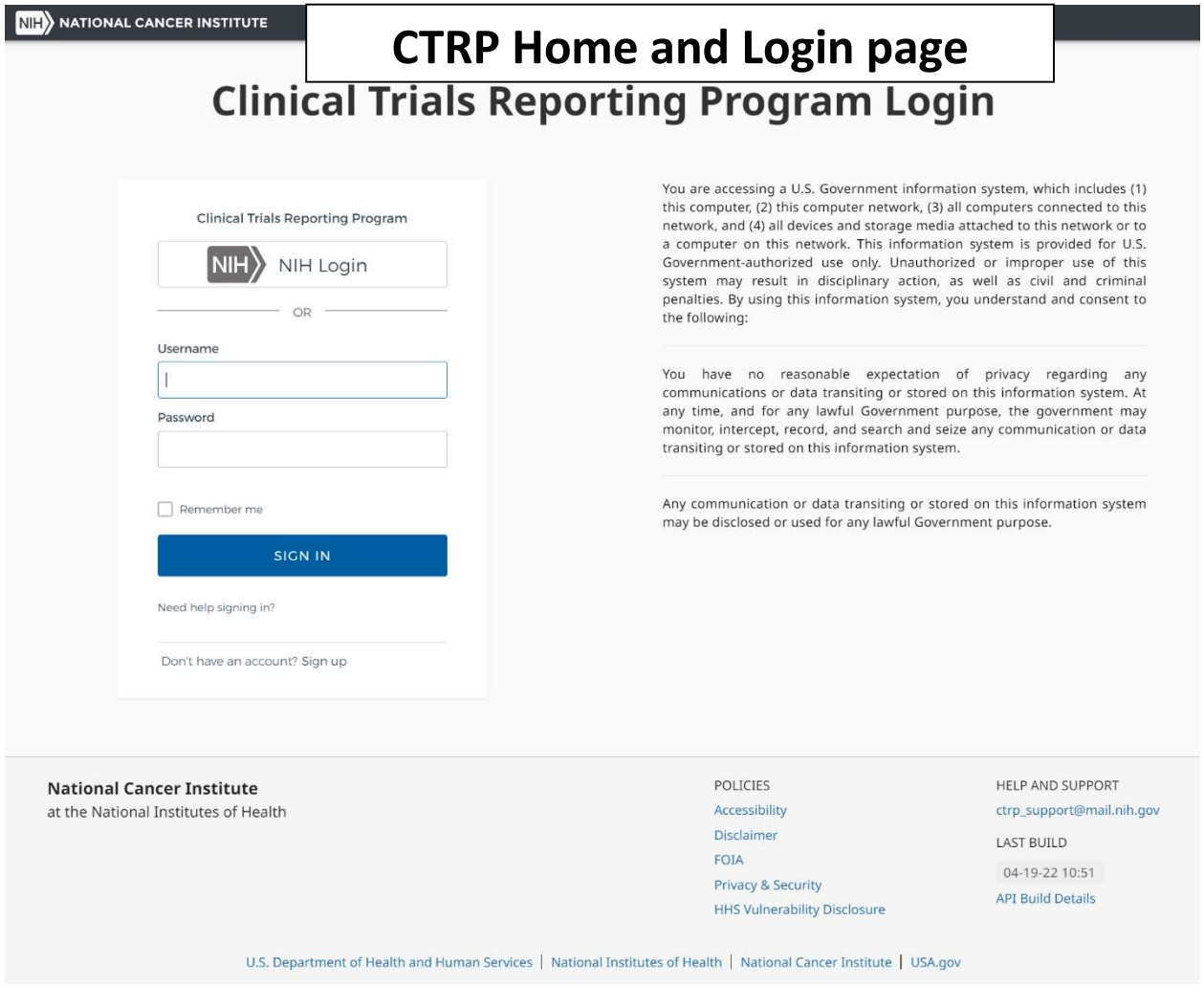
