BIOSPECIMEN PRE-ANALYTICAL VARIABLES (BPV) REQUEST FOR BIOSPECIMENS

Please complete this request form to access BPV biospecimens and submit the completed form to **bpv@vai.org**. The form will be reviewed and approved by the joint National Cancer Institute/Van Andel Research Institute access committee according to the criteria for approval (see page 4) prior to release for distribution.

Principal investigator informatio	n					
First name:	Middle initial:	Last name:				
Salutation:	Degree:	Title:				
Mailing address						
Institution:						
Department:						
Address 1:						
Address 2:						
City:	State:	Zip code:	Country:			
Phone no.:	Alt. phone no.:	Fax no.:	Email:			
Study information						
Study title:						
-	Are you currently funded for this research?					
	s this request for an NIH grant application?					
Funding source(s):						
If not, what is your plan and timel	ine for obtaining funding?					
How soon will you use samples, once received?						
Study objective (including specific aims):						
Brief justification:						
Describe how you plan to share the data and resources generated:						
Considerable						
Case details						
Case demographics						
Gender: Male Female Any						
Race:						
Ethnicity: Not Hispanic or La	_ ·	— ·				
Age range: Minimum:						
Case history (please list any inclusion and exclusion requirements):						

Biospecimen information						
Please check the appropriate tissue type below along with the desired sample preparation.						
Tissue type (check all that apply)						
Anatomic site	Tumor-primary	Normal adjacent	Tissue microarray	Case matched blood		
☐ Colon						
☐ Kidney						
Lung						
Ovary						
☐ Blood (not case matched)						
Total number of cases requested:						
Module requested:						
☐ Module I (time	in fixative)	le II (delay to fixation)	☐ Module V (freezi	ng/storage methods)		
		Module I (time in fixative)				
	Experimental protocol	Delay to fixation	Time in fixative			
	A		6 hours			
	В С	<1 hour	12 hours			
	C		23 hours			
D 72 hours Module II (delay to fixative)						
	Experimental protocol	Delay to fixation	Time in fixative			
	E	1 hour	12 hours			
	F	2 hours	11 hours			
	G		10 hours			
	H	12 hours	12 hours			
Module V (snap freezing method or storage method)						
	U	Dry ice	-80°C			
V W		Dry ice	-80°C			
	X	LN ₂	LN ₂			
Do you require access to digital images (check all that apply)? H&E IHC* *If IHC, check antigens of interest below:						
IHC antigens of interest						
	Colon	Kidney	Ovarian			
	☐ EGFR		P53			
☐ CDX-2			CK7			
β-catenin		☐ CD10 ☐	WT-1			
☐ Cytok			CA-125			
	☐ MSH6	☐ MUC-1 ☐	MUC-1			
	☐ P16 (CDKN2a		PAX-8			
			ER (6F11)			

Sample preparation details—frozen tissue
Frozen tissue prep type: Required OCT acceptable? Yes No Tissue scroll/ribbon? Other:
Tissue weight requested: mg
Slides: H&E slides (# requested per block:) Frozen sections (# requested per block:)
Nucleic acid: DNA RNA Minimum quantity: μg Maximum quantity: μg Minimum concentration: ng/μL Maximum concentration: ng/μL Minimum volume: μL
Sample preparation details—formalin tissue
Formalin tissue prep type: Required Scroll/ribbon DNA/RNA (see below) Other:
Slides:
Nucleic acid: DNA RNA Maximum quantity: μg Maximum quantity: μg Minimum concentration: ng/μL Maximum concentration: ng/μL Minimum volume: μL
Sample preparation details—blood
Blood prep type: Required Serum Plasma DNA/RNA (see below) Other:
Volume requested:
Nucleic acid: DNA RNA Minimum quantity: µg Maximum quantity: µg Minimum concentration: ng/µL Maximum concentration: ng/µL Minimum volume: µL
Other preparation details
Please describe any other details. Include specifics, as appropriate:
Are there any other considerations of which we should be aware?

CRITERIA USED FOR REQUEST APPROVAL

- For the tissues and blood products that were part of experimental protocols: The potential for the project to address an important question/problem or a critical barrier to progress in the field of biospecimen science. Requests to use the specimens for projects in other fields of interest (e.g., biomarker assay development) will be considered on a case by case basis. Blood and blood products for which there was not a matching experimental protocol will be broadly available for cancer research.
- The level of experience and expertise of the investigators to conduct the proposed analyses.
- The degree to which the requested samples are uniquely suited for the proposed study or whether other samples are equally appropriate.
- The degree to which the quantity of sample requested matches the intended use and the impact on remaining amounts.
- The degree to which the proposed study increases the overall value of the BPV resource.
- Whether the proposed molecular study has already been performed by the BPV Program and if so, would the proposed study be redundant, complementary or synergistic.
- Whether there are sufficient funds to support the proposed research.

Please submit your complete application and any questions via email to **bpv@vai.org**.