases		7028291	Indicate the number of para-aortic lymph nodes that have macrometastases present.
72	Number of para-aortic nodes with isolated tumor cells		7028293	Indicate the total number of para- aortic lymph nodes that have isolated tumor cells present.
73	Total number of para- aortic nodes examined		7028296	Indicate the total number of para- aortic lymph nodes assessed for the presence of metastases and/or tumor cells.
74	Number of para-aortic sentinel nodes examined		7028297	Indicate the total number of para- aortic sentinel lymph nodes assessed for the presence of metastases and/or tumor cells.
75	Additional pathologic findings	 □ None identified □ Atypical hyperplasia/endometrial intraepithelial neoplasia (EIN) □ Other (specify) 	7028278	Indicate additional pathologic findings present in the patient. Note: If the additional pathologic finding is not listed, proceed to Question 75a, otherwise, skip to Question 76.
75a	Specify additional pathologic findings		7028279	If not included in the previous list, specify any additional pathologic findings.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
76	Residual tumor	□ RO □ RX	2608702	Indicate the status of a tissue margin
		□ R1 □ RX □ R2 □ Not evaluated		following surgical resection.
77	Was ER IHC performed?	☐ Yes ☐ No ☐ Unknown	6062416	Indicate whether estrogen receptor (ER) expression was assessed by immunohistochemistry (IHC). Note: If ER IHC was performed, proceed to Question 78, otherwise, skip to Question 79.
78	ER expression by IHC	□ Positive□ Negative□ Equivocal	6690720	Indicate the status of estrogen receptor (ER) protein expression as determined by immunohistochemistry (IHC).
79	Was PR IHC performed?	☐ Yes ☐ No ☐ Unknown	6062504	Indicate whether progesterone receptor (PR) expression was assessed by immunohistochemistry (IHC). Note: If PR IHC was performed, proceed to Question 80, otherwise, skip to Question 81.
80	PR expression by IHC	☐ Positive ☐ Negative ☐ Equivocal	6063673	Indicate the status of progesterone receptor (PR) protein expression as determined by immunohistochemistry (IHC).
81	Was HER2 IHC performed?	☐ Yes ☐ No ☐ Unknown	6063454	Indicate whether HER2 expression was assessed by immunohistochemistry (IHC). Note: If HER2 IHC was performed, proceed to Question 82, otherwise, skip to Question 83.
82	HER2 expression by IHC	☐ Positive ☐ Negative ☐ Equivocal	2957563	Indicate the expression of HER2 as assessed by immunohistochemistry (IHC).
83	Was HER2 FISH/CISH performed?	☐ Yes ☐ No ☐ Unknown	6063447	Indicate whether HER2 was assessed by fluorescence in situ hybridization (FISH) or chromogenic in situ hybridization (CISH). Note: If HER2 FISH/CISH was performed, proceed to Question 84, otherwise, skip to Question 85.
84	HER2 status by FISH/CISH	☐ Amplified☐ Not amplified☐ Equivocal	2854089	Select the HER2 status as assessed by FISH/CISH.
85	Was MLH1 IHC performed?	☐ Yes ☐ No ☐ Unknown	6062411	Indicate whether MLH1 expression was assessed by immunohisto-chemistry (IHC). Note: If MLH1 IHC was performed, proceed to Question 86, otherwise, skip to Question 87.
86	MLH1 expression by IHC	□ Positive□ Negative□ Equivocal	6063668	Indicate the status of MLH1 protein expression as determined by immunohistochemistry (IHC).
87	MLH1 promoter methylation status	 □ MLH1 promoter hypermethylation present □ MLH1 promoter hypermethylation absent □ MLH1 promoter hypermethylation not assessed □ Cannot be determined 	6033150	Indicate the methylation status of the MLH1 promoter.
88	Was MSH2 IHC performed?	☐ Yes ☐ No ☐ Unknown	6062412	Indicate whether MSH2 expression was assessed by immunohistochemistry (IHC). Note: If MSH2 IHC was performed, proceed to Question 89, otherwise, skip to Question 90.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
89	MSH2 expression by IHC	□ Positive	6063669	Indicate the status of MSH2 protein
	,	☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
90	Was PMS2 IHC	·	6062413	Indicate whether PMS2 expression
	performed?	☐ Yes		was assessed by immunohisto-
	·	□ No		chemistry (IHC). Note: If PMS2 IHC was
		☐ Unknown		performed, proceed to Question 91,
				otherwise, skip to Question 92.
91	PMS2 expression by IHC	Positive	6063670	Indicate the status of PMS2 protein
		☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
92	Was MSH6 IHC	D V	6062414	Indicate whether MSH6 expression
	performed?	│ □ Yes │ □ No		was assessed by immunohisto-
				chemistry (IHC). Note: If MSH6 IHC was
		☐ Unknown		performed, proceed to Question 93, otherwise, skip to Question 94.
93	MSH6 expression by IHC	☐ Positive	6063671	Indicate the status of MSH6 protein
	Wishlo expression by file	☐ Negative	0003071	expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
94	Microsatellite instability	☐ Stable	6032776	Indicate the status of the tumor for
	, , , , , , , , , , , , , , , , , , , ,	Low	0002770	microsatellite instability. (High at
		☐ High		≥30% loci tested, low at 1%-29% loci
		☐ Equivocal		tested.)
95	Was p53 IHC performed?		6690705	Indicate whether p53 expression
	·	☐ Yes		was assessed by immunohisto-
		□ No		chemistry (IHC). Note: If p53 IHC was
		☐ Unknown		performed, proceed to Question 96,
				otherwise, skip to Question 97.
96	p53 expression by IHC	□ Normal	6690719	Indicate the status of p53 protein
		☐ Abnormal strong diffuse overexpression		expression as determined by
		(>90%)		immunohistochemistry (IHC).
		☐ Abnormal null expression (complete loss of		
		expression)		
07	W 4 C III C	☐ Cannot be determined	6600743	Indiants whether all comments
97	Was p16 IHC performed?	☐ Yes	6690713	Indicate whether p16 expression
		□ No		was assessed by immunohisto-
		☐ Unknown		chemistry (IHC). Note: If p16 IHC was performed, proceed to Question 98,
				otherwise, skip to Question 99.
98	p16 expression by IHC	☐ Positive	6690724	Indicate the status of p16 protein
		☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
99	Mismatch repair status	D Evidence of MAMP mutation by secures:	6002208	Indicate the mismatch repair gene
		☐ Evidence of MMR mutation by sequencing☐ Evidence of MMR protein loss by IHC		status.
		MMR loss evidence hypermutation		
		phenotype (>10 mutations/Mb)		
		□ No evidence of MMR alteration		
100	Was PTEN IHC	☐ Yes	6062415	Indicate whether PTEN expression
	performed?	□ No		was assessed by
		Unknown		immunohistochemistry (IHC).
101	PTEN expression by IHC	□ Positive	6063672	Indicate the status of PTEN protein
		□ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
102	Was PTEN mutation	│	6063446	Indicate whether PTEN mutation
	analysis performed?	□ No		analysis was performed. Note: If PTEN
		□ Unknown		mutation analysis was performed, proceed to Question 103, otherwise, skip
		L CHRIOWII		to Question 105.
L	I	<u> </u>		to Question 103.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
103	Was a mutation in PTEN	7, 2, 2, 2	6063529	Indicate whether a mutation in PTEN
	identified?	☐ Yes		was identified through mutation
		□ No		analysis. Note: if a PTEN mutation was
				identified, proceed to Question 104,
101				otherwise, skip to Question 105.
104	If PTEN mutation	□ R130G	6063736	Indicate the mutation identified for
	identified, which one?	│ □ R130Q │ □ R130L		the PTEN gene. Note: If the PTEN mutation is not listed, proceed to
		□ R130P		Question 104a, otherwise, skip to
		☐ Other (specify)		Question 105.
104a	Other PTEN mutation	(-) //	6816076	If the specific mutation of the PTEN
				gene identified was not included in
				the previous list, please specify the
				mutation
105	Was PIK3CA mutation	D V	6063445	Indicate whether PIK3CA mutation
	analysis performed?	☐ Yes ☐ No		analysis was performed. Note: If
		□ Unknown		PIK3CA mutation analysis was performed, proceed to Question 106,
		- CHRISWII		otherwise, skip to Question 108.
106	Was a mutation in		6063524	Indicate whether a mutation in
	PIK3CA identified?			PIK3CA was identified through
		Yes		mutation analysis. Note: If a PIK3CA
		□ No		mutation was identified, proceed to
				Question 107, otherwise, skip to Question 108.
107	If PIK3CA mutation	□ H1047R	7028281	Indicate the mutation identified for
	identified, which one?	□ R88Q □ E545K □ H1047L		the PIK3CA gene. Note: If the PIK3CA
		□ E542K □ E545D □ H1047Y □ E542A □ E545A □ H1047Y		mutation is not listed, proceed to
		☐ E542Q ☐ E545G ☐ H1047Q		Question 107a, otherwise, skip to
		□ F542V □ F5450 □ Other		Question 108.
		(specify)		
107a	Other PIK3CA mutation		7028282	If the specific mutation of the
				PIK3CA gene identified was not included in the previous list, please
				specify the mutation.
108	Was TP53 mutation		6063448	Indicate whether TP53 mutation
	analysis performed?	☐ Yes		analysis was performed. <i>Note: If TP53</i>
		□ No		mutation analysis was performed,
		☐ Unknown		proceed to Question 109, otherwise, skip
109	Was a mutation in TP53		6063523	to Question 111. Indicate whether a mutation in TP53
103	identified?		0003323	was identified through mutation
		Yes		analysis. Note: If a TP53 mutation was
		□ No		identified, proceed to Question 110,
				otherwise, skip to Question 111.
110	If TP53 mutation	□ R273H □ R248W	7028281	Indicate the mutation identified for
	identified, which one?	□ R273C □ R175H		the TP53 gene. Note: If the TP53 mutation is not listed, proceed to
		□ R2731 □ R175C		Question 110a, otherwise, skip to
		☐ R248Q ☐ Other (specify)		Question 111.
110a	Other TP53 mutation		6101683	If the specific mutation of the TP53
				gene identified was not included in
				the previous list, please specify the
				mutation.
	ımor Sample Information		1	<i>II</i>
111	Are you submitting a			If yes, proceed to question 112. If submitting a metastatic/recurrent
	primary tumor tissue sample for this case?	☐ Yes		tumor biospecimen, proceed to Question
	sample for tills case!	□ No		140.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
112	Primary tumor		6584265	Please provide a number to identify
	biospecimen ordinal			which biospecimen this is in the
				sequence. Note: This number should be "1".
113	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.
114	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 to the BPC.
115	Sample represents		6584730	Does this primary tumor specimen
	primary diagnosis?	□ Yes		represent the PRIMARY DIAGNOSIS
		□ No		for this Case ID3? Note: If no, proceed
				to Question 116, otherwise, skip to Question 117.
116	Specify the ICD-10 code		3226287	Provide the ICD-10 code for the
				primary tumor used to generate the
				model submitted to HCMI.
117	Tumor tissue sample	☐ Cryopreserved	5432521	Provide the method used to
	preservation method	□ FFPE		preserve the tumor tissue sample
		☐ Frozen ☐ OCT		collected for molecular characterization.
		☐ Snap frozen		Characterization.
118	Anatomic site from which	☐ Endometrium	4214629	Select the anatomic site of the
	the tumor was obtained	☐ Lower uterine segment		tumor tissue sample used to
		☐ Endometrial polyp		generate the model for HCMI. <i>Note:</i>
		☐ Adenomyosis		If the tissue or organ not listed, proceed
		☐ Other (specify)		to Question 118a. Otherwise, skip to Question 119.
118a	Other anatomic site from		5946219	If not provided in the previous list,
	which the tumor was			provide the anatomic site of the
	obtained			tumor tissue sample used to
110			2122511	generate the model for HCMI.
119	Method of cancer sample	☐ Total hysterectomy and bilateral salpingo-	3103514	Provide the procedure performed to
	procurement	oophorectomy Radical hysterectomy		obtain the primary tumor tissue. Note: If the method of procurement is
		☐ Simple hysterectomy		not listed, proceed to Question 119a,
		☐ Supracervical hysterectomy		otherwise, skip to Question 120.
		☐ Endometrial biopsy		
		☐ Hysteroscopy		
		☐ Dilation and curettage (D&C)		
1100	Chacifu the other method	☐ Other (specify)	2006720	Specification proceedings appropriate
119a	Specify the other method of tumor sample		2006730	Specify the procedure performed to obtain the primary tumor tissue, if
	procurement			not included in the previous list.
120	Number of days from		3288495	Provide the number of days from the
	index date to date of			index date to the date of the
	tumor sample			procedure that produced the tumor
	procurement			tissue submitted for HCMI.
121	Tumor tissue type	☐ Primary	3288124	Provide the primary tumor tissue
		☐ Additional Primary ☐ NOS		type for this sample.
Primary Tu	ımor Model Information			
122	Primary model		6594596	Please provide a number to identify
	biospecimen ordinal			which biospecimen this is in the
				sequence. Note: This number is
				expected to be "1".

