Initial Case Quality Control Form

Testicular

<u>Instructions:</u> This form should be completed for all cases submitted for TCGA, prior to the shipment of samples to the BCR.

Questions regarding this form should be directed to the Tissue Source Site's primary Clinical Outreach Contact at the BCR.

Tissue Source Site (TSS) acknowledges that the Biospecimen Core Resource (BCR) may confirm that the diagnosis of the frozen biospecimen is consistent with the primary diagnosis reported by the TSS through histopathology examination in the BCR laboratory. If the BCR identifies a possible discrepancy, the TSS authorizes the BCR to report these patient results to the TSS by means of a formal report in confidential email format for the quality assurance program of the TSS to address.

Γissue	Source Site (TSS):	TSS ID: TSS Unique Patient ID: Interview	ver Name:	Interview Date / /
#	Question	Entry Alternatives		Working Instructions
Tum	or Information			
		Histologic Diagnosis	Percent	Indicate the confirmed histologic diagnosis of the tumor
		□ Seminoma	%	submitted for TCGA. 3081934 (Histologic Diagnosis), 3729998 (Percentage)
	Histologic Diagnosis of	☐ Non-Seminoma - Choriocarcinoma	%	The listed histologies are the only histologic types being accepted
1*	Frozen Tumor Submitted for	☐ Non-Seminoma - Embryonal carcinoma	%	for this TCGA study. Recurrent tumors are NOT accepted.
1"	TCGA	□ Non-Seminoma – Yolk Sac Tumor	%	
	Check all that apply	☐ Non-Seminoma – Teratoma (Mature)	%	Note: Spermatocytic seminoma cases are excluded from this study.
		☐ Non-Seminoma – Teratoma (Immature)	%	
		Total	100%	1
2*	Tumor Presentation	☐ Primary (primary untreated malignant biospecimen)		Indicate the type of tumor submitted for TCGA. 3288124 This is a biospecimen that has not been treated with chemotherapy or radiation prior to resection. If a metastatic tumor is being submitted for a triplet case, please complete the Metastatic CQCF.
3*	Anatomic Site of Malignant Specimen	□ Testis		Indicate the anatomic site of the frozen tumor biospecimen submitted for TCGA. 4132152
4*	Tumor Laterality	□ Right □ Left		Indicate the laterality if the frozen tumor biospecimen submitted for TCGA was located in a paired site. 827
5*	Date of Cancer Sample Procurement	Month Day	 Year	Provide the date of the procedure performed to obtain the malignant tissue submitted for TCGA. 3008197 (month), 3008195 (day), 3008199 (year)
6*	Method of Cancer Sample Procurement	□ Orchiectomy		Indicate the procedure performed to obtain the malignant tissue submitted for TCGA. 3103514
7*	Country Where Cancer Sample was Procured			Provide the country where the tissue submitted for TCGA was procured. 3203072
8*	Race	 □ American Indian or Alaska Native A person having origins in any of the original peoples of North and South America), and who maintains tribal affiliation or community attachment. □ Asian A person having origins in any of the original peoples of the far East, Sout subcontinent including, for example, Cambodia, China, India, Japan, Koree Philippine Islands, Thailand, and Vietnam. □ White 	heast Asia, or in the Indian	Provide the patient's race using the defined categories. 2192199

