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	Follow-Up: Brain	
Fissue 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. This form should be used for the following Brain Cancers: Embryonal Tumor, Medulloblastoma, Diffuse Midline Glioma, and Lower Grade Glioma.

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).
2	103		3643012	Provide the ncivii-specific allohymized ib (ibs).
3	Index date		6154722	Select the reference date used to calculate time
		☐ Initial pathologic diagnosis		intervals (e.g. days to treatment). Date of initial
		☐ Sample procurement		pathologic diagnosis is the HCMI standard and should be used unless it is unavailable. If an
		☐ First patient visit		alternative index date is used, indicate it here and
				use it for all interval calculations.
	Patient Status		<u>_</u>	
4	Number of days		3008273	Provide the number of days from the index date
	from index date to			to the last date of follow-up with the patient or
	date of last			last contact with the medical record.
5	follow-up Vital status		5	Indicate whether the patient is alive, dead, or lost
5	Vital Status	☐ Alive	3	to follow-up at the date of last contact.
		☐ Dead		Note: If the patient is deceased, continue to Question
		☐ Lost to follow-up		6, otherwise skip to Question 8.
6	Number of days		3165475	Provide the number of days from the index date
	from index date to			to the date of death.
	date of death	_		
7	Cause of death	Related to this cancer	2554674	Indicate the patient's cause of death.
		□ Non-cancer related		
		☐ Related to another cancer☐ Other (specify)		
		☐ Unknown		
7a	Other cause of		4783275	If the cause of death is not included in the
	death			provided list, specify the cause of death.
8	Disease status at	☐ No evidence of disease	2188290	Provide the last known state of the patient's
	follow-up	☐ Stable disease		tumor up to the point of current follow-up data
		☐ Progressive disease		submission.
Troatmont	Information	☐ Unknown		
9	Information Was surgery		2978013	Indicate whether surgery was performed to treat
	performed as part	☐ Yes	2373323	the primary tumor.
	of the primary	□ No		Note: If the patient did not receive surgical treatment,
	disease treatment	☐ Unknown		skip to Question 11.
	plan?			
10	Number of days		3008335	Provide the number of days from the index date
	from index date to			to the date of surgical treatment.
	date of surgical			
11	treatment Was systemic		3397567	Indicate whether the patient received systemic
11	adjuvant therapy	☐ Yes	3337307	adjuvant pharmaceutical therapy.
	administered?	□ No		Note: If the patient did have systemic adjuvant
		☐ Unknown		therapy, the <u>Pharmaceutical Supplemental Form</u>
				should be completed.
12	Was adjuvant	□ Vos	2005312	Indicate whether the patient had adjuvant
	radiation therapy	│ □ Yes │ □ No		radiation therapy.
	administered?	□ Unknown		Note: If the patient had adjuvant radiation therapy, the <u>Radiation Supplemental Form</u> should be
				completed.

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Completed By:	Completion Date (MM/DD/YYYY):	1		7

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 Text	Data Entry Options		CDE ID	Instruction Text
Question	Question Text	Data Entry Options		CDE ID	IIISTI GETTOIT TEXT
1	Was cytotoxic chemotherapy administered?	☐ Yes ☐ No ☐ Unknown		5628399	Indicate whether the patient received cytotoxic chemotherapy. Note: If cytotoxic chemotherapy was administered, proceed to the "Cytotoxic Chemotherapy" section, Questions 4-7.
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 8-11.
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 12-15.
Cytotoxic Cl	hemotherapy			1	
4	Chemotherapeutic administered	□ Bevacizumab □ Carboplatin □ Carmustine □ Cisplatin □ Cyclophosphamide □ Cytarabine □ Etoposide □ Hydroxyurea □ Irinotecan	□ Lomustine □ Panobinostat □ Prednisone □ Procarbazine □ Temozolomide □ Vincristine □ Vorinostat □ Other (specify) □ Chemotherapy not given	2853873	Select the chemotherapeutic(s) used for therapy. Note: Questions 4-7 are repeatable as needed to capture each individual chemotherapeutic administered. If the chemotherapeutic is not included in the provided list, proceed to Question 4a, otherwise, skip to Question 5.
4a	Other		8 -	2514640	If the adjuvant therapy is not included in the
5	chemotherapeutic Days from index date to start of pharmaceutical treatment			5102411	provided list, specify adjuvant therapy. Provide the number of days from the index date to the date of initiation of treatment with adjuvant pharmaceutical therapy.
6	Days from index date to last known date of pharmaceutical treatment			65167	Provide the number of days from the index date to the last known date of pharmaceutical treatment.
7	Is the patient still receiving treatment?	☐ Yes ☐ No ☐ Unknown		6379568	Indicate whether the patient is still undergoing treatment.
Immunothe		ı			
8	Other Immunotherapy administered			2953828	Specify the immunotherapy. Note: Questions 8-11 are repeatable as needed to capture each individual immunotherapy administered.
9	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.

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	Follow-Up: Brain	
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
10	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.
11	Is the patient still receiving treatment?	☐ Yes ☐ No ☐ Unknown	6379568	Indicate whether the patient is still undergoing treatment.
Targeted Ti	herapy			
12	Targeted therapy		4308476	Specify targeted therapy. Note: Questions 12-15 are repeatable as needed to capture each individual targeted therapy
13	Days from index date to start of targeted therapy treatment		5102411	Provide the number of days from the index date to the date of initiation of treatment with targeted therapy.
14	Days from index date to last known date of targeted therapy treatment		65167	Provide the number of days from the index date to the last known date of targeted therapy treatment.
15	Is the patient still receiving treatment?	☐ Yes ☐ No ☐ Unknown	6379568	Indicate whether the patient is still undergoing treatment.

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Completed By:	Completion Date (MM/DD/YYYY):		1 3 6 5 S	

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.

Radiation Supplemental Form

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.