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

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

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# 1. Would you consider your practice to be primarily:

academic (primary activity within medical school teaching hospital or													
research institute)													
				,									
	_			,			. 15						

- private laboratory (not at hospital)
- private hospital (not affiliated with medical center or research institute)
- laboratory affiliated with managed health organization

# 2. What is the approximate annual surgical pathology

specimen volume at the laboratory where you practice (if multiple, give largest):
○ <10K
○ 10-25K
○ >25-50K
○ >50K
3. What proportion of these specimens are gynecologic?
○ <10%
O 10-20%
>20-50%
>50-100%
4. Does your laboratory have a sub-specialty sign-out with a designated gynecologic section?
○ No
○ Yes
<ul> <li>5. Does your laboratory receive risk-reducing surgery specimens from women at high-risk for gynecologic disease/cancer?</li> <li>No</li> <li>Yes</li> </ul>

5a. If you answered yes above, estimate annual number?

6. How are specimens for the following specific indications (a-d below) processed?

# a. High-Grade Serous Cancer Stage I, II, IIIAi

	No	Yes, partial	Yes, total
SEE-Fim (Sectioning and Extensively Examining of the Fimbria)	0	0	0
submit fimbria	0	0	0
submit ovaries	0	0	0
endometrium (if uterus removed)	0	0	0

# b. Risk-reducing salpingo-oophorectomy or salpingectomy

	No	Yes, partial	Yes, total
SEE-Fim (Sectioning and Extensively Examining of the Fimbria)	0	0	0
submit fimbria	0	0	0
submit ovaries	0	0	0
endometrium (if uterus removed)	0	0	0

c. Surgery for benign indications, first sections reveal equivocal or definite STIC (serous tubal intraepithelial carcinoma), epithelial atypia in ovary or clinically occult cancer

No Yes, Yes, total

	m (Sectioning and Extensively xamining of the Fimbria)	0	0	0		
_	submit fimbria	0	0	0		
	submit ovaries	0	0	0		
endo	metrium (if uterus removed)	0	0	0		
d. Surge	ery for benign indications, f	irst sectio	ons reviev	wed are		
		No	Yes, partial	Yes, total		
	m (Sectioning and Extensively xamining of the Fimbria)	0	0	0		
	submit fimbria	0	0	0		
	submit ovaries	0	0	0		
endo	metrium (if uterus removed)	0	0	0		
	your laboratory stain secti markers (check all that ap		nbria for I	Ki67, p53		
On ev	very 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 ea	arly stage high-grade serous cand	er?				
Marke	ers not evaluated					

8. 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 salpingo-oophorectomy or salpingectomy, early high-grade serous cancer (HGSC), serous tubal intraepithelial carcinoma (STIC), and a minor percentage (10%) of total benign tubes and ovaries.

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O Yes

# 9. Other comments/clarifications?

Your answer

**SUBMIT** 

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