OMB No.: 0925-XXXX Expiration Date: XX/XX/XXXX.

Public reporting burden for this collection of information is estimated to average 10 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. **An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.** Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: NIH, Project Clearance Branch, 6705 Rockledge Drive, MSC 7974, Bethesda, MD 20892-7974, ATTN: PRA (0925-XXXX).

Survey to assess the feasibility of establishing a gynecologic specimen bank for research

 Would you consider your practice to be primarily:

- o academic (primary activity within medical school teaching hospital or research institute)
- o private laboratory (not at the hospital)
- o private hospital (not affiliated with a medical center or research institute)
- O laboratory affiliated with managed health organization
- What is the approximate annual surgical pathology specimen volume at the laboratory where you practice (if multiple, give largest):
 - o <10K
 - o 10-25K
 - o >25-50K
 - o >50K
- What proportion of these specimens are gynecologic?
 - o <10%
 - o 10-20%
 - o >20-50%
 - o >50-100%
- Does your laboratory have a subspecialty sign-out with a designated gynecologic section?
 - o No
 - o Yes
- Does your laboratory receive risk-reducing surgery specimens from women at high-risk for gynecologic disease/cancer? If so, estimate annual number?
 - o No
 - O Yes (annual number: ____)

How are specimens for the following specific indications processed?

Clinical Indication	SEE-Fim				
	(Sectioning and				
	Extensively	Submit fimbria	Submit ovaries	Endometrium	
	Examining of the				
	Fimbria)				
High-Grade Serous Cancer	o No	o No	o No	o No	
Stage I, II, IIIAi	o Yes	o Yes (total)	o Yes (total)	o Yes (total)	
		o Yes (partial)	o Yes (partial)	o Yes (partial)	
Risk-reducing salpingo-	o No	o No	o No	o No	
oophorectomy or	o Yes	o Yes (total)	o Yes (total)	o Yes (total)	
salpingectomy		o Yes (partial)	o Yes (partial)	o Yes (partial)	
Surgery for benign	o No	o No	o No	o No	

indications, first sections	o Yes	O Yes (total)	O Yes (total)	O Yes (total)
reveal equivocal or		o Yes (partial)	o Yes (partial)	o Yes (partial)
definite STIC (serous		,		,
tubal intraepithelial				
carcinoma), epithelial				
atypia in ovary or				
clinically occult cancer				
Surgery for benign	o No	0 No	o No	o No
indications, first sections	o Yes	o Yes (total)	o Yes (total)	o Yes (total)
reviewed are negative		o Yes (partial)	o Yes (partial)	o Yes (partial)

- Does your laboratory stain sections of fimbria for Ki67, p53 or other markers (check all that apply):
 - On every specimen?
 - On every risk-reducing salpingo-oophorectomy or salpingectomy specimen?
 - On selected risk-reducing salpingo-oophorectomy or salpingectomy specimens based on H&E review?
 - On early stage high-grade serous cancer?
 - Markers not evaluated
- Would your laboratory consider providing de-identified blocks and matched pathology reports to a
 national specimen bank organized by the National Cancer Institute (NCI) to provide access to
 researchers throughout the world? Possible specimens would include risk-reducing salpingooophorectomy or salpingectomy, early high-grade serous cancer (HGSC), serous tubal
 intraepithelial carcinoma (STIC), and a minor percentage (10%) of total benign tubes and ovaries.
 - o No
 - o Yes

ourer committee	ts/clarifications			