Public reporting burden for this collection of information is estimated to average 5 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 (0925-0624). Do not return the completed form to this address.

Filling out PDF Forms

This PDF form contains "roll-over or double-click " help functionality.

This form allows you to enter data directly onto the screen. After completing the form, you are able to print the document so that you can fax/mail the document.

To fill out a form:

- 1. Select the hand tool. $\langle n \rangle$
- 2. Position the pointer inside a field, and click to type text.
- 3. After entering text or selecting a check box, do one of the following:

- Press tab to accept the form field change and go to the next form field.

- Press Shift+Tab to accept the form field change and go to the previous form field.

- Press Enter (Windows) or Return (Mac OS) to accept the form field change and deselect the current form field.

4. Once completed, print the form.

Cancer Trials Support Unit

Acknowledgment Form (for sites utilizing the NCI CIRB)

Email, Mail or Fax to: Cancer Trials Support Unit (CTSU) ATTN: Coalition of Cancer Cooperative Groups (CCCG) Suite1100 1818 Market Street Philadelphia, PA 19103 FAX: 1-215-569-0206 <u>CTSURegulatory@ctsu.coccg.org</u>

This form is **NOT** an IRB approval document. The form does **NOT** require an IRB approval nor IRB signature, but may be completed by the **local site research personnel**. By signing this form, you are informing the CTSU that each site listed below is currently covered by the NCI CIRB and that the CTSU should enter either a **continuing renewal** or **amendment** approval for the site(s) & protocol identified.

1) Protocol #:	2) Protocol Version Date (Required for Amendments Only):			
	/			
		m m d d	m m d d y y y y	
3) Institution Name (<i>List all institutions covered by IRB approval that will conduct this study. Attach complete list if necessary.</i>)	4) NCI Institution	5 & 5a) OHRP Federalwide Assurance Number		
that will conduct this study. Allach complete list if necessary.)	Code	FWA	FWA Expiration Date	
	eoue	(mm/dd/yyyy)	_	
Ex: University of State	ALXXX	FWA00000123	03/01/2006	
6) Principal Investigator:	7) NCI Investigator #:			
	0) OUDD IDD Desistration Number (for the NCI CIDD)			
8) Approval Type:	9) OHRP IRB Registration Number (for the NCI CIRB): Adult-Late Phase Emphasis Board: IRB00000781 Adult-Early Phase Emphasis Board: IRB00009430 Pediatric Board: IRB00009430			
Amendment Renewal				
10) Comments:	1			
The person signing below certifies that the information provided above is correct.				
Questions #1 through #9 must be completed for this form to be accepted.				
11) Name of Person Signing Form:	12) Site Role:			
		1		
13) Title (if applicable):	14) Phone Number:			
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15) Signature:	16) Date:			
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