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
123	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.
124	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.
125	Model represents primary diagnosis?	☐ Yes ☐ No	6584730	Does this model represent the primary diagnosis for this Case ID3?
126	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.
127	Model's primary tumor biospecimen ordinal		6584265	Enter the biospecimen ordinal of the PRIMARY TUMOR TISSUE from which this model is derived.
Treatment	Information			
128	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 129-137. Radiation therapy is addressed in Questions 138-139.
129	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 130-131. Hormone therapy is addressed in Questions 132-133. Immunotherapy is addressed in Questions 134-135. Targeted therapy is addressed in Questions 136-137.
130	Neoadjuvant chemotherapeutic regimen	□ Doxorubicin □ Ifosfamide □ Liposomal doxorubicin □ Etoposide □ Cisplatin □ Gemcitabine □ Paclitaxel □ Docetaxel □ Carboplatin □ Other (specify)	2853313	Select all chemotherapeutics used for neoadjuvant therapy. Note: If neoadjuvant chemotherapy was not given, skip to Question 132. If the neoadjuvant chemotherapeutic regimen is not listed, proceed to Question 130a, otherwise, skip to Question 131.
130a	Other neoadjuvant chemotherapeutic regimen		62694	If the neoadjuvant therapy is not included in the provided list, specify neoadjuvant therapies administered.
131	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.
132	Hormone therapy	□ Medroxyprogesterone acetate □ Anastrozole □ Megestrol acetate □ Exemestane □ Tamoxifen □ Progestational □ Goserelin IUD □ Leuprolide □ Other (specify)	7028259	Select the hormone therapy administered to the patient. Note: If hormone therapy was not administered, skip to Question 134. If the hormone therapy is not listed, proceed to Question 132a, otherwise, skip to Question 133.
132a	Other hormone therapy		2405358	If the hormone therapy is not included in the provided list, specify hormone therapy.
133	Days to hormone therapy treatment from index date		5102411	Provide the number of days from index date to the date of treatment with hormone therapy.