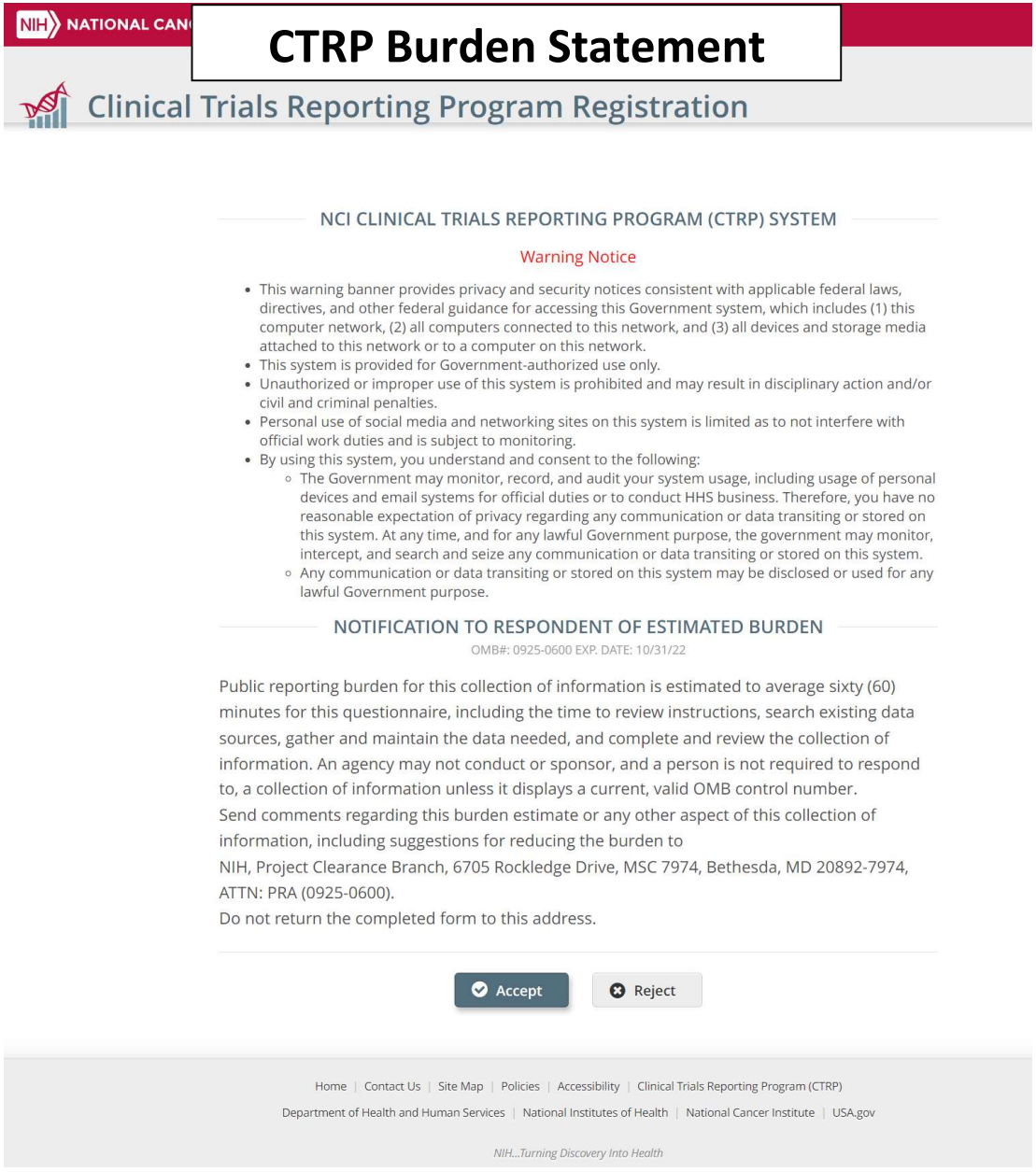
NCI CTRP Attachment 3a

