

Adverse Events

Page: Adverse Events: Assessment - Cycle 01

Instructions: Cycle length and corresponding instructions will vary per protocol. Final cycle of therapy. Report events existing during this cycle and then recur during subsequent cycles. Report hospitalization events from the previous cycle.

protocol system. Record any observed adverse events not listed on the blank lines at the end. Date is in DD MON YYYY format. Explain any blank dates or fields in the Comments section.

Reporting period start date? Day 1 of this cycle

Reporting period end date? Day 1 of the next cycle. If final cycle, date of the EOT toxicity assessment (outlined in the protocol, generally at least 30 days after last dose of study drug).

Were adverse events assessed during this reporting period? Yes No

If yes, did the patient experience any adverse events during this reporting period? Yes No

Date of most recent adverse event assessment

Comments This should always be at the END of the reporting period. It is the date of the last AE assessment prior to starting the next cycle of Tx. It will almost always be the same date as Day 1 of the next cycle.

If you're not sure, click the Save button. Note that edit checks will still fire.

Save this form, but don't submit to SWOG yet.

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