Follow-Up Form Endometrial (UCEC)

V4.02 071912

Instructions: The Follow-up Form is to be completed 12 months after a case enters the Biospecimen Core Resource (BCR). All information provided on this form includes activity from the "Date of Last Contact" provided on the TCGA Enrollment Form to the most recent date of contact with the patient. This form should only be completed by the Tissue Source Site if updated information can be provided to TCGA. Please direct any questions to the Clinical Outreach team at the BCR.

Please note the following definitions for the "Unknown" and "Not Evaluated" answer options on this form.

Unknown: This answer option should only be selected if the TSS does not know this information after all efforts to obtain the data have been exhausted. If this answer option is selected for a question that is part of the TCGA required data set, the TSS must complete a discrepancy note providing a reason why the answer is unknown.

Not Evaluated: This answer option should only be selected by the TSS if it is known that the information being requested cannot be obtained. This could be because the test in question was never performed on the patient or the TSS knows that the information requested was never disclosed.

Tissue Source Site (TSS): TSS Id	lentifier: TSS Unique Patient Identifier:	

Completed By (Interviewer Name on OpenClinica): ____

_Completed Date: ____

General Information			
#	Data Element	Entry Alternatives	Working Instructions
1	Has this TSS received permission from the NCI to provide time intervals as a substitute for requested dates on this form?	□ Yes □ No	Please note that the time intervals must be recorded in place of dates where designated throughout this form if you have selected "yes" in the box. Provided time intervals must begin with the date of initial pathologic diagnosis (i.e., biopsy or resection). Only provide Interval data if you have received permission from the NCI to provide time intervals as a substitute for requested dates on this form.
2	Is this Patient Lost to Follow-up?	□ Yes □ No	Indicate whether the patient is lost to follow-up, as defined by the ACoS Commission on Cancer. This only includes cases where updated follow-up information has not been collected within the past 15 months and all efforts to contact the patient have been exhausted (this includes reviewing the Social Security death index). If the patient is lost to follow-up, the remaining questions can be left unanswered. <u>61333</u> If the patient is deceased and a TCGA follow-up form has not yet been completed, the answer to this question should be "no," and the remaining applicable questions should be completed.

Follow-Up Information

#	Data Element	Entry Alternatives	Working Instructions
3	Adjuvant (Post- Operative) Radiation Therapy	□ Yes □ No □ Unknown	Indicate whether the patient had adjuvant/ post- operative radiation therapy. <u>2005312</u> If the patient did have adjuvant radiation, the Radiation Supplemental Form should be completed.
4	Adjuvant (Post- Operative) Pharmaceutical Therapy	☐ Yes □ No □ Unknown	Indicate whether the patient had adjuvant/ post- operative pharmaceutical therapy. <u>3397567</u> If the patient did have adjuvant pharmaceutical therapy, the Pharmaceutical Supplemental Form should be completed.
5	Has the patient ever taken menopausal hormone therapy?	 Current User Former User Never Used Unknown 	Indicate whether the patient, at any time, used menopausal hormone therapy. <u>3012813</u>
6	Has the patient ever taken oral contraceptives?	 Current User Former User Never Used Unknown 	Indicate whether the patient, at any time, used oral contraceptives. <u>3104217</u>

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#	Data Element	Entry Alternatives	Working Instructions
		Current User	Indicate whether the patient, at any time, used Tamoxifen.
	Has the patient ever	Former User	3104234
7	taken Tamoxifen?	□ Never Used	
	taken ramoxiten.		
			Indicate whether the patient has a history of hypertension.
0	Unantonaion		2183378
8	Hypertension	Unknown	2100070
			Indicate whether the patient has, at any time, been diagnosed
	Has the patient ever	□ Yes	with diabetes by a physician. This includes borderline and
9	been diagnosed with		gestational diabetes.
	diabetes by a physician?		<u>2716085</u>
			Provide the number of full term pregnancies the patient has
		□1	had. <u>3012512</u>
10	Number of full term		<u>3012312</u>
10	pregnancies		
		□ 4+	
		Unknown	
	Has the nationt had	□ Yes	Indicate whether the patient has a history of colorectal cancer.
11	Has the patient had colorectal cancer?	🗖 No	<u>2684753</u>
	colorectal calleer?	🗖 Unknown	
	Tumor Status	□ Tumor free	Indicate whether the patient was tumor/disease free at the
12	(at time of last contact or	With tumor	date of last contact or death.
12	death)		<u>2759550</u>
			Indicate whether the neticut was living on decased at the date
13	Vital Status	□ Living	Indicate whether the patient was living or deceased at the date of last contact.
15	(at date of last contact)	□ Deceased	5
			If the patient is living, provide the date of last contact with the
		/ /	patient (as reported by the patient, medical provider, family
14	Date of Last Contact	$(month)^* (day) (year)^*$	member, or caregiver).
			<u>2897020</u> (month), <u>2897022</u> (day), <u>2897024</u> (year)
			Provide the number of days from the date the patient was
	Normhan a GDarra Grann		initially diagnosed pathologically with the disease described
	Number of Days from		on this form to the date of last contact.
15	Date of Initial		<u>3008273</u>
	Pathologic Diagnosis to		Only provide Interval data if you have received permission from
	Date of Last Contact		the NCI to provide time intervals as a substitute for requested
			dates on this form.
			If the patient is deceased, provide the date of death. 2897026, (month) 2897028 (day), 2897030 (year)
16	Date of Death	$(month)^* (day) (year)^*$	$\frac{2697020}{2697020}$, (11011(1) $\frac{2697020}{2697020}$ (uay), $\frac{2697030}{2697030}$ (year)
		(month)* (day) (year)*	
			Provide the number of days from the date the patient was
	Number of Days from		initially diagnosed pathologically with the disease described
	Date of Initial		on this form to the date of death.
17	Pathologic Diagnosis to		<u>3165475</u>
	Date of Death		Only provide Interval data if you have received permission from
	Date of Death		the NCI to provide time intervals as a substitute for requested
			dates on this form.
		Progressive Disease	Provide the patient's response to their initial first course treatment.
	Measure of success of	□ Stable Disease	<u>2786727</u>
10	outcome <u>at the</u>	Partial Response	
18	<u>completion of initial</u>	Complete Response	
	first course treatment	□ Not Applicable (Treatment Ongoing)	
			Indicate the notion to measure of success at the time
	What was the measure	Progressive Disease	Indicate the patient's measure of success at the time this follow-up form is completed.
10	of success at Date of	Stable Disease	<u>3033278</u>
19	Last Contact provided	Partial Response	0000270
	on this form?	Complete Response	
	,	🗖 Unknown	