NCI CTRP Registration Workflow and Screen Shots

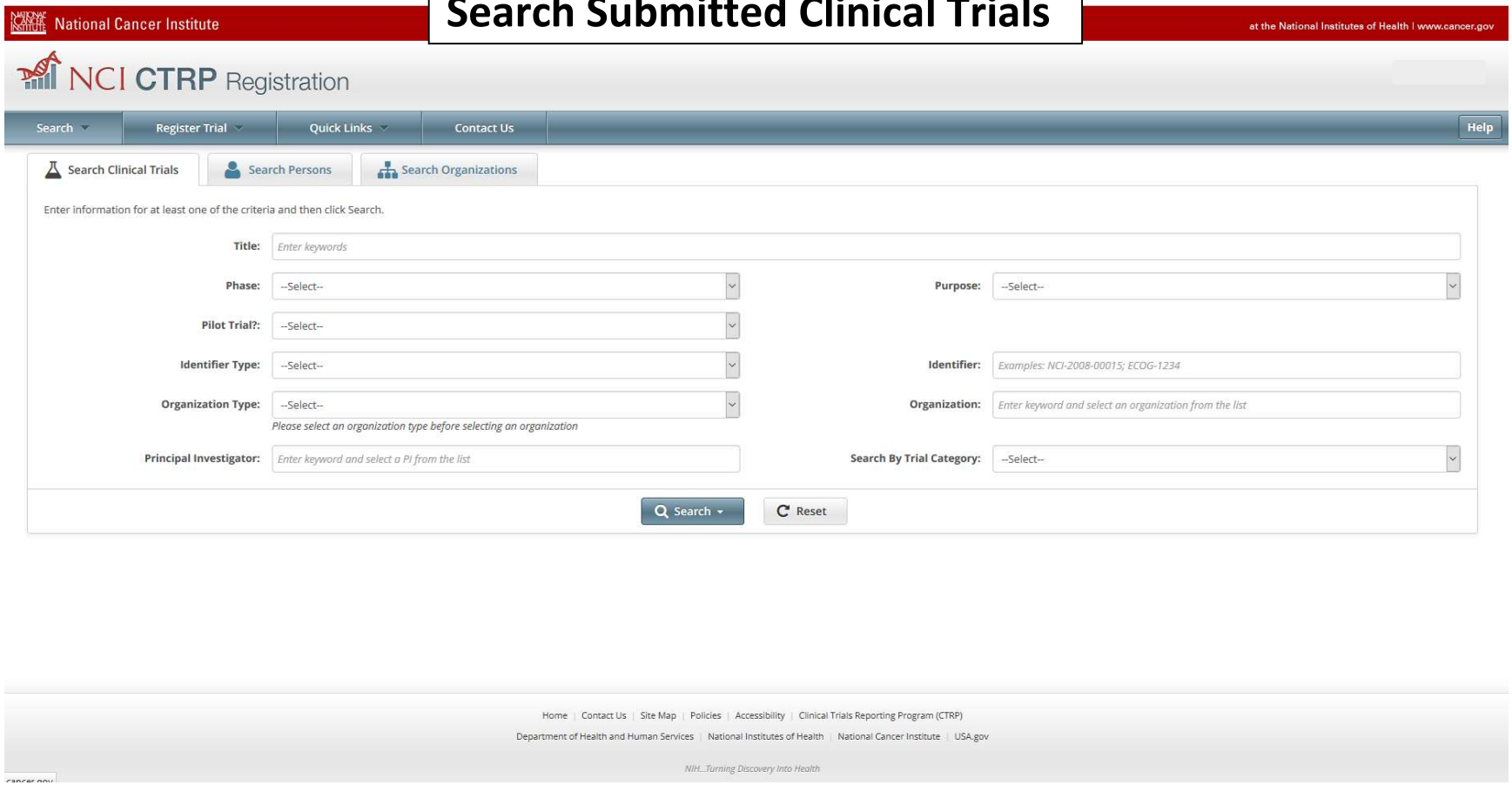
CTRP Home and Login page



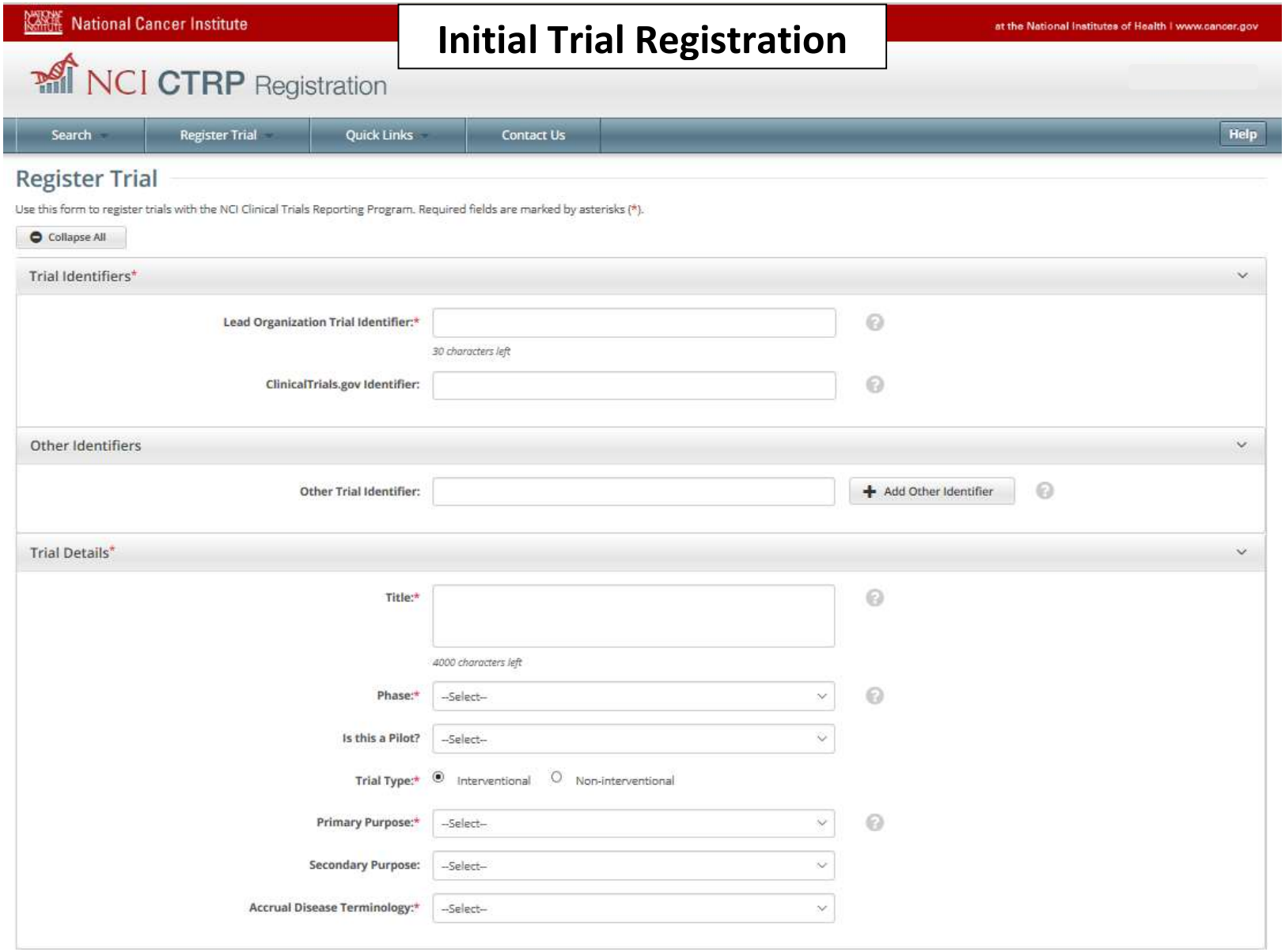
CTRP Burden Statement

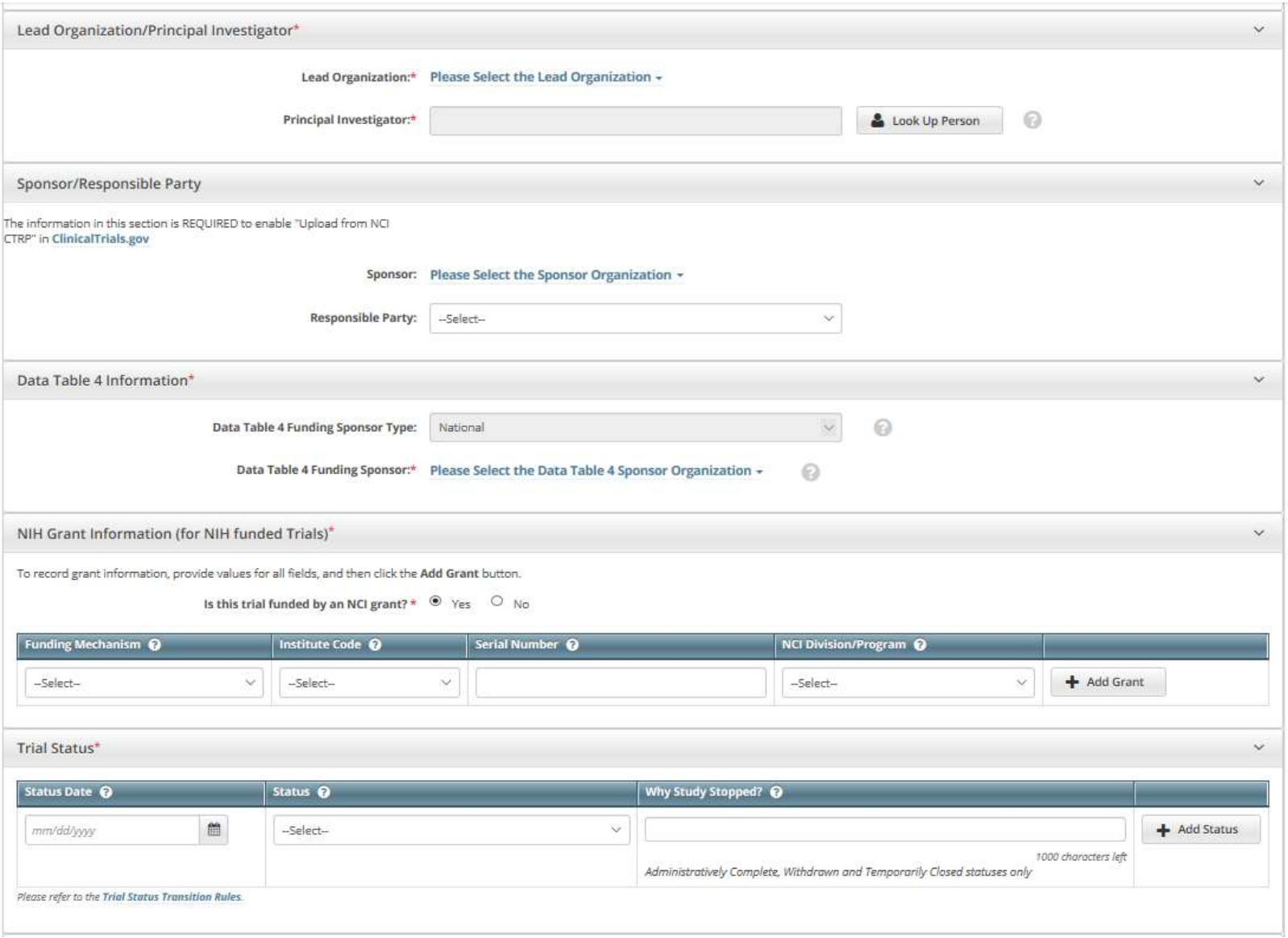


Search Submitted Clinical Trials



Initial Trial Registration




Lead Organization/Principal Investigator*
Lead Organization:* Please Select the Lead Organization
Principal Investigator:*
Look Up Person
Sponsor/Responsible Party
The information in this section is REQUIRED to enable "Upload from NCI CTRP" in ClinicalTrials.gov
Sponsor: Please Select the Sponsor Organization
Responsible Party: --Select--
Data Table 4 Information*
Data Table 4 Funding Sponsor Type: National