#	Question	Entry Alternatives	Working Instructions			
		 A person having origins in any of the original peoples of the far Europe, the Middle East, or North Africa. Black or African American A person having origins in any of any of the black racial groups of Africa. Terms such as "Haitian" or "Negro" can be used in addition to "Black or African American." Native Hawaiian or other Pacific Islander	Provide the patient's ethnicity using the defined categories.			
9	Ethnicity	 Not Hispanic or Latino A person not meeting the definition of Hispanic or Latino. Hispanic or Latino A person of Mexican, Puerto Rican, Cuban, Central or South American or other Spanish culture or origin, regardless of race. Not Evaluated Not provided or available. Unknown Could not be determined or unsure. 	2192217			
10*	Vessel Used	☐ Cryovial ☐ Cassette ☐ Biospecimen Storage Bag ☐ Cryomold ☐ Other, specify	Indicate the type of vessel used to ship the tissue to the Biospecimen Core Resource (BCR) for TCGA. 3081940			
11	Other Vessel Used		If the vessel used to ship the tissue to the BCR is not included in the provided list, specify the vessel used. 3288137			
12*	Is tumor sample being submitted for macrodissection?	□ Yes □ No	Indicate whether the tumor sample submitted to the BCR is intended to undergo macrodissection after the BCR receives the sample. 3521908			
13*	Was sample prescreened at site?	□ Yes	Indicate whether the sample submitted to the BCR was prescreened at the TSS. 3081942			
Tumo	or Slides Submitted					
<u>12</u>	Types of Slides Submitted Check all that apply	□ Physical Top Slide □ Physical FFPE Slide □ Digital Top Slide Image	Indicate the type(s) of slide(s) submitted to the BCR. 3521909 Top Slide Definition: Slide cut directly from frozen biospecimen = mirror image of inked surface			
<u>13</u>	Slide/Digital Image ID #		Provide the slide ID for each slide (physical and digital image) submitted to the BCR. 2321277			
Tumo	Tumor Sample Information If the TSS is submitting multiple pieces of the same primary tumor for this case; complete the following information for each piece of tumor sent to the BCR.					
14	Tumor Identifier		Provide the TSS unique tumor ID. If multiple pieces of tumor are submitted, each tumor needs a unique ID. 3288096			
<u>15</u>	Weight of Frozen Tumor	(mg)	Provide the weight of the tumor sample submitted for TCGA. 3081946 Weight can be estimated based on the size of the tumor submitted.			
<u>16</u>	Tumor Nuclei %	(%)	Provide the percent of tumor nuclei for the sample submitted for TCGA. 2841225 Check with the BCR to confirm the current acceptable TCGA metrics.			

#	Question		Entry Alternatives		Working Instructions	
<u>17</u>	Necrosis %	(%)			Provide the percent of necrosis for the sample submitted for TCGA. 2841237 Check with the BCR to confirm the current acceptable TCGA metrics.	
Norn	nal Information A normal contr	rol must be present to qualify.				
18	Type(s) of Normal Control Check all that apply	☐ Whole Blood ☐ Buffy Coat ☐ Lymphocytes	☐ Extracted DNA from Blood☐ Extracted DNA from Saliva or☐ Non-Neoplastic Control Tissu		Indicate the type of normal control submitted for this case. 3081936	
Norm	nal Control: Whole Blood					
19	Method of Normal Sample Procurement	☐ Blood Draw			Indicate the procedure performed to obtain the normal control sample submitted for TCGA. 3288147	
20	Date of Normal Sample Procurement	 Month		Year	Provide the date of the procedure performed to obtain the normal control submitted for TCGA. 3288195 (month), 3288196 (day), 3288197 (year)	
<u>21</u>	Normal Identifier				Provide the TSS unique normal ID. If multiple normal control samples are submitted, each normal control needs a unique ID. 3288138	
Norm	nal Control: Buffy Coat/ Lymph	ocytes				
22	Method of Normal Sample Procurement	☐ Blood Draw			Indicate the procedure performed to obtain the normal control sample submitted for TCGA. 3288147	
23	Date of Normal Sample Procurement	 Month		Year	Provide the date of the procedure performed to obtain the normal control submitted for TCGA. 3288195 (month), 3288196 (day), 3288197 (year)	
	Normal Control Type	☐ Buffy Coat☐ Lymphocytes	•		Indicate the type of normal control submitted for TCGA. 3081936	
24	Normal Identifier				Provide the TSS unique normal ID. If multiple normal control samples are submitted, each normal control needs a unique ID. 3288138	
Normal Control: Extracted DNA from Blood or Saliva						
25	Method of Normal Sample Procurement	☐ Blood Draw ☐ Oragene ☐ Other, specify			Indicate the procedure performed to obtain the normal control sample submitted for TCGA. 3288147	
26	Other Method of Normal Sample Procurement				If the procedure performed to obtain the normal sample is not included in the provided list, specify the method used. 3288151	
27	Date of Normal Sample Procurement	 Month		Year	Provide the date of the procedure performed to obtain the normal control submitted for TCGA. - 3288195 (month), 3288196 (day), 3288197 (year)	

