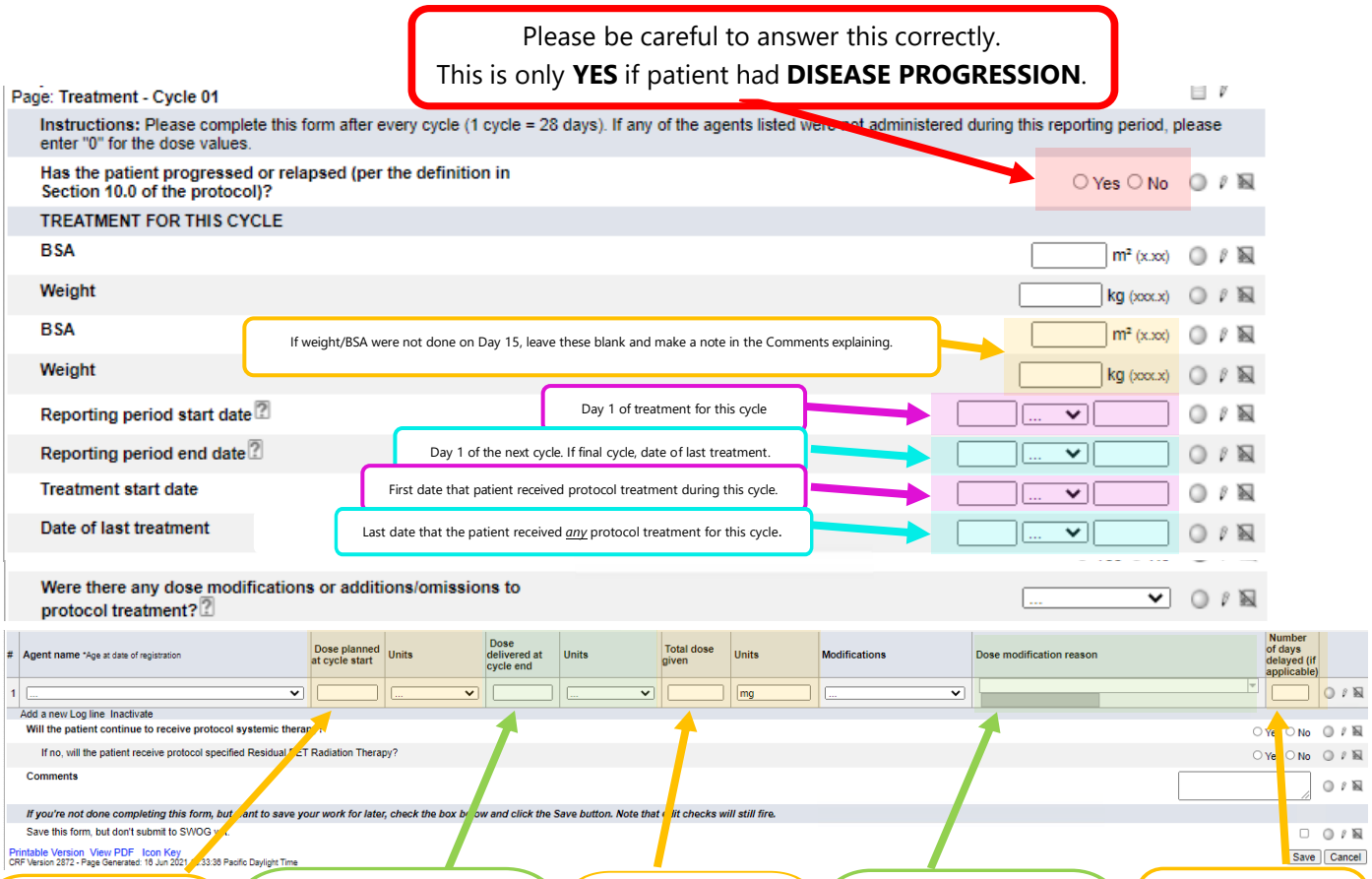
EXAMPLE Treatment Form:

*Actual data entry fields **may vary per study***

Version Date: 4/24/2026



This is the dose planned for the first dose of the cycle. If there were no dose modifications, this will be the protocol specified dose.

This is the dose delivered for the last planned dose. If there were no dose modifications, this will be exactly the same as Dose Planned at Cycle Start. If one of the scheduled doses was not given at all, this will be zero.