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Question	Question Text	Data Entry Options		CDE ID	Instruction Text
134	Immunotherapy	, , , , , , , , ,		7028261	Select the immunotherapy
		☐ Filgrastim ☐ Other (specify)			administered to the patient. Note: If immunotherapy was not administered, skip to Question 136. If the immunotherapy is not listed, proceed to Question 134a, otherwise, skip to Question 135.
134a	Specify other immunotherapy			2953828	Provide the name of the immunotherapy administered to the patient.
135	Days to immunotherapy treatment from index date		-	5102411	Provide the number of days from the index date to the date of treatment with immunotherapy.
136	Targeted Therapy	□ Bevacizumab □ Everolimus □ Ridaforolimus □ Other (specify)		7028260	Select the targeted therapy administered to the patient. Note: If targeted therapy was not administered, skip to Question 138. If the targeted therapy regimen is not listed, proceed to Question 136a, otherwise, skip to Question 137.
136a	Specify targeted therapy			4308476	Provide the name of the targeted therapy administered to the patient.
137	Days to targeted therapy treatment from index date		-	5102411	Provide the number of days from the index date to the date of treatment with targeted therapy.
138	Radiation therapy administered type	□ 2D conventional □ 3D conformal □ Brachytherapy HDR □ Brachytherapy LDR □ IMRT □ Proton Beam	Stereotactic Body RT Stereotactic Radiosurgery WBRT Other (specify) Unspecified Not applicable	3028890	Provide the type of radiation therapy that was administered to the patient. Note: If radiation therapy was not administered, proceed to Question 140. If the radiation therapy is not listed, proceed to Question 138a, otherwise, skip to Question 139.
138a	Other radiation therapy		-	2195477	If the radiation therapy type is not included in the provided list, specify the type.
139	Days to radiation treatment from index date		-	5102411	Provide the number of days from the index date to the date of treatment with radiation therapy.
	/Recurrent Tumor Biospecin	nen Information		1	
140	Are you submitting a metastatic/recurrent tumor tissue sample?	☐ Yes ☐ No			Indicate whether a metastatic/ recurrent tumor biospecimen was collected for this ID3 case. Note: If yes, proceed to Question 141. If submitting an OTHER tissue sample, proceed to Question 232.
141	Metastatic/recurrent tissue biospecimen ordinal			6584266	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.
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.