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#	Question	Entry Alternatives	Working Instructions	
<u>28</u>	Normal Identifier		Provide the TSS unique normal ID. If multiple normal control samples are submitted, each normal control needs a unique ID. 3288138	
<u>29</u>	Extracted DNA Quantity	(μg)	Provide the quantity (µg) of the normal control sample sent to the BCR for TCGA. $\underline{3288185}$	
<u>30</u>	Extracted DNA Quantification Method		Provide the quantification method of the normal control sample sent to the BCR for TCGA. 3288186	
<u>31</u>	Extracted DNA Concentration	(μg/μL)	Provide the concentration (µg/ µL) of the normal control sample sent to the BCR for TCGA. $$\underline{3288187}$	
<u>32</u>	Extracted DNA Volume	(μL)	Provide the volume (μL) of the normal control sample sent to the BCR for TCGA. 3288188	
Norm	al Control: Non-Neoplastic Co	ntrol Tissue		
33	Method of Normal Sample Procurement	□ Orchiectomy	Indicate the procedure performed to obtain the normal control sample submitted for TCGA. 3288147	
34	Date of Normal Sample Procurement	Month Day Year	Provide the date of the procedure performed to obtain the normal control submitted for TCGA. 3288195 (month), 3288196 (day), 3288197 (year)	
<u>35</u>	Normal Identifier		Provide the TSS unique normal ID. If multiple normal control samples are submitted, each normal control needs a unique ID. 3288138	
<u>36</u>	Anatomic Site of Non- Neoplastic Control Tissue	□ Epididymis □ Spermatic Cord	If the normal control type is normal tissue, indicate the anatomic site of the non-neoplastic control tissue submitted for TCGA. 3081938	
	Proximity of Normal Tissue to Tumor	☐ Distal (> 2cm) from the primary tumor	If the normal control type is normal tissue, confirm that the submitted normal tissue was at least 2cm away from the primary tumor. 3088708 Adjacent (< 2cm) Normal Tissue is not accepted for this tissue type. Unknown Normal Tissue is not acceptable for this tissue type.	
<u>37</u>	Normal Slide ID#		If the normal control type is normal tissue, provide the slide ID for the physical top slide OR the digital slide image of the normal control being sent to the BCR. 3288217	
Verification: By providing the below information, the Principal Investigator acknowledges that the information provided by the institution is true and correct and has been quality controlled.				

#	Question	Entry Alternatives		Working Instructions			
Tissue throug	Pathology Review Tissue Source Site (TSS) acknowledges that the Biospecimen Core Resource (BCR) may confirm that the diagnosis of the frozen biospecimen is consistent with the primary diagnosis reported by the TSS through histopathology examination in the BCR laboratory. If the BCR identifies a possible discrepancy, the TSS authorizes the BCR to report these patient results to the TSS by means of a formal report in confidential email format for the quality assurance program of the TSS to address.						
38	Name of Pathologist			Provide the name of the Pathologist that provided the information for all previous sections. 3288225			
39	Date of Pathologist Review	Month Day	Year	Provide the date of the pathology review performed by the TSS pathologist above. 3462941 (month), 3462917 (day), 3462960 (year)			
Princ	ipal Investigator/Authorized	Designee Confirmation					
40	Percent Tumor Nuclei meets TCGA metrics?	□ Yes □ No		Confirm that the malignant sample submitted to the BCR meets the current tumor nuclei metrics for TCGA. 3288520 Check with the BCR to confirm the current acceptable TCGA metrics. If submitting for macrodissection, please contact the BCR prior to shipment.			
41	Percent Necrosis meets TCGA metrics?	□ Yes □ No		Confirm that the malignant sample submitted to the BCR meets the current necrosis metrics for TCGA. 3288524 Check with the BCR to confirm the current acceptable TCGA metrics. If submitting for macrodissection, please contact the BCR prior to shipment.			
42	De-Identified Pathology Report Submitted?	□ Yes □ No		Confirm that a de-identified pathology report will be sent to BCR prior to or with the shipment of the physical samples. 3288292			
43	Is the histologic diagnosis on the CQCF (as determined by the TSS pathology review of the TCGA frozen section top slide) consistent with the histology listed in the final diagnosis on the pathology report?	□ Yes □ No		Confirm that the diagnosis provided on this CQCF for the tumor sample being submitted to TCGA is consistent with the diagnosis found on the patient's pathology report for the tumor being sent to the BCR. 3288300 If "yes," skip related question below. The diagnosis is considered to be consistent if at least one of the following criteria are met: 1) Diagnosis on the CQCF is identical to the pathology report for the tumor being sent to the BCR. 2) Diagnosis on the CQCF includes as least one of the subtypes listed on the pathology report and all subtypes on the pathology report are acceptable for TCGA. 3) Diagnosis on the CQCF is "histology, NOS" (i.e., Adenocarcinoma, NOS) and the pathology report lists a specific subtype within the same histologic group 4) Diagnosis on the CQCF indicates "Mixed Subtype" and the pathology report lists two or more acceptable subtypes, provided that percent subtype(s) meet applicable TCGA disease-specific requirements.			

