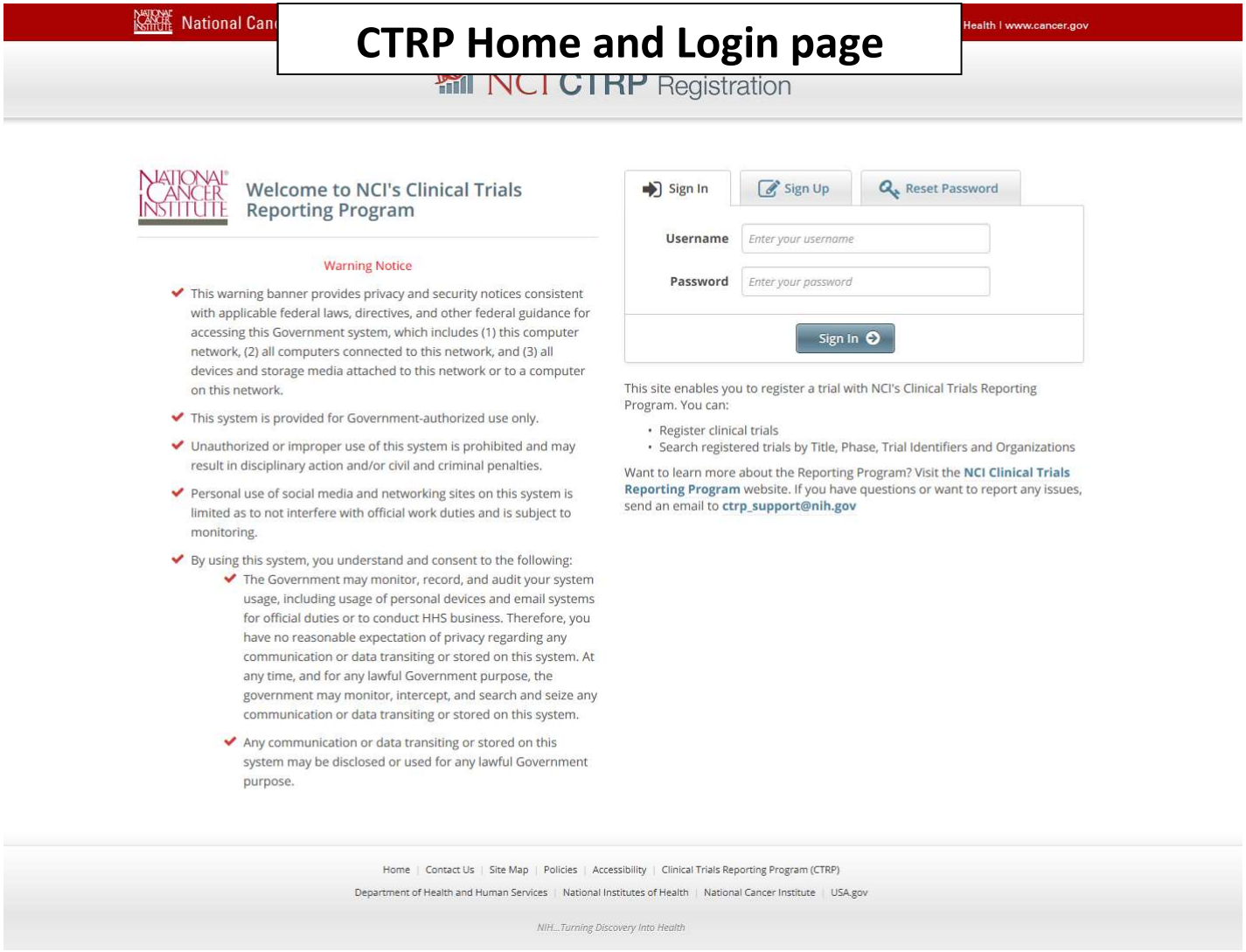
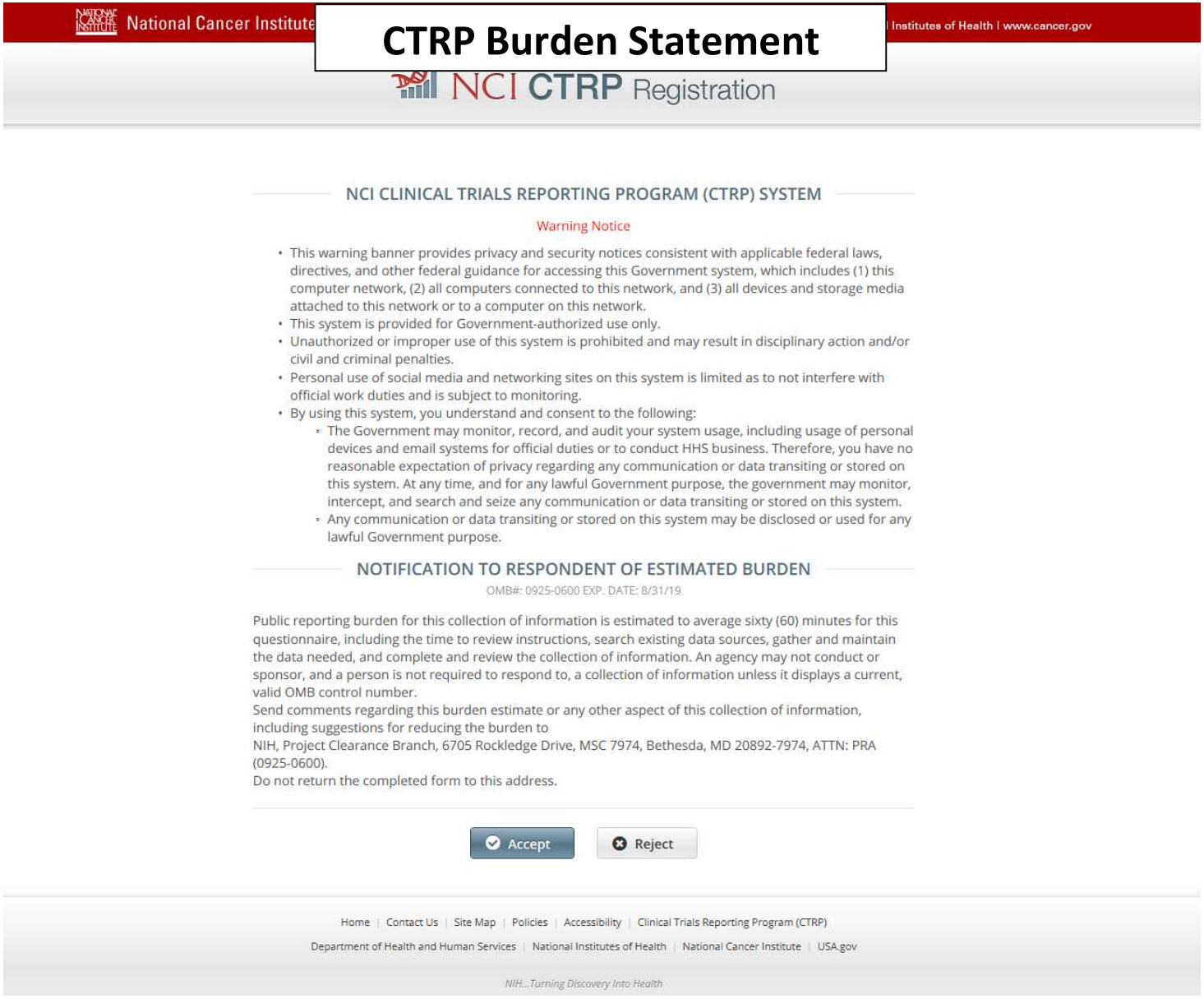
