Anyone with access to a study can run an Adverse Event (AE) Summary Report or Other Reportable Information and Occurrence (ORIO) Summary Report to see how many and what types of AEs/ORIOs have occurred.

**eResearch Study Workspace**

1. Click either AE Summary Report or ORIO Summary Report from the AE/ORIO tab of the approved Study Workspace.

**Note:** Links to all individual AEs/ORIOs associated with the study are also listed.

The Adverse Event Summary Report lists all UM Serious AEs and UM Non-Serious AEs that have been submitted (but not necessarily approved) and includes information such as the Submission ID and Status (State) of the submission.

The report also includes Total # of... tables that list only AEs that have been submitted and approved.