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#	Question	Entry Alternatives	Working Instructions
44	If the diagnosis on this form is not consistent with the provided pathology report, indicate the reason for the inconsistency.	 □ Macrodissection performed at TSS to select for a region containing an acceptable TCGA diagnosis (see note at right) □ Pathology analysis at TSS determined a specific histological subtype different from original pathology report (see note at right) □ Pathology review of frozen section for TCGA determined histological subtype different from the pathology report (see note at right) 	If the diagnosis provided on this form is not consistent with the diagnosis found on the pathology report provided, specify a reason for this inconsistency. 3288315 If a TSS pathology review of the TCGA committed sample resulted in a different histological subtype than what is documented on the original pathology report, an amendment to the pathology report should be submitted when the sample is shipped to the BCR; or in the absence of an amended pathology report, the TSS must complete and submit an electronic copy of the "TCGA Pathologic Diagnosis Discrepancy Form". In the case of diagnosis modifications, institution protocol should be followed for proper quality assurance.
45	History of Testicular Cancer	☐ Yes, history of metachronous contralateral testicular cancer ☐ Yes, history of synchronous contralateral testicular cancer ☐ No, primary tumor submitted to TCGA is the only TGCT diagnosis to date ☐ Unknown	If the patient does have a history of prior/synchronous malignancy, specify whether they have a history of testicular cancer. 3729780
46	History of Other Malignancy (Not including prior diagnoses of Testicular Cancer)	□ None □ History of Prior Malignancy □ History of Synchronous Malignancy □ Both History of Synchronous and Prior Malignancy	Indicate whether the patient has a history of malignancies. If the patient has any history, including synchronous or bilateral malignancies, please complete an "Other Malignancy Form" for each malignancy diagnosed prior to the procurement of the tissue submitted for TCGA. 3382736 If the patient has a history of multiple diagnoses of basal or squamous cell skin cancer, complete an OMF for the first diagnosis for each of these types.
47	History of Neoadjuvant Treatment for Tumor Submitted for TCGA	□ None □ Radiation prior to sample procurement* □ Pharmaceutical treatment prior to sample procurement* □ Both pharmaceutical treatment and radiation prior to sample procurement*	Indicate whether the patient received therapy for this cancer prior to the sample procurement of <i>the tumor submitted for TCGA</i> . If the patient did receive treatment for this cancer prior to procurement, the TSS should contact the BCR for further instruction. 3382737 *Systemic therapy and certain localized therapies (those administered to the same site as the TCGA submitted tissue) given prior to the procurement of the sample submitted for TCGA are exclusionary.
48	Consent Status	☐ Formally Consented ☐ Exemption 4* ☐ Consented by Death ☐ Waiver*	Indicate whether the patient was formally consented, consented by death, or if the case has an exemption or waiver for consent. If the submitting institution's IRB has approved consent for TCGA, consent requirements have been met. 3288361 *Exemptions and waivers for consent must be approved by NCI.
Date	of Consent		
49	Date of Consent	Month Day Year	If the patient was formally consented, provide the date of consent. 3081955 (Month), 3081957 (Day), 3081959 (Year)

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#	Question		Entry Altern	natives	Working Instructions	
Date	Date of Death Do not complete date of death, if patient formally consented.					
52	Date of Death				If the patient consented by death, provide the date of death. 2897026 (Month), 2897028 (Day), 2897030 (Year)	
		Month	Day	Year		

		,
		//
Principal Investigator or Designee Signature	Print Name	Date

I acknowledge that the above information provided by my institution is true and correct and has been quality controlled.