V1.0

Follow-Up: Pediatric Liver

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

Form Notes: A Follow-Up Form should be completed for each HCMI case upon notice of model establishment and molecular characterization success from Leidos. All information provided on this form should include activity from the "Date of Last Contact" provided on the HCMI Enrollment Form to the most recent date of contact with the patient or the patient's medical record.

tient Status Number of days from index date to date of last follow-up Vital status	☐ Initial pathologic diagnosis ☐ Sample procurement ☐ First patient visit	2003301 5845012 6154722 3008273	Provide the patient's ID2 (this ID will only be used by IMS for internal quality control). Provide the HCMI-specific anonymized ID (ID3). 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.
tient Status Number of days from index date to date of last follow-up	☐ Sample procurement	6154722	Provide the HCMI-specific anonymized ID (ID3). 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.
tient Status Number of days from index date to date of last follow-up	☐ Sample procurement	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.
tient Status Number of days from index date to date of last follow-up	☐ Sample procurement		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.
Number of days from index date to date of last follow-up	☐ Sample procurement	3008273	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.
Number of days from index date to date of last follow-up	☐ Sample procurement	3008273	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.
Number of days from index date to date of last follow-up		3008273	alternative index date is used, indicate it here and use it for all interval calculations.
Number of days from index date to date of last follow-up	——————	3008273	use it for all interval calculations.
Number of days from index date to date of last follow-up		3008273	
Number of days from index date to date of last follow-up		3008273	
from index date to date of last follow-up		3000273	Provide the number of days from the index date
date of last follow-up			to the last date of follow-up with the patient or
follow-up			last contact with the medical record.
/ital status			last contact with the medical record.
vitai status	□ Alive	5	Indicate whether the patient is alive, dead, or lost
	□ Dead		to follow-up at the date of last contact.
	☐ Lost to follow-up		Note: If the patient is deceased, continue to
		2.65.55	Question 6, otherwise skip to Question 8.
•		31654/5	Provide the number of days from the index date to the date of death.
			to the date of death.
Cause of death	☐ Related to this cancer	2554674	Indicate the patient's cause of death.
oudse or death.		255 167 1	maisure the patient s sause of acutin
	☐ Related to another cancer		
	☐ Other (specify)		
	☐ Unknown		
Other cause of		4783275	If the cause of death is not included in the
death			provided list, specify the cause of death.
		2188290	Provide the last known state of the patient's
follow-up			tumor up to the point of current follow-up data
	_		submission.
formation	L CHRIGWII		
Was surgery		2978013	Indicate whether surgery was performed to treat
performed as part	☐ Yes		the primary tumor.
of the primary	□ No		Note: If the patient did not receive surgical
disease treatment	☐ Unknown		treatment, skip to Question 11.
plan?		2000225	
·		3008335	Provide the number of days from the index date
			to the date of surgical treatment.
_			
		3397567	Indicate whether the patient received systemic
adjuvant therapy	□ Yes		adjuvant pharmaceutical therapy.
administered?	□ No		Note: If the patient did have systemic adjuvant
	☐ Unknown		therapy, the Pharmaceutical Supplemental Form
			should be completed.
Was adjuvant	_	2005312	Indicate whether the patient had adjuvant
radiation therapy			radiation therapy.
administered?			Note: If the patient had adjuvant radiation
	⊔ Unknown		therapy, the Radiation Supplemental Form should be completed.
frdC CdDfc FeV pod pN frdtiVaa V r	other cause of eath risease status at ollow-up ormation vas surgery erformed as part of the primary isease treatment lan? Illumber of days rom index date to ate of surgical reatment vas systemic djuvant therapy dministered?	rom index date to ate of death ause of death ause of death ause of death Related to this cancer Non-cancer related Related to another cancer Other (specify) Unknown Other cause of eath Disease status at Stable disease Progressive disease Unknown Other specify Unknown Oth	rom index date to ate of death ause of eath ause of death ause of disease ause of dise

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	Follow-Up: Pediatric Liver		Y. 1
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Completed By:	Completion Date (MM/DD/YYYY):		

Pharmaceutical Supplemental Form

Form Notes: A Pharmaceutical Supplemental Form should be completed for each HCMI case for which the patient received adjuvant pharmaceutical therapy. All information provided on this form should include activity from the "Date of Last Contact" provided on the HCMI Enrollment Form to the most recent date of contact with the patient or the patient's medical record.

