



## 5.5.1 Create Event Definitions

When you create an Event Definition, you specify its parameters and select the CRFs for it. You do not have to specify CRFs for the Event Definition when you first create it, but can select the CRFs later. To create an Event Definition for the current Study:

1. Access the [Build Study page](#) by selecting **Tasks > Build Study**.
2. In the Actions column of the table, for the **Create Event Definitions** task, click the Add icon

*The Create Study Event Definition page opens.*

**Create Study Event Definition for Docetaxel in Patients With Completely Resected NSCLC**

- A **Scheduled event** is one that is expected to occur for each subject as part of the ordinary progress of the study.
- An **Unscheduled event** is not expected to occur, but may occur as circumstance dictates.
- Scheduled and unscheduled events typically occur at some particular time.
- A **Common event** collects data forms, but is not expected to be associated with a particular time.
- The **Repeating flag** indicates that this type of study event can occur repeatedly within the containing study.
- The **Category attribute** is typically used to indicate the study phase appropriate to this type of study event. Examples might include Screening, PreTreatment, Treatment, and FollowUp.

\* indicates required field.

Name:  \*

Description:

Repeating:  Yes  No

Type:

Category:

3. Complete the dialog box, referring to the field descriptions included in the dialog box. For example:
    - **Name** (of Event Definition): Follow Up Visit
    - **Description**: Each scheduled session to administer the treatment.
    - **Repeating**: Yes (because a Subject will complete the Event multiple times in the Study; each time, the user completes a new set of forms for the Subject for that occurrence of the Event).
    - **Type**: Scheduled (because the Event will be scheduled for specific dates and times). Note that an Event whose Type is **Unscheduled** might or might not occur (like an Adverse Event), while an Event whose Type is **Common** is expected to occur but at an unknown time.
    - **Category**: Treatment.
  4. Click **Continue**.
- The Define Study Event - Select CRF(s) page opens. It lists all CRFs created in your OpenClinica system.*

### Define Study Event - Select CRF(s)

Please select the CRF(s) you would like to make available in this study event.

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CRF Name	Date Created	Owner	Date Updated	Last Updated by	Selected
Adverse Events	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>
Agent Administration	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>
Concomitant Medications	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>
Eligibility	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>
Physical Exam	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>
Verification of Informed Consent	05-Jul-2011	agoodwin	05-Jul-2011	agoodwin	<input type="checkbox"/>

5. In this step, you select the CRFs for the Event Definition. To help locate CRFs to include, click a column header to sort by that column, or use the Find field. You do not have to select CRFs at this point--you can complete the Event Definition without them, then add the CRFs later if you like. If you want to select the CRFs now, click the checkbox in the Selected column for each CRF you want to include in the Event Definition, then click Continue. For example, select the Physical Exam CRF, which will be completed for each Followup Visit Event in the Study. *The Define Study Event - Selected CRF(s) - Select Default Version page opens. It lists configuration options for each CRF you selected.*

### Define Study Event - Selected CRF(s) - Select Default Version

**Physical Exam**

Required:  Double Data Entry:  Password Required:  Default Version:

Hide CRF:  Source Data Verification:

Choose Null Values (What is Null Value?)

NI <input type="checkbox"/>	NA <input type="checkbox"/>	UNK <input type="checkbox"/>	NASK <input type="checkbox"/>
ASKU <input type="checkbox"/>	NAV <input type="checkbox"/>	OTH <input type="checkbox"/>	PINF <input type="checkbox"/>
NINF <input type="checkbox"/>	MSK <input type="checkbox"/>	NP <input type="checkbox"/>	NPE <input type="checkbox"/>

6. Specify the configuration parameters for each CRF you selected, referring to the following information as needed. Note that you can specify different parameters for the CRFs at different sites; for more information, see [Create and Modify Sites](#).
- **Required:** Select if the CRF must be completed for the Event, or do not select if the CRF is optional for the Event.
  - **Double Data Entry:** Select if the data for the CRF must be entered twice. For more information on how this works, see [Double Data Entry](#).
  - **Password Required:** Select if a user must supply their OpenClinica password when marking a CRF complete at the end of data entry.
  - **Default Version:** If more than one version of the CRF exists, such as for multiple languages, select the default version for the Event. You can limit the available versions for a Site when you create or modify the Site as part of building the Study.
  - **Hide CRF:** Select to make the CRF viewable only at the Study level, meaning Sites cannot access the CRF. Typically, you hide a CRF when the data is not entered using the

OpenClinica web interface but rather is imported from files.

- **Source Data Verification:** Select one of the values from the drop-down list. Select 100% Required, Partially Required, or Not Required to help you organize CRFs in the Source Data Verification table. When you specify Not Applicable, the CRFs are not included in Source Data Verification table. For more information, see [Source Data Verification](#).
- **Choose Null Values: Please Do Not Use This Feature.** This functionality will be removed in future versions of OpenClinica. If you need to specify selections to define null values (for example "Not Done" or "Unknown," add those selections to the Response Options Text and Response Values columns of the [CRF template](#).

7. Click **Continue**.

*The Confirm Event Definition page opens.*

8. Verify the Event Definition information, then click **Confirm and Finish** to complete this Event Definition.

Confirm Event Definition Creation							
Name:	Completion Visit						
Description:	Final visit when Subject finishes treatment as part of the Study.						
Repeating:	No						
Type:	Scheduled						
Category:							
CRFs							
Name	Required	Double Data Entry	Password Required	Hide CRF	Default Version	Source Data Verification	Null Values
Physical Exam	Yes	Yes	Yes	Yes	English	100% Required	NI,
Confirm and Finish		Confirm and Create Another Definition			Cancel		

*The Build Study page opens. The Alerts and Messages sidebar panel displays a message that the Event Definition was successfully added. In the Create Event Definitions row, the Count has increased by 1, reflecting the Event Definition you just completed.*

Functional approval by Laura Keita. Signed on 2016-03-30 4:36PM

Approved for publication by Ben Baumann. Signed on 2016-03-31 10:13AM

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