	Enrollment: Endometrium	College Co	1	20	Car	7
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
144	Metastatic/recurrent tumor tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ Frozen ☐ 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 metastasis/ recurrence		6132218	Provide the number of days from the index date to the date of diagnosis of metastatic/recurrent disease.
146	Method of metastatic/ recurrent cancer sample procurement	□ Total hysterectomy and bilateral salping oophorectomy □ Radical hysterectomy □ Simple hysterectomy □ Supracervical hysterectomy □ Bilateral salpingo-oophorectomy □ Right salpingo-oophorectomy □ Left salpingo-oophorectomy □ Salpingo-oophorectomy □ Left oophorectomy □ Left oophorectomy □ Left salpingectomy □ Right salpingectomy □ Left salpingectomy □ Right salpingectomy □ Right salpingectomy □ Right salpingectomy □ Peritoneal salpingectomy □ Peritoneal biopsies □ Peritoneal washing □ Lymphadenectomy □ Core needle biopsy □ Fine needle aspirate □ Other Method (specify)		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.
147	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.
148	Metastatic/recurrent site	□ Endometrium □ Soft tissue □ Right ovary □ Left ovary □ Left ovary □ Left ovary □ Right fallopian □ tube □ Left fallopian □ Left tubo-ova □ Lung □ Liver □ Adrenal gland □ Brain □ Bone □ Other (specify	tube arian rian mph	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 148a, otherwise, skip to Question 149.
148a	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.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
149	Site of relapse	☐ Local ☐ Distant	2002506	If the primary tumor relapsed,
450	105.40	☐ Regional ☐ Not applicable	22222	provide the site of relapse.
150	ICD-10 code		3226287	Provide the ICD-10 code for the metastatic/recurrent tumor used to
				generate the model submitted to
				HCMI.
151	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.
152	Maintenance and/or		6119066	Provide the name(s) of the
	consolidation therapy			maintenance and/or consolidation
	administered prior to			therapy administered to the patient
	collection of metastatic/			prior to the collection of the
	recurrent tissue			metastatic/recurrent tissue used to
153	Days to start of		5102411	develop the model. Provide the number of days from the
155	maintenance and/or		3102411	index date to the date maintenance
	consolidation therapy			and/or consolidation therapy
	from index date			started.
154	Days to last known		5102431	Provide the number of days from the
	administration date of			index date to the last known date of
	maintenance and/or consolidation therapy			maintenance and/or consolidation
	from index date	·		therapy.
155	Is the patient still		6379568	Indicate whether the patient is still
	receiving treatment?	│ □ Yes │ □ No		undergoing maintenance and/or
		□ Unknown		consolidation therapy.
450			212222	
156	Disease status	☐ No evidence of disease	2188290	Provide the disease status following maintenance and/or consolidation
		☐ Progressive disease		therapy.
		☐ Stable disease		
		☐ Unknown		
		es for Metastatic/Recurrent Tumor Prognosis or Re		
157	Other tissue/organ involvement	☐ Not applicable ☐ Not identified	2793699	Select all sites of other tissue/organ involvement. This does not include
	involvement	☐ Right ovary		the primary tumor site.
		☐ Left ovary		the primary turnor site.
		☐ Ovary (side not specified)		
		☐ Right fallopian tube		
		☐ Left fallopian tube		
		Fallopian tube (side not specified)		
		☐ Vagina ☐ Right parametrium		
		☐ Left parametrium		
		☐ Parametrium (side not specified)		
		☐ Pelvic wall		
		☐ Bladder wall		
		☐ Bladder mucosa		
		☐ Rectal wall ☐ Bowel mucosa		
		☐ Omentum		
		☐ Pelvic lymph node(s)		
		☐ Peri-aortic lymph node(s)		
		☐ Other organ/tissue (specify)		
		Cannot be determined		

