


Public reporting burden for this collection of information is estimated to vary from 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 (0925-0753). 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. 
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.

Submit to the CTSU
Regulatory Office via the
Regulatory Submission
Portal: www.ctsu.org.

<p>Cancer Trials Support Unit</p> <p><i>Optional form 1</i></p> <p>Withdraw from Protocol Participation</p>	<p><i>Email, Mail or</i> Cancer Trials Support ATTN: Coalition of Cancer Cooperative Groups (CCCG) Suite 1100 1818 Market Street Philadelphia, PA 19103 FAX: 1-215-569-0206 CTSURegulatory@ctsu.ccccg.org</p>
Institution Name (<i>List all institutions permanently closing this study</i>)	NCI Institution Code : (ALXXX)
Protocol Title: (<i>Shortened version acceptable</i>)	Protocol Number:
Rationale for Study Closure: (<i>Select the answer that best describes the closure</i>)	
1) No subjects were accrued at the institution(s) listed above, and the protocol is now closed to accrual by the sponsoring organization.	<input type="checkbox"/>
2) No subjects were accrued at the institution(s) listed above, and the study has been closed with the local IRB with no anticipation of future accrual (<i>If the protocol is re-opened at a later date the site must submit an initial approval</i>).	<input type="checkbox"/>
3) All subjects accrued at local institutions have completed treatment and follow-up and no further accruals are anticipated at the institution(s) listed above.	<input type="checkbox"/>
4) IRB/Ethics board responsibilities for the protocol listed above are being transferred to another IRB. Review responsibilities at IRB # _____ are being transferred to IRB# _____ (<i>Please note that initial review of this protocol from the new IRB must be submitted to the CTSU. Submission of this form only documents withdraw of the approval from the originating IRB.</i>)	<input type="checkbox"/>
Date of IRB/Ethics Board Action: ___ ___ ___ mm dd yyyy	
<i>The institutional staff signing below certifies that the information provided above is correct.</i>	
Name of Signatory:	Name of approving Organization:
Title of Signatory:	Phone (_____) _____ - _____
Signature:	Date: ____/____/____ m m d d y y y y