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 (0925-0753). 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.

RTOG-0834 CTSU DATA TRANSMITTAL FORM

For Post-Enrollment Data Submissions

- Use this form for post-enrollment data submissions; do not use this form to submit site registration/patient enrollment documents
- Record only one patient and protocol per transmittal form
- Ensure Patient ID and Protocol ID are recorded on each page of each item included
- Ensure pages are in proper sequence (2-sided forms must be copied by site before faxing)
- Do not fax more than 50 pages in one submission
- Changes to data initiated by the site must be reported on the Data Correction Form
- Submit updated data with a new CTSU Data Transmittal Form and new date

·			
Date:	Total # Pages Faxed:		Patient ID#:
(dd-mm-yyyy)	(Including Transmittal)		(EORTC Sequential IDENT. No.)
Site Name:			NCI CTEP Code:
(Institution)			(Internal ID)
Site Address:			INST. No:
Transmittal Completed By:			Phone #:
Email address:			
The item(s) listed below should be faxed to CT			
Item(s) Attached	Number of pages		Visit
Query Form (Query)			
☐ Data Correction Form (DCF)			
☐ Local Pathology / Genetic Testing (Form 2)		☐ Before Random	ization
On Study Form (Form 5)		☐ Before 1 st treatm baseline forms	ent administration (Send his wi h o her
☐ Hematology Form (Form 6)		Baseline, All Arr Within 4 weeks	ns: before randomization
		During Radiothe Week 1, 2, 3, 4,	erapy, Arms 2 & 4: and 5 for TMZ administration
		End of Radiothe Week 6	erapy, Arms 2 & 4:
		After the end of 4 weeks after the	Radiotherapy, All Arms: e end of Radiotherapy
		Adjuvant TMZ, A	Arms 3 & 4: ssments
☐ Biochemistry Form (Form 7)		Baseline, All Arr Within 4 weeks	ms: before randomization
		During Radiothe Week 4	erapy, Arms 2 & 4:
		End of Radiothe Week 6	erapy, Arms 2 & 4:
			Radiotherapy, All Arms: e end of Radiotherapy
		Adjuvant TMZ, A	Arms 3 & 4: ssments

Contact Information: Westat, CTSU Data Operations Center, 1-888-823-5923

Form Version: July 2018

Number	Visit
or pages	Baseline, All Arms: Within 4 weeks before randomization
	During Radiotherapy, All Arms: Week 1, 2, 3, 4 and 5
	End of Radiotherapy: Week 6
	After the end of Radiotherapy, All Arms: 4 weeks after the end of Radiotherapy AND thereafter for every 3 months until disease progression At disease progression
	Adjuvant TMZ, Arms 3 & 4: Additional Assessments
	 Within 4 weeks before randomization 4 Weeks after Radiotherapy Thereafter every 3 months until disease progression or death At disease progression Follow up
	* For patients participating in this component Baseline Thereafter for yearly intervals until tumor progression or death
	☐ At the end of Radiotherapy
	
	Arms 2 & 4 only: at the end of concomitant chemotherapy
	Arms 3 & 4 only: After each cycle of Adjuvant Chemotherapy Cycle:
	4 Weeks after end of Radiotherapy Thereafter every 3 months until disease progression At disease progression
	 End of Protocol Treatment (or in case patient is not randomized) Arms 3 & 4 only At disease progression
	☐ Due every 3 months after disease progression and until patient's death
	Number of pages

For CTSU use only: Short Name shown in (brackets)