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
157a	Specify other tissue/organ involvement		7029867	If not included in the previous list, specify the other site of other tissue/organ involvement. This does not include the primary tumor site.
158	Peritoneal/Ascitic fluid	 □ Not submitted/unknown □ Negative for malignancy (normal/benign) □ Atypical and/or suspicious □ Malignant (positive for malignancy) □ Unsatisfactory/nondiagnostic 	6690681	Indicate the outcome of cytological analysis for peritoneal/ascitic fluid.
159	Lymph nodes	 □ No lymph nodes submitted or found □ All lymph nodes negative for tumor cells □ Positive lymph nodes 	7028287	Indicate whether the lymph nodes are involved with endometrial cancer as determined by pathologic assessment. Note: If lymph nodes are positive, proceed to Question 160, otherwise, skip to Question 170.
160	Number of pelvic nodes with micrometastases		7028289	Indicate the number of pelvic lymph nodes that have micrometastases present.
161	Number of pelvic nodes with macrometastases		7028288	Indicate the number of pelvic lymph nodes that have macrometastases present.
162	Number of pelvic nodes with isolated tumor cells		7028290	Indicate the number of pelvic lymph nodes that have isolated tumor cells present.
163	Total number of pelvic nodes examined		7028294	Indicate the total number of pelvic lymph nodes assessed for the presence of metastases and/or tumor cells.
164	Number of pelvic sentinel nodes examined		7028295	Indicate the total number of pelvic sentinel lymph nodes assessed for the presence of metastases and/or tumor cells.
165	Number of para-aortic nodes with micrometastases		7028292	Indicate the number of para-aortic lymph nodes that have micrometastases present.
166	Number of para-aortic nodes with macrometastases		7028291	Indicate the number of para-aortic lymph nodes that have macrometastases present.
167	Number of para-aortic nodes with isolated tumor cells		7028293	Indicate the total number of para- aortic lymph nodes that have isolated tumor cells present.
168	Total number of para- aortic nodes examined		7028296	Indicate the total number of para- aortic lymph nodes assessed for the presence of metastases and/or tumor cells.
169	Number of para-aortic sentinel nodes examined		7028297	Indicate the total number of para- aortic sentinel lymph nodes assessed for the presence of metastases and/or tumor cells.
170	Residual tumor	□ R0 □ RX □ R1 □ Not evaluated □ R2	2608702	Indicate the status of a tissue margin following surgical resection.
171	Was ER IHC performed?	☐ Yes ☐ No ☐ Unknown	6062416	Indicate whether estrogen receptor (ER) expression was assessed by immunohistochemistry (IHC). Note: If ER IHC was performed, proceed to Question 172, otherwise, skip to Question 173.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
172	ER expression by IHC		6690720	Indicate the status of estrogen
		Positive		receptor (ER) protein expression as
		□ Negative		determined by immunohisto-
		☐ Equivocal		chemistry (IHC).
173	Was PR IHC performed?		6062504	Indicate whether progesterone
	·	☐ Yes		receptor (PR) expression was
		□ No		assessed by immunohistochemistry
		☐ Unknown		(IHC). Note: If PR IHC was performed,
				proceed to Question 174, otherwise, skip
174	DD average in a built		6062672	to Question 175.
174	PR expression by IHC	☐ Positive	6063673	Indicate the status of progesterone
		☐ Negative		receptor (PR) protein expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
175	Was HER2 IHC		6063454	Indicate whether HER2 expression
	performed?	☐ Yes		was assessed by immunohisto-
		□ No		chemistry (IHC). Note: If HER2 IHC was
		☐ Unknown		performed, proceed to Question 176,
		_		otherwise, skip to Question 177.
176	HER2 expression by IHC	□ Positive	2957563	Indicate the expression of HER2 as
		□ Negative		assessed by immunohistochemistry
177	Was HER2 FISH/CISH	☐ Equivocal	6063447	(IHC). Indicate whether HER2 was assessed
1//	performed?		0003447	by fluorescence in situ hybridization
	performed:	□ Yes		(FISH) or chromogenic in situ
		□ No		hybridization (CISH). Note: If HER2
		☐ Unknown		FISH/CISH was performed, proceed to
				Question 178, otherwise, skip to
				Question 179.
178	HER2 status by FISH/CISH	☐ Amplified	2854089	Select the HER2 status as assessed
		☐ Not amplified ☐ Equivocal		by FISH/CISH.
179	Was MLH1 IHC	Lquivocai	6062411	Indicate whether MLH1 expression
1/3	performed?		0002411	was assessed by
	perrennea.	│ □ Yes │ □ No		immunohistochemistry (IHC). Note: If
		│ □ No │ □ Unknown		MLH1 IHC was performed, proceed to
		O O O O O O O O O O O O O O O O O O O		Question 180, otherwise, skip to
100				Question 181.
180	MLH1 expression by IHC	Positive	6063668	Indicate the status of MLH1 protein
		☐ Negative		expression as determined by immunohistochemistry (IHC).
181	MLH1 promoter	☐ Equivocal ☐ MLH1 promoter hypermethylation present	6033150	Indicate the methylation status of
101	methylation status	☐ MLH1 promoter hypermethylation present	0033130	the MLH1 promoter.
		☐ MLH1 promoter hypermethylation not		the mana promoten
		assessed		
		☐ Cannot be determined	<u>l</u>	
182	Was MSH2 IHC		6062412	Indicate whether MSH2 expression
	performed?	☐ Yes		was assessed by
		□ No		immunohistochemistry (IHC). Note: If
		☐ Unknown		MSH2 IHC was performed, proceed to
				Question 183, otherwise, skip to Question 184.
183	MSH2 expression by IHC	☐ Positive	6063669	Indicate the status of MSH2 protein
	2 2p. 233.577 27 1170	☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
184	Was PMS2 IHC		6062413	Indicate whether PMS2 expression
	performed?	☐ Yes		was assessed by immunohisto-
		□ No		chemistry (IHC). Note: If PMS2 IHC was
		☐ Unknown		performed, proceed to Question 185,
	<u> </u>		1	otherwise, skip to Question 186.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
185	PMS2 expression by IHC	☐ Positive	6063670	Indicate the status of PMS2 protein
		☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
186	Was MSH6 IHC	□ Vee	6062414	Indicate whether MSH6 expression
	performed?	☐ Yes ☐ No		was assessed by immunohisto-
		☐ Unknown		chemistry (IHC). Note: If MSH6 IHC was performed, proceed to Question 187,
		- Cinkilowii		otherwise, skip to Question 188.
187	MSH6 expression by IHC	☐ Positive	6063671	Indicate the status of MSH6 protein
		☐ Negative		expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
188	Microsatellite instability	☐ Stable	6032776	Indicate the status of the tumor for
		Low		microsatellite instability. (High at
		High		≥30% loci tested, low at 1%-29% loci
189	Was p53 IHC performed?	☐ Equivocal	6690705	tested.) Indicate whether p53 expression
109	was poor inc periorilleur	☐ Yes	0090703	was assessed by immunohisto-
		□ No		chemistry (IHC). Note: If p53 IHC was
		☐ Unknown		performed, proceed to Question 190,
				otherwise, skip to Question 191.
190	p53 expression by IHC	□ Normal	6690719	Indicate the status of p53 protein
		☐ Abnormal strong diffuse overexpression		expression as determined by
		(>90%) ☐ Abnormal null expression (complete loss of		immunohistochemistry (IHC).
		expression)		
		☐ Cannot be determined		
191	Was p16 IHC performed?		6690713	Indicate whether p16 expression
	' '	☐ Yes		was assessed by immunohisto-
		□ No		chemistry (IHC). Note: If p16 IHC was
		☐ Unknown		performed, proceed to Question 192,
192	p16 expression by IHC	☐ Positive	6690724	otherwise, skip to Question 193. Indicate the status of p16 protein
132	pro expression by inc	☐ Negative	0030724	expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
193	Mismatch repair status	☐ Evidence of MMR mutation by sequencing	6002208	Indicate the mismatch repair gene
	·	☐ Evidence of MMR protein loss by IHC		status.
		☐ MMR loss evidence hypermutation		
		phenotype (>10 mutations/Mb)		
		☐ No evidence of MMR alteration		
194	Was PTEN IHC	☐ Yes ☐ No	6062415	Indicate whether PTEN expression
	performed?	☐ Unknown		was assessed by immunohisto- chemistry (IHC).
195	PTEN expression by IHC	☐ Positive	6063672	Indicate the status of PTEN protein
200	The same of the sa	☐ Negative	0000072	expression as determined by
		☐ Equivocal		immunohistochemistry (IHC).
196	Was PTEN mutation	_	6063446	Indicate whether PTEN mutation
	analysis performed?	☐ Yes		analysis was performed. Note: If PTEN
		□ No		mutation analysis was performed,
		☐ Unknown		proceed to Question 197, otherwise, skip to Question 199.
197	Was a mutation in PTEN		6063529	Indicate whether a mutation in PTEN
	identified?	☐ Yes		was identified through mutation
		□ No		analysis. Note: if a PTEN mutation was
				identified, proceed to Question 198,
100	If DTEN we station	П вазов	6063736	otherwise, skip to Question 199.
198	If PTEN mutation	R130G R130P	6063736	Indicate the mutation identified for
	identified, which one?	☐ R130Q ☐ Other (specify) ☐ R130L		the PTEN gene. Note: If the PTEN mutation is not listed, proceed to
		L NIJOL		Question 198a, otherwise, skip to
				Question 199.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
198a	Other PTEN mutation		6816076	If the specific mutation of the PTEN
				gene identified was not included in
				the previous list, please specify the mutation.
199	Was PIK3CA mutation		6063445	Indicate whether PIK3CA mutation
133	analysis performed?	□ Yes	0003443	analysis was performed. <i>Note: If</i>
	, ,	□ No		PIK3CA mutation analysis was
		☐ Unknown		performed, proceed to Question 200,
200	Was a mutation in		6063524	otherwise, skip to Question 202. Indicate whether a mutation in
200	PIK3CA identified?		0003324	PIK3CA was identified through
	· ····································	□ Yes		mutation analysis. Note: If a PIK3CA
		□ No		mutation was identified, proceed to
				Question 201, otherwise, skip to
201	If PIK3CA mutation		7028281	Question 202. Indicate the mutation identified for
201	identified, which one?	☐ R88Q ☐ E545K ☐ H1047R ☐ H1047L	7020201	the PIK3CA gene. <i>Note: If the PIK3CA</i>
	, , , , , , , , , , , , , , , , , , , ,	∐ E542K		mutation is not listed, proceed to
		LI E542A		Question 201a, otherwise, skip to
		☐ E542Q ☐ E545G ☐ Other ☐ E542V ☐ E545Q ☐ Other		Question 202.
		(specify)		
201a	Other PIK3CA mutation		7028282	If the specific mutation of the
				PIK3CA gene identified was not
				included in the previous list, please
202	Was TP53 mutation		6063448	specify the mutation. Indicate whether TP53 mutation
202	analysis performed?	□ Yes	0003440	analysis was performed. <i>Note: If TP53</i>
		□ No		mutation analysis was performed,
		☐ Unknown		proceed to Question 203, otherwise, skip
203	Was a mutation in TP53		6063523	to Question 205. Indicate whether a mutation in TP53
	identified?	☐ Yes		was identified through mutation
		□ No		analysis. Note: If a TP53 mutation was
				identified, proceed to Question 204,
204	If TP53 mutation	—	7028281	otherwise, skip to Question 205. Indicate the mutation identified for
204	identified, which one?	□ R273H □ R248W	7020201	the TP53 gene. <i>Note: If the TP53</i>
		□ R273C □ R175H □ R273S		mutation is not listed, proceed to
		□ R273I □ R175C		Question 204a, otherwise, skip to
		☐ R248Q ☐ Other (specify)		Question 205.
204a	Other TP53 mutation		6101683	If the specific mutation of the TP53
				gene identified was not included in
				the previous list, please specify the
				mutation.
Additional 205		r Biospecimen Information (if applicable)	<u> </u>	A bigspagimen obtained from a single
205	Are you submitting an additional metastatic/			A biospecimen obtained from a single site at a single timepoint in
	recurrent tumor tissue			progression that is portioned for both
	sample?			sequencing and model generation
				counts as 1 single tumor specimen. A
		□ Yes		biospecimen obtained from another site or at a later timepoint in
		□ No		progression that is portioned for both
				sequencing and model generation
				counts as a second single tumor
				specimen. Note: If yes, proceed to Question 206, otherwise, skip to
				Question 222.

