

Sponsor Name	Protocol Number
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A. PATIENT INFORMATION (REQUIRED)			
Patient ID		Second Patient ID	
DOB (MM/DD/YYYY)	Race / Ethnicity	Sex	M      F

B. ORDERING PHYSICIAN INFORMATION (REQUIRED)		
Site / Institution Name		Site Number
Research Partnership		Street Address, Unit
City	Postal Code	Country
Phone		Fax
Principal Investigator/ Ordering Physician		Email Address (required for report delivery)

C. TESTING OPTIONS		Optional add-on tests:	MMR IHC	PD-L1 IHC <sup>1</sup>	HRD	Tumor Origin	DPYD	
xT Solid Tumor + Normal* – 648 genes ↳ Add Concurrent xF Liquid Biopsy* – 105 genes	(Uses normal match blood sample)	Conversion to xF Liquid Biopsy 105 genes – If concurrent testing is not selected, you can opt-in to one of the following: Convert to xF <a href="#">immediately</a> Convert to xF <a href="#">after additional tissue request</a>						
	xT Solid Tumor Only* – 648 genes							
	xT Hematologic Malignancy* – 648 genes	(FFPE)						
	xF Liquid Biopsy* – 105 genes	(Non-hematologic malignancies only)	<sup>1</sup> PD-L1 clone 22c3 is the default. For different clones, please select all that apply: 22c3                      28-8                      SP142					
xE Whole Exome Tumor + Normal* – 19,433 genes	(Requires normal match sample)							

\*For cancers determined to be ovarian, breast, prostate or pancreatic (at pathology review), this includes an order for a separate BRCA1/2 – Tumor Analysis.

D. SPECIMEN RETRIEVAL					
xT or xE Solid Tumor	Option 1 – Specific specimen requested (Please provide specimen details below).		Option 2 – Let the submitting pathologist choose specimen.		Option 3 – Biopsy to be scheduled for:
	Pathology Lab Name				
	Case Number	Block #	Solid Tumor Collection Date	Check here if the pathology lab is <u>not</u> part of the treatment team.	
xT or xE Normal	Blood    Saliva	Date of Collection	Section A must be completed for these options.	Send saliva kit to patient <b>for xT Normal only</b>	Previously Submitted
xF Liquid Biopsy	Blood				
xT Hematologic Malignancy	FFPE (Bone Marrow Biopsy, Bone Marrow Clot, Lymph Node, or other involved tissue)	Date of Collection	Section A must be completed for these options. <b>Please see specimen instructions for details.</b>	Send saliva kit to patient	

E. CURRENT DIAGNOSIS						
NSCLC	Melanoma	Prostate	Colorectal Carcinoma	Ovarian	Breast	Other:
Disease Status (select all that apply):			Metastatic	Refractory	Relapse	Other:
ICD-10 Primary Diagnosis Code(s)		Additional Details			Stage	

**SIGNATURE** I certify that I have explained to the patient the purpose, risks and benefits of the test(s) being ordered. My signature below is my certification of medical necessity for the test and further certifies that I am authorized to order the test(s) and have obtained from the patient informed consent that meets the requirements of applicable law for Tempus to: (a) perform the test described in this form; (b) obtain, receive, and release, test results and any corresponding medical information as necessary for reimbursement or the processing of insurance claims; and (c) collect, use, and retain samples and information obtained from the patient, including the test results, for an indefinite period of time, including de-identifying such information and disclosing the de-identified information for other purposes (as described on the back of this form).

G. ORDERING INVESTIGATOR/HEALTHCARE PROVIDER SIGNATURE	
Ordering Physician's Signature	Date (MM/DD/YYYY)
Printed Name	

H. FORM COMPLETED BY	
Name	Date (MM/DD/YYYY)
Email	