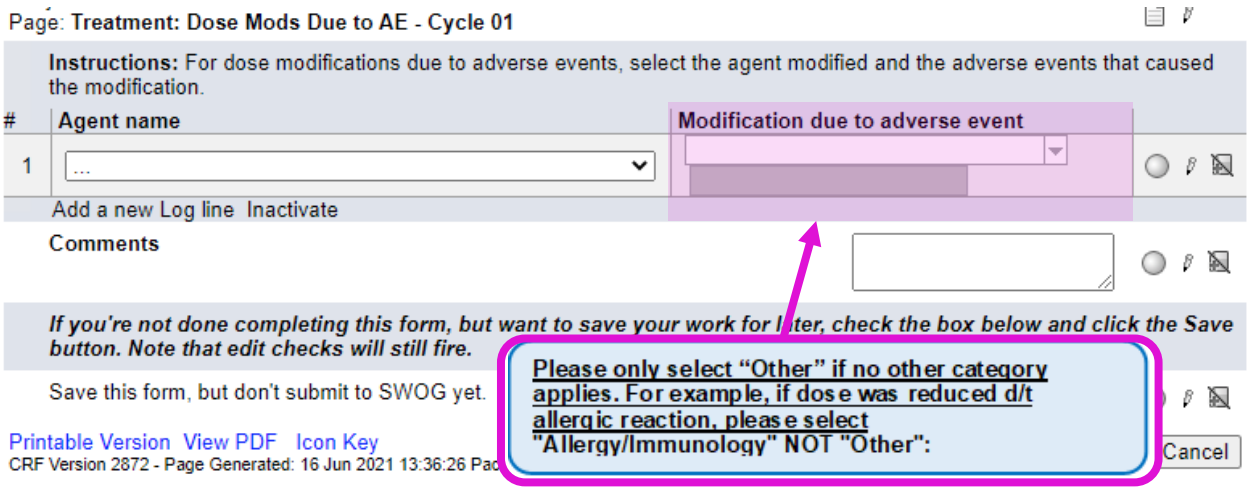
This is the actual dose that the patient received for the entire cycle.

Do not select "Other" unless no other dose mod reason applies. For example, if dose was modified due to an AE, you must select Adverse Event for the Dose modification reason.

If dose was not delayed, leave this BLANK

General Notes:

1. All protocol therapy agents must be reported in every cycle, *unless* the agent was permanently discontinued in a previous cycle.
2. Reporting Period Start Date and Treatment start date should be the same date; the day that ppt first received any treatment in the cycle. Study calendar (Protocol Section 9) footnotes allow for a 48-hour window for labs.
3. Actual body weight of a patient should always be used to calculate body surface area (BSA) for dosing. (See also: Section I.2 of SWOG Policy #38: Research Calculations for Clinical Trials, accessible from: https://www.swog.org/Policy38).
4. All dose modifications must be documented in the treatment table, and in the "Comments" section of the Treatment Form unless otherwise indicated below.
5. Generally, the following do NOT need to be reported as dose modifications (for most protocols):
   a. Doses given within an allowable window (such as: dose delayed for 2 days for administrative reasons and protocol specifies an allowable variance of +/- 3 days. Note: Where a window is NOT specified in the protocol, SWOG defers to the allowable windows specified in the SWOG Best Practices document, accessible from: https://www.swog.org/clinical-trials/protocol-workbench).
   b. Dose adjusted due to ≥ 10% change in weight. (See also: Section I.4 of SWOG Policy #38: Research Calculations for Clinical Trials, accessible from: https://www.swog.org/Policy38).
   c. Dose rounded (unless dose was changed by ≥ 10% from the calculated dose). (See also: Section I.3 of SWOG Policy #38: Research Calculations for Clinical Trials, accessible from: https://www.swog.org/Policy38).


Page: Treatment: Dose Mods Due to AE - Cycle 01
Instructions: For dose modifications due to adverse events, select the agent modified and the adverse events that caused the modification.
#
Agent name
Modification due to adverse event
1
...
Add a new Log line Inactivate
Comments
If you're not done completing this form, but want to save your work for later, check the box below and click the Save button. Note that edit checks will still fire.
Save this form, but don't submit to SWOG yet.
Printable Version View PDF Icon Key
CRF Version 2872 - Page Generated: 16 Jun 2021 13:36:26 Pac
Please only select "Other" if no other category applies. For example, if dose was reduced d/t allergic reaction, please select "Allergy/Immunology" NOT "Other":
Cancel