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	Tumor Event Information Complete this section if the patient had a new tumor event. If the patient did not have a new tumor event (or if the TSS does not know) indicate this in the question below, and the remainder of this section can be skipped. ote: The New Tumor Event section on OpenClinica can be completed multiple times, if the patient had multiple New Tumor Events.			
#	Data Element	Entry Alternatives	Working Instructions	
20	New Tumor Event After Initial Treatment?	□ Yes □ No □ Unknown	Indicate whether the patient had a new tumor event (e.g. metastatic, recurrent, or new primary tumor) after initial treatment. 3121376 If the patient did not have a new tumor event or if this is unknown, the remaining questions can be skipped.	
<u>21</u>	Type of New Tumor Event	 Locoregional Recurrence Distant Metastasis New Primary Tumor 	Indicate whether the patient's new tumor event was a locoregional recurrence, a distant metastasis or a new primary tumor. A new primary tumor is a tumor with a different histology as the tumor submitted to TCGA. 3119721	
<u>22</u>	Site of New Tumor Event	LungBrainBoneUnknownLiverOther, specify	Indicate the site of this new tumor event. 3108271	
<u>23</u>	Other Site of New Tumor Event		If the site of the new tumor event is not included in the provided list, describe the site of this new tumor event. 3128033	
<u>24</u>	Date of New Tumor Event	/// (month)* (day) (year)*	If the patient had a new tumor event, provide the date of diagnosis for this new tumor event. 3104044 (month), 3104042 (day), 3104046 (year)	
<u>25</u>	Number of Days from Date of Initial Pathologic Diagnosis to Date of New Tumor Event After Initial Treatment		Provide the number of days from the date the patient was initially diagnosed pathologically with the disease to the date of new tumor event after initial treatment. <u>3392464</u> Only provide Interval data if you have received permission from the NCI to provide time intervals as a substitute for requested dates on this form.	
<u>26</u>	Additional Surgery for New Tumor Event:	☐ Yes ☐ No ☐ Unknown	Using the patient's medical records, indicate whether the patient had surgery for the new tumor event in question. <u>3427611</u>	
<u>27</u>	Month of Additional Surgery for New Tumor Event	(month)* (day) (year)*	If the patient had surgery for the new tumor event, provide the month this surgery was performed. <u>3427612 (month), 3427613 (day), 3427614 (year)</u>	
<u>28</u>	Number of Days from Date of Initial Pathologic Diagnosis to Date of Additional Surgery for New Tumor Event		Provide the number of days from the date the patient was initially diagnosed pathologically with the disease described on this form to the date of additional surgery for new tumor event (loco-regional). <u>3008335</u> Only provide Interval data if you have received permission from the NCI to provide time intervals as a substitute for requested dates on this form.	
<u>29</u>	Procedure Type for New Tumor Event	 Excisional Biopsy Incisional Biopsy Surgical Resection Unknown Other Method, Specify Below 	If the patient had surgery for the new tumor event, provide the type of procedure performed for this tumor. 3125097	
<u>30</u>	Other Procedure Type for New Tumor Event		If the procedure for the new tumor event was not included in the list provided, indicate the type of procedure performed. <u>3125102</u>	
<u>31</u>	Residual Tumor After surgery for New Tumor Event	 RX: The presence of residual tumor or margin status cannot be assessed. R0: No residual tumor and negative microscopic margins in resected specimen. R1: Microscopic residual tumor. No gross residual disease but positive microscopic margins. 	Using the patient's pathology/laboratory report, select the residual tumor status after the surgical resection for the new tumor event. 3104081	

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#	Data Element	Entry Alternatives	Working Instructions
		 R2: Macroscopic residual tumor. Grossly visible residual disease. Unknown 	
<u>32</u>	Additional treatment for New Tumor Event: <i>Radiation Therapy</i>	☐ Yes □ No □ Unknown	Indicate whether the patient received radiation treatment for this new tumor event. <u>3427615</u>
<u>33</u>	Additional treatment for New Tumor Event: <i>Pharmaceutical Therapy</i>	☐ Yes □ No □ Unknown	Indicate whether the patient received pharmaceutical treatment for this new tumor event. <u>3427616</u>

Principal Investigator or Designee Signature

Print Name

Date