Enrollment: Wilms Tumor

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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.
Normal Con	ntrol Information			
4	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.
Tumor Tissu	ue Collected for Mo	lecular Characterization, Sample Information		
5	Tumor tissue sample preservation method	☐ FFPE ☐ Fresh ☐ OCT ☐ Snap frozen	5432521	Provide the method used to preserve the tumor tissue sample collected to be used for molecular characterization.
Cancer Mod	del Information		l	
6	Anatomic site of tumor from which model was derived	☐ Abdomen ☐ Ascites ☐ Kidney ☐ Lung ☐ Lymph node ☐ Other (specify)	4214629	Indicate the anatomic site of the tumor tissue used to generate the model for the HCMI. Note: If the anatomic site of tumor tissue is not listed, proceed to Question 6a, otherwise, skip to Question 7.
ба	Other anatomic site		5946219	If the anatomic site for the tumor submitted to HCMI is not included on the provided list, specify the anatomic site.
7	Method of cancer sample procurement	☐ Core needle biopsy ☐ Excisional biopsy ☐ Fine needle aspiration ☐ Incisional biopsy ☐ Tumor resection ☐ Other (specify)	3103514	Indicate the procedure performed to obtain the tumor tissue used to generate the model for HCMI. Note: If the method of sample procurement is not listed, proceed to Question 7a, otherwise, skip to Question 8.
7a	Other method of sample procurement		2006730	If the procedure performed to obtain the tumor tissue is not included in the provided list, specify the procedure.
8	Number of days from index date to date of cancer 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.

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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
9	ICD-10 code for	□ C64.1 □ C78.0	3226287	Provide the ICD-10 code for the tumor used to
	model tumor	□ C64.2 □ C78.7		generate the model submitted to HCMI.
		□ C64.9 □ C79.8		Note: If the ICD-10 code is not listed, proceed to
		☐ C77.9 ☐ Other (specify)		Question 9a, otherwise, skip to Question 10.
9a	Other ICD-10 code		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.
10	Tumor tissue type	☐ Premalignant	3288124	Provide the tumor tissue type for the
		☐ Primary		biospecimen used to produce the model for the
		☐ Recurrent		HCMI.
		☐ Metastatic		Note: If 'Metastatic' is selected, continue to
		☐ Additional primary		answer through Question 18. If the tissue is not
		□ NOS		'Metastatic', skip to Question 19.
Metastatic	Model Information (o	nly complete Questions 11-18 if 'Metastatic' wa	s selected in	Question 10)
11	Age at diagnosis		6032752	Provide the age (in days) of the patient when
	of metastasis			diagnosed with metastatic disease. If the
				patient's age is greater than 32,507 days (89
				years), please enter 32,507.
12	Number of days		6132218	Provide the number of days from the index date
	from index date to			to the date of diagnosis of metastatic disease.
	date of diagnosis			
	of metastasis			
13	Metastatic site	☐ Abdomen	6119068	Select the site from which the metastatic tissue
		Liver		used to develop the model was derived.
		Lung		Note: If the metastatic site is not listed, proceed
		☐ Lymph node(s); regional		to Question 13a, otherwise, skip to Question 14.
		Lymph node(s); distant		
		☐ Other (specify)		
13a	Other metastatic		3128033	If not included in the previous list, specify the site
	site			from which the metastatic tissue used to develop
			5110055	the model was derived.
14	Maintenance		6119066	If applicable, provide the name of the
	and/or			maintenance and/or consolidation therapy
	consolidation			administered to the patient prior to the collection
	therapy			of the metastatic tissue used to develop the
	administered			model.
	prior to collection			Note: If maintenance and/or consolidation
	of metastatic			therapy was not administered, skip to Question
15	tissue		F102411	Provide the number of days from the index date
15	Days from index date to start of		5102411	to the date maintenance and/or consolidation
	maintenance			therapy started.
	and/or consolidation			
16	therapy Days from index		65167	Provide the number of days from the index date
10	date to last known		03107	to the last known date of maintenance and/or
	date of			consolidation therapy.
	maintenance			consolidation therapy.
	and/or			
	consolidation			
	therapy treatment			
17	Is the patient still	☐ Yes	6379568	Indicate whether the patient is still undergoing
1/	receiving	□ No	03,3300	treatment.
	treatment?	☐ Unknown		ir caunciii.
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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
18	Disease status	☐ No evidence of disease	2188290	Provide the disease status following maintenance
		☐ Progressive disease		and/or consolidation therapy.
		☐ Stable disease		
		☐ Unknown		
Patient Info	rmation			
19	Gender	☐ Male	2200604	Provide the patient's gender using the defined
		☐ Female		categories. Identification of gender is based upon
		☐ Unspecified		self-report and may come from a form,
		- Onspecified		questionnaire, interview, etc.
20	Height		649	Provide the patient's height, in centimeters.
21	Weight		651	Provide the patient's weight, in kilograms.
22	Body mass index		2006410	If the patient's height and weight are not
	(BMI)			collected, provide the patient's body mass index
				(BMI).
23	Race		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
		☐ American Indian or Alaska Native		maintains tribal affiliation or community attachment. Asian: A person having origins in any of the peoples of
		☐ Asian		the Far East, Southeast Asia, or in the Indian
		☐ Black or African American		subcontinent including, for example, Cambodia, China,
		☐ Native Hawaiian or other Pacific Islander		India, Japan, Korea, Malaysia, Pakistan, the Phillippine
		☐ White		Islands, Thailand, and Vietnam.
		☐ Unknown		Black or African American: A person having origins in
		☐ Not allowed to collect		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.
24	Ethnicity		2192217	Provide the patient's ethnicity using the defined
		□ Hispania or Latina		categories.
		☐ Hispanic or Latino		Hispanic or Latino: A person of Mexican, Puerto Rican,
		☐ Not Hispanic or Latino ☐ Unknown		Cuban, Central or South American or other Spanish
		□ Not allowed to collect		culture or origin, regardless of race.
		Not allowed to collect		Not Hispanic or Latino: A person not meeting the
2-			2006076	definition of Hispanic or Latino.
25	Year of birth		2896954	Provide the year of the patient's birth. If the
				patient was born prior to 1928, insert the date
26	Familia Int.	П. С	F02222	1928.
26	Family history of	☐ Same	5832923	Has a first-degree relative of the patient been
	cancer	□ Different		diagnosed with a cancer of the same or a
		□ None		different type?
2=	6 11 11 1	☐ Unknown	2404575	
27	Smoking history	☐ Lifelong non-smoker (<100 cigarettes	2181650	Indicate the patient's history of tobacco smoking
		smoked in a lifetime)		as well as their current smoking status using the
		☐ Current smoker (includes daily and non-		defined categories.
		daily smokers)		
		☐ Current reformed smoker (duration not		
		specified)		
		☐ Current reformed smoker for >15 years		
	ĺ	☐ Current reformed smoker for ≤15 years		

