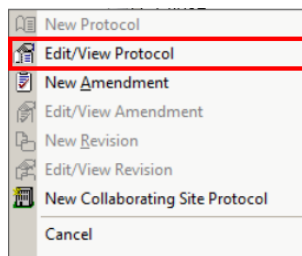


PowerTrials: POM – define protocol parameters

Quick reference guide

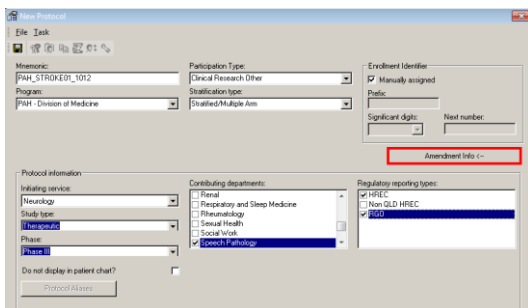
Once a protocol is created within *POM*, the protocol parameters will need to be defined within the *Amendment Info* section. Refer to *POM 2 Create New Protocol QRG* if the protocol has not yet been created.

1. Within *POM*, right click on the protocol or relevant amendment under the *Initiating Service Folder*.
2. Select *Edit/View Protocol*.




The *Edit Protocol* window will open.

3. Click the *Amendment Info* button, if required.



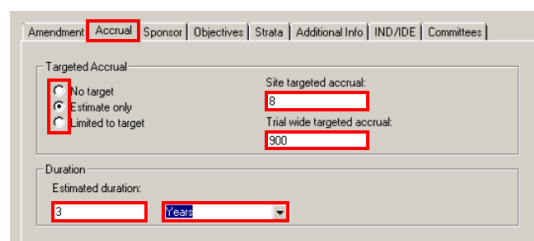
4. Complete the *Amendment* tab if not yet done:
 - *Title*: This is to be the full name of the study.
 - *Reason for amendment*: Initial protocol will be selected for a newly created protocol
 - *Treatment modalities*: Select the appropriate treatment method being used in the study

- *Diseases*: This will be the primary system in which the study is being conducted for, e.g Neurology will be Diseases – Nervous System.



5. Click the *Accrual* tab and complete information required:

- *Targeted Accrual*: Select the appropriate option if there is Targeted Accrual. If *Limited to Target* is selected, the system will not allow the user to enrol more than the number stated.
- *Site targeted accrual*: Input target number of patients for this site
- *Trial wide targeted accrual*: If the study is a multi-facility trial, input the trial-wide target for patient accrual
- *Duration*: Record the estimated duration of the research study



6. Click the *Sponsor* tab and complete information required:

- *Primary sponsor*: Search for and input the primary sponsor
 - If a sponsor is not available in the list, a job will need to be logged to have them added.
- *Funded?*: Tick checkbox if study is funded



- **Grant number:** Type Grant number, if applicable.
- **Support type:** Tick the elements the sponsor will provide
- **Additional Sponsors:** Can be added by clicking New and completing the same information.

7. Click the **Objectives** tab and complete information, if required (this is optional).

- Click **New**
- Enter **Objective Number**
- Select **Type**
- Type **Objective Statement**
- Click **OK**

8. Click the **Strata** tab and complete Arm/Cohort information if applicable. Please refer to the **POM 4 Strata and Cohorts QRG**.

9. Click the **Additional Info** tab and answer the questions displayed:

10. Click the **IND/IDE** tab and complete information required:

- Click **New**
- Click the appropriate button - **IND/IDE:**
 - **IND:** IND Number, Catalog Name, Drug Name
 - **IDE:** IDE Number, Device Name and IDE Type
- Click **Add**

11. Click the **Committees** tab and tick the relevant committee/s. Protocols will need to enter in both the Ethics and Governance Committees unless an exemption has been granted.

12. Click the **Save** icon to save the protocol settings.

13. To exit the **Edit Amendment** window, select **File** from the menu, then the **Exit** option.