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Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
206	Metastatic/recurrent		6584266	Please provide a number to identify
	tissue biospecimen ordinal			which biospecimen this is in the sequence. The first biospecimen
	Ordinal			should be number "1," the second
				should be number "2," etc.
207	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.
208	BPC submitter ID (if		6584919	Please provide the BPC-generated ID
	available)			for this sample as it will appear on
				the Sample Submission Form
209	Metastatic/ recurrent		5432521	transmitted to the BPC. Provide the method used to
203	tumor tissue sample	☐ Cryopreserved ☐ OCT	3432321	preserve the metastatic/recurrent
	preservation method	☐ FFPE ☐ Snap frozer	ı	tumor tissue sample collected for
		☐ Frozen — Shap Heles		molecular characterization.
210	Number of days from		6132218	Provide the number of days from the
	index date to date of diagnosis of additional			index date to the date of diagnosis of additional metastatic/recurrent
	metastasis/ recurrence			disease.
211	Method of metastatic/	☐ Total hysterectomy and bilateral salpingo	o- 6587389	Indicate the procedure performed to
	recurrent cancer sample	oophorectomy		obtain the metastatic/recurrent
	procurement	☐ Radical hysterectomy☐ Simple hysterectomy		tumor tissue. Note: If the method of procurement is not listed, proceed to
		☐ Supracervical hysterectomy		Question 211a, otherwise, skip to
		☐ Bilateral salpingo-oophorectomy		Question 212.
		☐ Right salpingo-oophorectomy		
		Left salpingo-oophorectomy		
		☐ Salpingo-oophorectomy, side not specific☐ Right oophorectomy	ea	
		☐ Left oophorectomy		
		☐ Bilateral salpingectomy		
		☐ Right salpingectomy		
		☐ Left salpingectomy ☐ Salpingectomy, side not specified		
		☐ Vaginal cuff resection		
		□ Omentectomy		
		☐ Peritoneal biopsies		
		☐ Peritoneal washing		
		☐ Lymphadenectomy☐ Core needle biopsy		
		☐ Fine needle aspirate		
		☐ Other Method (specify)		
211a	Other method of cancer		6587390	If the procedure performed to
	sample procurement			obtain the tumor tissue is not
				included in the provided list, specify the procedure.
212	Number of days from		3288495	Provide the number of days from the
	index date to date of			index date to the date of the
	metastatic/ recurrent			procedure that produced the
	sample procurement			metastatic/recurrent tumor tissue submitted for HCMI.
213	Metastatic/ recurrent site		6587394	Select the site from which the
		☐ Endometrium ☐ Soft tissue ☐ Right ovary	330,334	metastatic/recurrent tissue used to
		☐ Lower uterine ☐ Left ovary		develop the model was derived.
		segment		Note: If the metastatic/recurrent site is not listed, proceed to Question 213a,
		tube		otherwise, skip to Question 214.
				otnerwise, skip to Question 214.