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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
•	nor Diagnosis Infor				
28	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.
29	Patient age on index date			6379572	Provide the age (in days) of the patient on the index date. If the patient's age is greater than 32,507 days (89 years), please enter 32,507.
30	Morphology	☐ 8960/3 (Wilms tumo☐ Other (specify)	r)	3226275	Using the patient's pathology/laboratory report, provide the ICD-O-3 histology code of the primary tumor. Note: If the morphology is not listed, proceed to Question 30a, otherwise, skip to Question 31.
30a	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.
31	Tissue or organ of origin	☐ Kidney ☐ Other (specify)		3427536	Using the patient's pathology/laboratory report, select the primary site of the disease. Note: If the tissue or organ of origin is not listed, proceed to Question 31a, otherwise, skip to Question 32.
31a	Other tissue or organ of origin	□ Abdomen □ Accessory sinus □ Adrenal gland □ Anus □ Appendix □ Bladder □ Bone □ Breast □ Connective, subcutaneous and other soft tissues □ Esophagus □ Eye □ Gallbladder □ Gum □ Head, face or neck □ Heart □ Kidney □ Larynx □ Lip □ Liver □ Lung □ Lymph node □ Male genital organs □ Mediastinum □ Meninges □ Mouth □ Nasal cavity □ Nasopharynx □ Nervous system □ Oropharynx	Other ill-defined sites Ovary Palate Pancreas Penis Peripheral nerves and autonomic nervous system of trunk Peritoneum Pharynx Pituitary gland Prostate gland Rectosigmoid junction Renal pelvis Retroperitoneum Skin Small intestine Spinal cord Spleen Stomach Testis Thymus Thyroid gland Tongue Tonsil Trachea Unknown primary Utrinary system Uterus Vagina Vulva	3427536	If the primary site of the disease is not included on the previous list, select the primary site of the disease.

