## "I"EMPUS International Requisition Form (De-Identified) — 2024.04.30 Phone: +1800.739.4137 | Fax: +1800.893.0276 | support@tempus.com

	ation is incomplete or missing, testir		33.00111							
A.PATIENT INFORMATION										
Primary Patient ID			Second	Secondary Patient ID						
DOB (DD/MM/YYYY) Biologic		Biological Sex	Race/Ethnicity							
B.ORDERING PHYSICIAN INFORMATION										
Distributor	agal nama)				Phone					
			g Physician (full le							
Facility Name		Tempus	Account #	or report delivery)		Fax				
Facility Address (Street, Unit)				Postal Code		Country				
Additional person to be copied										
Name			ах	Facility Name						
C.TESTING OPTIONS										
Common test combinations Test descriptions				Specimen required		Optional ac	tional add-on tests (select all that apply):			
xT (DNA) & xR (RNA): Solid Tur	xT (DNA) & xR (RNA): Solid Tumor/Normal XT: 648-gene DN xR: whole transc		th normal match;		FFPE Tissue; Normal: Blood or Saliva		xT Solid Tu	mor, xR LDT	xT Normal	
Add xF liquid bionsu at time of order		scriptome KIVA sequen	enig test.		Normal. Blood of Salive	•	FFPE Tissu	e	Blood or Saliva	
Add xF liquid biopsy at time of order, based on the following:  1 believe it is medically necessary to order a liquid biopsy test concurrently with a solid tumor tissue test because of one or more of the following reasons: (a) guidelines support the use of testing in this disease state; (b) turnaround time for tissue result may delay a treatment decision for my patient; (c) the tissue is at risk to fail (e.g. small tissue, archived tissue) and I may not have a timely result to make  22C3 DEFAULT  UGT1A1									DPYD UGT1A1	
a treatment decision for my patient; (d) genomic heterogeneity may cause the patient's available tissue to not be completely representative, and I want to make sure I have a complete mutation profile.								OOTIA		
If I have not already ordered an xF test above, I opt to convert my xT solid tumor order to an xF liquid biopsy test if necessary:  By converting immediately OR After an additional tissue request is attempted								SP142 SP263 MMR IHC		
xT (DNA) & xR (RNA): Solid Tumor xT: 648-gene DNA sequencing test			R: whole transcriptom	ne RNA sequencing test.	FFPE Tissue			+ FISH <sup>1,2</sup>		
xE (DNA) & xR (RNA): Solid Tumor/Normal XE: over 19,000-gene whole ex RI: whole transcriptome RNA				ith normal match;	FFPE Tissue;			C FDA¹		
Individual test options		Normal: Blood or Saliva			igin (RNA)					
xR (RNA Only): Solid Tumor		FFPE Tissue 1 Powered by NeoGenomics. 2 For more information about reflex to FISH, please see								
xT (DNA Only): Solid Tumor 648-gene DNA sequencing test.				FFPE Tissue			Tempus' Reference Lab Logistics Overview at Tempus.com. 3 Normal sample is required for ovarian or breast cancers			
xF (Liquid Biopsy):			23-gene liquid biopsy test for solid tumors.			Blood (Streck)				
xE (DNA Only): Solid Tumor/Normal Over 19,000-gene whole ex			sequencing test with	FFPE Tissue; Normal: Blood or Saliva						
D.SPECIMEN RETRIEVAL See Tempus' specimen guidelines for collection instructions and further details.										
FFPE Tissue										
Option 1:	Option 2:	Option 3:		Pathology Lab (Name, Ci	ty)					
Specific specimen requested	Let the submitting pathologist choose specimen									
	choose specimen	Biopsy to be sche		Case Number	Block#		Date of Collection		on	
Blood				Saliva						
Sample previously submitted Date of Collection:				Send saliva kit to patient Date of Collection:						
E.CURRENT DIAGNOSIS										
Breast NSCLC	Pancreatic Other:		Primary ICD-10	Codes (C & D codes only)	9	Stage I	III	Other:		
Colorectal Ovarian	Prostate					II	IV			
Disease Status (select all that apply):			Has the patient had any type of transplant?							
Metastatic Relapse Other:  Refractory Recurrent			No Yes —Type:							
F.PHYSICIAN SIGNATURE AND CONSENT										
I certify that the patient has received an explanation of the purpose, risks, and benefits of the ordered test(s). My signature below certifies medical necessity of the test(s) (including that the test results will inform the treatment plan) and further certifies that I am authorized to order the test(s) and that the patient has provided informed consent that meets the requirements of applicable law for Tempus or its reference lab to: (a) collect and use the patient's samples (including genetic material) and health information										
law for Tempus or its reference lab to: (a) co and perform the ordered test(s); (b) obtain reimbursement or the processing of insura	Printed Name (full legal name) Today's Date (DD/MM/YY				D/MM/YYYY)					