Data Table 4 Funding Sponsor:* Please Select the Data Table 4 Sponsor Organization
NIH Grant Information (for NIH funded Trials)*
To record grant information, provide values for all fields, and then click the Add Grant button.
Is this trial funded by an NCI grant? * Yes No
Funding Mechanism
Institute Code
Serial Number
NCI Division/Program
--Select--
--Select--
--Select--
Add Grant
Trial Status*
Status Date
Status
Why Study Stopped?
mm/dd/yyyy
--Select--
1000 characters left
Administratively Complete, Withdrawn and Temporarily Closed statuses only
Add Status
Please refer to the Trial Status Transition Rules.

Trial Dates*

Trial Start Date:* mm/dd/yyyy ○ Actual ○ Anticipated

Primary Completion Date: * mm/dd/yyyy ○ Actual ○ Anticipated

Completion Date: mm/dd/yyyy ○ Actual ○ Anticipated

*Please refer to **Trial Status Rules for Start and Completion dates.***

FDA IND/IDE Information for applicable trials

To record IND/IDE information, provide values for all fields, and then click the **Add IND/IDE** button.

IND/IDE Types	IND/IDE Number	IND/IDE Grantor	IND/IDE Holder Type	NIH Institution, NCI Division/Program Code (if applicable)	Availability of Expanded Access	Expanded Access Record (if applicable)	
-Select-		-Select-	-Select-	-Select-	Unknown		+ Add IND/IDE

‹ Scroll left/right to view full table ›

Regulatory Information

The information in this section is REQUIRED to enable "Upload from NCI CTRP" in **ClinicalTrials.gov**