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	ce Site (TSS) Name: By:		nent: Endometrium HCMI Identifier (ID3): on Date (MM/DD/YYYY):		
213a	Other metastatic/	□ Exta-abdominal lymph node(s) □ Peritoneum □ Lung □ Liver □ Adrenal gland □ Brain □ Bone	☐ Left fallopian tube ☐ Right tubo-ovarian ☐ Left tubo-ovarian ☐ Pelvic lymph node(s) ☐ Para-aortic lymph node(s) ☐ Other (specify)	6587395	If not included in the previous list,
213a	recurrent site			6587395	specify the site from which the metastatic/recurrent tissue used to develop the model was derived.
214	Site of relapse	☐ Local ☐ Regional ☐ Distant ☐ Not applicable		2002506	If the primary tumor relapsed, provide the site of relapse.
215	ICD-10 code			3226287	Provide the ICD-10 code for the metastatic/recurrent tumor used to generate the model submitted to HCMI.
216	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

217

218

219

220

221

Maintenance and/or

consolidation therapy

administered prior to

recurrent tissue

Days to start of

from index date

from index date

Disease status

Is the patient still receiving treatment?

Days to last known

maintenance and/or

consolidation therapy

administration date of

☐ Yes

□ No

☐ Unknown

□ No evidence of disease

□ Progressive disease

☐ Stable disease☐ Unknown

maintenance and/or consolidation therapy

collection of metastatic/

HCMI.

started.

therapy.

therapy.

Provide the name(s) of the

prior to the collection of the

and/or consolidation therapy

develop the model.

maintenance and/or consolidation

therapy administered to the patient

metastatic/recurrent tissue used to

Provide the number of days from the

index date to the date maintenance

Provide the number of days from the

index date to the last known date of maintenance and/or consolidation

Indicate whether the patient is still

Provide the disease status following

maintenance and/or consolidation

undergoing maintenance and/or

consolidation therapy.

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6379568

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Additional Metastatic/Recurrent Tumor Prognostic/Predictive/Lifestyle Features for Additional Metastatic/Recurrent Tumor Prognosis or Responsiveness to Treatment (Note: Questions 157-204 may be repeated to capture clinical molecular characterization information for additional metastatic/recurrent biospecimens.)

Metastati	c/Recurrent Tumor Model Inj	formation		
222	METASTATIC/ RECURRENT model		6594587	Please provide a number to identify which biospecimen this is in the
				'
	biospecimen ordinal			sequence. Note: The first biospecimen
				should be number "1," the second
				should be number "2," etc.