Question	Question Toyt	Data Entry Ontions	CDE ID	Instruction Toyt
Question 1	Question Text Was cytotoxic	Data Entry Options	5628399	Instruction Text Indicate whether the patient received cytotoxic
1	chemotherapy administered?	☐ Yes ☐ No ☐ Unknown	2028399	chemotherapy. Note: If cytotoxic chemotherapy was administered, proceed to the "Cytotoxic Chemotherapy" section, Questions 2-5.
2	Was immunotherapy (cellular and immune checkpoint) administered?	☐ Yes ☐ No ☐ Unknown	3057655	Indicate whether the patient received immunotherapy. Note: If immunotherapy was administered, proceed to the "Immunotherapy" section, Questions 6-9.
3	Was targeted therapy (small molecule inhibitors and targeted antibodies) administered?	☐ Yes ☐ No ☐ Unknown	2785850	Indicate whether the patient received targeted therapy. Note: If targeted therapy was administered, proceed to the "Targeted Therapy" section, Questions 10-13.
Cytotoxic Cl	nemotherapy		,	,
2	Chemotherapeutic administered	□ 5-fluorouracil □ Carboplatin □ Cisplatin □ Doxorubicin □ Etoposide □ Ifosfamide □ Pirarubicin □ Vincristine □ Vinorelbine □ 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)	2853873	Select the chemotherapeutic used for therapy. Note: Questions 2-5 are repeatable as needed to capture each individual chemotherapeutic administered. If the chemotherapeutic is not included in the provided list, proceed to Question 2a, otherwise, skip to Question 3.
2a	Other chemotherapeutic		2514640	If the adjuvant therapy is not included in the provided list, specify adjuvant therapy.
3	Days from index date to start of pharmaceutical treatment		5102411	Provide the number of days from the index date to the date of initiation of treatment with adjuvant pharmaceutical therapy.
4	Days from index date to last known date of		65167	Provide the number of days from the index date to the last known date of pharmaceutical treatment.

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	pharmaceutical			
Question	treatment Question Text	Data Fatas Ontions	CDE ID	Instruction Text
5	•	Data Entry Options ☐ Yes	6379568	Indicate whether the patient is still undergoing
Э	Is the patient still		03/9508	,
	receiving	□ No		treatment.
Immunothe	treatment?	Unknown		
6	Immunotherapy		2185614	Select the immunotherapy administered.
Ü	administered		2103014	Note: Questions 6-9 are repeatable as needed to capture each individual immunotherapy administered.
7	Days from index date to start of immunotherapy treatment		5102411	Provide the number of days from the index date to the date of the initiation of treatment with immunotherapy.
8	Days from index date to last known date of immunotherapy treatment		65167	Provide the number of days from the index date to the last known date of immunotherapy treatment.
9	Is the patient still	☐ Yes	6379568	Indicate whether the patient is still undergoing
	receiving	□ No		treatment.
	treatment?	☐ Unknown		
Targeted T	herapy		•	
10	Targeted therapy administered	□ Sorafenib	6010754	Select the targeted therapy administered. Note: Questions 10-13 are repeatable as needed to capture each individual targeted therapy
		☐ Other (specify)		administered.
		(opea)		If the targeted therapy is not included in the
				provided list, proceed to Question 10a,
				otherwise, skip to Question 11.
10a	Other targeted therapy		4308476	If the targeted therapy is not included in the provided list, specify the therapy.
11	Days from index		5102411	Provide the number of days from the index date
11	date to start of		3102411	to the date of initiation of treatment with
	targeted therapy			targeted therapy.
	treatment			targeted therapy.
12	Days from index		65167	Provide the number of days from the index date
	date to last known		3320,	to the last known date of targeted therapy
	date of targeted			treatment.
	therapy treatment			
13	Is the patient still	☐ Yes	6379568	Indicate whether the patient is still undergoing
-	receiving	□ No		treatment.
	treatment?	☐ Unknown		

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Completed By:	Completion Date (MM/DD/YYYY):	

Radiation Supplemental Form

Form Notes: A Radiation Supplemental Form should be completed for each HCMI case for which the patient received adjuvant radiation therapy. All information provided on this form should include activity from the "Date of Last Contact" provided on the HCMI Enrollment Form to the most recent date of contact with the patient or the patient's medical record.

Question	Question Text	Data Entry Options	CDE ID	Instruction Text
1	Radiation therapy administered type	□ 2D conventional □ 3D conformal □ Brachytherapy HDR □ Brachytherapy LDR □ IMRT □ Proton Beam □ Stereotactic Body RT □ Stereotactic Radiosurgery □ WBRT □ Other (specify) □ Unspecified	3028890	Provide the type of adjuvant radiation therapy that was administered to the patient, if not collected on the enrollment form for this patient. Note: If the radiation therapy type is not included in the provided list, proceed to Question 1a, otherwise, skip to Question 2.
1a	Other radiation therapy		3028890	If the radiation therapy type is not included in the provided list, specify the type.
2	Days from index date to start of adjuvant radiation therapy treatment		5102411	Provide the number of days from the index date to the date of treatment with adjuvant post-operative radiation therapy.
3	Days from index date to last known date of adjuvant radiation therapy treatment		65167	Provide the number of days from the index date to the last known date of radiation therapy treatment.
4	Is the patient still receiving treatment?	☐ Yes ☐ No ☐ Unknown	6379568	Indicate whether the patient is still undergoing treatment.