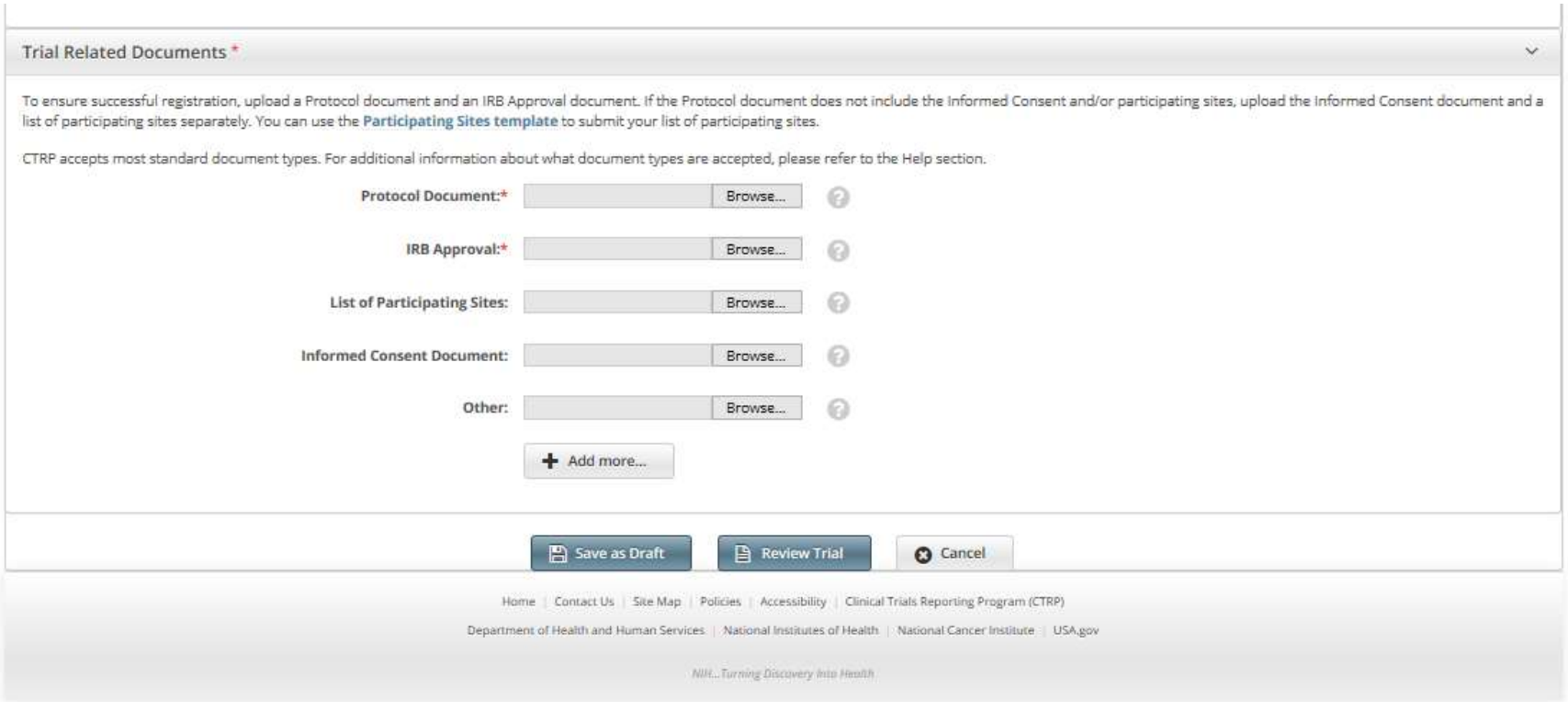
Studies a U.S. FDA-regulated Drug Product :

Studies a U.S. FDA-regulated Device Product :

Product Exported from the U.S :

FDA Regulated Intervention Indicator :

Data Monitoring Committee Appointed Indicator :


Trial Related Documents *
To ensure successful registration, upload a Protocol document and an IRB Approval document. If the Protocol document does not include the Informed Consent and/or participating sites, upload the Informed Consent document and a list of participating sites separately. You can use the Participating Sites template to submit your list of participating sites.
CTRP accepts most standard document types. For additional information about what document types are accepted, please refer to the Help section.
Protocol Document:*
Browse...
IRB Approval:*
Browse...
List of Participating Sites:
Browse...
Informed Consent Document:
Browse...
Other:
Browse...
Add more...
Save as Draft
Review Trial
Cancel
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Department of Health and Human Services | National Institutes of Health | National Cancer Institute | USA.gov
NIH...Turning Discovery Into Health