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Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):
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Question	Question Text	Data Entry Options		CDE ID	Instruction Text
223	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.
224	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 to the BPC.
225	Model's METASTATIC/			6586035	Enter the CMDC Sample ID of the
	RECURRENT tumor tissue				METASTATIC/RECURRENT tissue
	CMDC sample ID				from which this model is derived.
226	Model's METASTATIC/			6584266	Enter the biospecimen ordinal
	RECURRENT tumor tissue				of the METASTATIC/RECURRENT
	biospecimen ordinal				tissue from which this model is
					derived.
Additiona	l Metastatic/Recurrent Biosp	ecimen Tumor Model Info	rmation (if applicable)	ı	
227	METASTATIC/			6594587	Please provide a number to identify
	RECURRENT model			0004007	which biospecimen this is in the
	biospecimen ordinal				sequence. Note: The first biospecimen
	2100pconnen ordinar				should be number "1," the second
					should be number "2," etc.
228	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.
229	BPC submitter ID (if			6584919	Please provide the BPC-generated ID
223	available)			0384313	for this sample as it will appear on
	avaliable)				the Sample Submission Form
					transmitted to the BPC.
230	Model's METASTATIC/			6586035	Enter the CMDC Sample ID of the
230	RECURRENT tumor tissue			0380033	· ·
					METASTATIC/RECURRENT tissue from which this model is derived.
224	CMDC sample ID			CE042CC	
231	Model's METASTATIC/			6584266	Enter the biospecimen ordinal of the
	RECURRENT tumor tissue				METASTATIC/RECURRENT tissue
	biospecimen ordinal				from which this model is derived.
	specimen Information	T		ı	
232	Are you submitting an				Indicate whether an OTHER tissue
	OTHER tissue sample?	☐ Yes			sample (e.g. pre-malignant, non-
		□ No			malignant, or dysplastic tissue, etc.)
					was collected for HCMI for this case.
					Note: If yes, proceed to Question 233.
233	OTHER tissue			6584267	Please provide a number to identify
	biospecimen ordinal				which biospecimen this is in the
					sequence. Note: The first biospecimen
					should be number "1," the second
224	CNADC			CE0C005	should be number "2," etc.
234	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.
235	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 to the BPC.
236	OTHER tissue sample			5432521	Provide the method used to
	preservation method	☐ Cryopreserved	□ ост		preserve the OTHER tissue sample
		☐ FFPE			collected for molecular
		☐ Frozen	☐ Snap frozen		characterization.
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
237	Other method of cancer sample procurement	☐ Endometrial biopsy ☐ Hysteroscopy ☐ Dilation and curettage (D&C) ☐ Fine needle aspirate ☐ Core needle biopsy ☐ Other Method (specify)	6587398	Provide the procedure performed to obtain the OTHER tissue. Note: If the method of procurement is not listed, proceed to Question 237a, otherwise, skip to Question 238.
237a	Specify method of OTHER tissue sample procurement		6587399	Specify the procedure performed to obtain the OTHER tissue.
238	Tissue type	☐ Pre-malignant ☐ Other (specify)	64784	Indicate the OTHER tissue type. Note: If the OTHER tissue type is not listed, proceed to Question 238a, otherwise, skip to Question 239.
238a	Specify tissue type		64785	Specify the OTHER tissue type if not in the provided list.
239	Anatomic site of OTHER tissue	☐ Endometrium ☐ Lower uterine segment ☐ 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 239a, otherwise, skip to Question 240.
239a	Specify anatomic site of OTHER tissue		6584916	Specify the site of OTHER tissue, if not in the previous list.
240	ICD-10 code		3226287	Provide the ICD-10 code for the OTHER tissue used to generate the model submitted to HCMI.
241	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 Informa	ation (if applicable)		
242	Are you submitting an additional OTHER tissue sample?	☐ Yes ☐ No		Indicate whether an additional OTHER tissue sample (pre-malignant, non-malignant, or dysplastic tissue, etc.) is being submitted for HCMI for this case. Note: If yes, proceed to Question 243, otherwise, skip to Question 253.
243	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.
244	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.
245	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.
246	OTHER tissue sample preservation method	☐ Cryopreserved ☐ FFPE ☐ OCT ☐ Frozen ☐ Snap frozen	5432521	Provide the method used to preserve the OTHER tissue sample collected for molecular characterization.
247	Other method of cancer sample procurement	 □ Endometrial biopsy □ Hysteroscopy □ Dilation and curettage (D&C) □ Fine needle aspirate 	6587398	Provide the procedure performed to obtain the OTHER tissue. Note: If the method of procurement is not listed, proceed to Question 247a, otherwise, skip to Question 248.

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		☐ Core needle biopsy☐ Other Method (specify)		
247a	Specify method of OTHER tissue sample procurement	——————————————————————————————————————	6587399	Specify the procedure performed to obtain the OTHER tissue.
248	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.
249	Tissue type	☐ Pre-malignant ☐ Other (specify)	64784	Indicate the OTHER tissue type. Note: If the OTHER tissue type is not listed, proceed to Question 249a, otherwise, skip to Question 250.
249a	Specify tissue type		64785	Specify the OTHER tissue type if not in the provided list.
250	Anatomic site of OTHER tissue	☐ Endometrium ☐ Lower uterine segment ☐ 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 250a, otherwise, skip to Question 251.
250a	Specify anatomic site of OTHER tissue		6584916	Specify the site of OTHER tissue, if not in the previous list.
251	ICD-10 code		3226287	Provide the ICD-10 code for the OTHER tissue used to generate the model submitted to HCMI.
252	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.
	ue Model Information			
253	OTHER tissue model biospecimen ordinal		6594590	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.
254	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.
255	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.
256	Model's OTHER tissue CMDC sample ID		6586035	Enter the CMDC Sample ID of the OTHER tissue from which this model is derived.
257	Model's OTHER tissue biospecimen ordinal		6584267	Enter the biospecimen ordinal of the OTHER tissue from which this model is derived.
	Other Tissue Model Informa	tion (if applicable)	T -	
258	OTHER tissue model biospecimen ordinal		6594590	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.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
259	CMDC model ID		6586036	Please provide the CMDC model ID
				for this sample as it will appear on
				tubes and the Sample Submission
		<u> </u>		Form transmitted to the BPC.
260	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 to the BPC.
261	Model's OTHER tissue		6586035	Enter the CMDC Sample ID of the
	CMDC sample ID			OTHER tissue from which this model
				is derived.
262	Model's OTHER tissue		6584267	Enter the biospecimen ordinal of the
	biospecimen ordinal			OTHER tissue from which this model
				is derived.