# Accountability Logs

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Device Accountability Log
Investigators are responsible for maintaining strict control over investigational devices to ensure that the device is used only for subjects enrolled in the study. The Device Accountability Log helps maintain study device inventory and can be included in the regulatory binder.

Drug Accountability Log
Investigators are responsible for maintaining strict control over investigational drugs to ensure that the drug/biologic is used only for subjects enrolled in the study. The Drug or Biologic Dispensing/Accountability Log helps maintain an inventory of drugs used during the study and can be included in the regulatory binder.

Specimen Log
Safety laboratory assessments are usually part of most clinical trials that involve an investigational test article. Biological materials might be sent to the local laboratory or to a central laboratory. The Specimen Log helps track samples that are collected from study subjects.

Additional log resources can be found on the Partner's Healthcare website here.

Need assistance or have clinical study management questions? Please contact ResearchGo.

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- Group 1
  - Clinical Research Information Systems
  - Clinical Research Business Partners
- Group 2
  - Office of Research Administration
  - Jonsson Comprehensive Cancer Center
- Group 3
  - Office of Human Subjects Protection
  - CareConnect Website





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