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Tissue Source Site (TSS) Name:	HCMI Identifier (ID3):	Constitution of the second	3	18	1	1	
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
32	Histological type		3294805	Provide the traditional surgical pathology text
				description of the histological tumor type.
33	Histology	☐ Favorable	4358735	Using the patient's pathology/laboratory report,
		☐ Unfavorable		select the histology of the tumor submitted to
		☐ Unknown		the HCMI.
34	Prior malignancy	☐ Yes	5832924	Indicate whether the patient has a history of
	(of the same	□ No		prior malignancy of the same cancer type.
	cancer type)	☐ Unknown		
35	Prior malignancy	☐ Yes	5878828	Indicate whether the patient has a history of
	(other cancer	□ No		prior malignancy of a different cancer type.
	type)	☐ Unknown		
36	Tumor stage	☐ Stage I	6013641	Using the patient's pathology/laboratory report,
	(pathological)	□ γ-Stage I		select the pathological stage as defined by
		☐ Stage II		SIOP/COG/NWTSG.
		□ γ-Stage II		
		☐ Stage III		
		□ γ-Stage III		
		☐ Stage IV		
37	Metastasis at	☐ Metastatic	3438571	Indicate whether there was evidence of
	diagnosis	☐ Non-metastatic (confirmed)		metastasis at the time of diagnosis of the
	assessment status	☐ Non-metastatic (unconfirmed)		primary tumor.
38	Metastatic site(s)	☐ Abdomen	3029815	Indicate all the site(s) of metastasis at the time
	at diagnosis	Liver		of diagnosis of the primary tumor.
		Lung		Note: If the metastatic site(s) is not listed,
		☐ Lymph node(s); regional		proceed to Question 38a, otherwise, skip to
		☐ Lymph node(s); distant		Question 39.
		☐ Other (specify)		
38a	Specify metastatic	· , , , ,	3128033	If the site(s) of metastasis at the time of
	site(s)			diagnosis of the primary tumor is not included in
				the provided list, specify the site(s).
39	Site of relapse	☐ Local	2002506	If the primary tumor relapsed, select all sites of
	'	☐ Regional		relapse.
		☐ Distant		Note: If the primary tumor did not relapse,
		☐ Not applicable		select 'Not applicable'.
40	Tumor weight	, ,	3184957	Provide the weight in grams of the primary
	(gross pathology)			tumor.
41	Tumor laterality	☐ Bilateral	827	Indicate the side of the body on which the
	,	□ Left		primary tumor was located.
		☐ Right		,
		☐ Unilateral; side not specified		
		☐ Midline		
		□ Unknown		
Prognostic/	Predictive/Lifestyle F	eatures for Tumor Prognosis or Responsiveness	to Treatment	
42	Anaplasia	☐ Absent	4925534	Indicate the presence of anaplasia in the
		☐ Focal		primary tumor.
		☐ Diffuse		·
		☐ Unknown		
43	Wilms tumor	☐ Beckwith-Wiedemann syndrome (BWS)	3078770	Indicate whether the patient has any known
	related clinical	☐ Denys-Drash syndrome (DD)	33.37.73	clinical syndromes associated with Wilms tumor.
	syndrome name	☐ Hemihypertrophy (HH)		The second secon
	,	☐ WAGR syndrome (WAGR)		
		Other		
		□ None		
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Enrollment: Wilms Tumor



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
Treatment I	nformation			
44	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: Radiation therapy is addressed in Questions 52-53. Pharmaceutical therapy is addressed in Questions 45-51.
45	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 46-47. Immunotherapy is addressed in Questions 48-49. Targeted therapy is addressed in Questions 50-51.
46	Neoadjuvant chemotherapeutic regimen	 □ Cyclophosphamide □ Dactinomycin □ Doxorubicin □ Etoposide □ Ifosfamide, Carboplatin, Cisplatin □ Irinotecan □ Vincristine □ Vincristine, actinomycin-D, cyclophosphamide (VAC) □ Vincristine, doxorubicin, cyclophosphamide, ifosfamide, etoposide (VDC/IE) □ Vincristine, actinomycin-D, cyclophosphamide, vincristine, irinotecan (VAC/VI) □ Ifosfamide, carboplatin, etoposide (ICE) □ Vincristine, irinotecan, temozolomide (VIT) □ High-dose methotrexate, doxorubicin, cisplatin (MAP) □ Other (specify) □ Chemotherapy not given 	2853313	Select all chemotherapeutics used for neoadjuvant therapy. Note: If neoadjuvant chemotherapy was not given, skip to Question 48. If the neoadjuvant chemotherapeutic regimen is not listed, proceed to Question 46a, otherwise, skip to Question 47.
46a	Other neoadjuvant chemotherapeutic regimen		62694	If the neoadjuvant therapy is not included in the provided list, specify neoadjuvant therapy.
47	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.
48	Immunotherapy name, specify		2185614	Specify the name of the immunotherapy administered. Note: If immunotherapy was not given, skip to Question 50.
49	Days to immunotherapy treatment from index date		5102411	Provide the number of days from the index date to the date of treatment with immunotherapy.

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·	Enrollment: Wilms Tumor	
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Question	Question Text	Data Entry Options	CDE ID	Instruction Text
50	Targeted molecular therapy name, specify		2842797	Specify the targeted therapy administered to the patient. Note: If targeted therapy was not given, skip to Question 52.
51	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.
52	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, skip the remaining questions. If the radiation therapy is not listed, proceed to Question 52a, otherwise, skip to Question 53.
52a	Other radiation therapy	·	2195477	If the radiation therapy type is not included in the provided list, specify the type.
53	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.