NCI CTRP Attachment 3b

NCI CTRP Update Workflow and Screen Shots

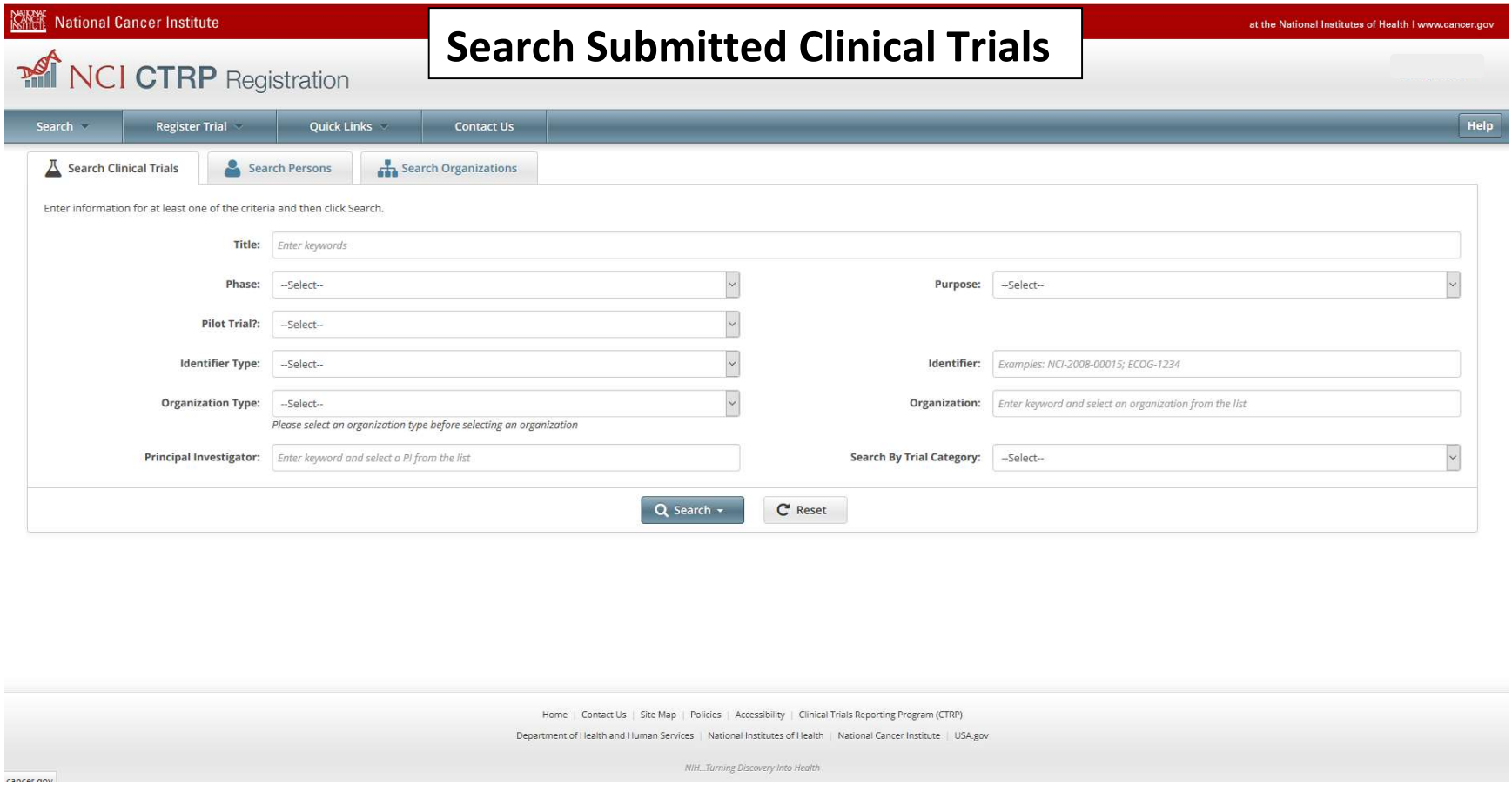
CTRP Home and Login page



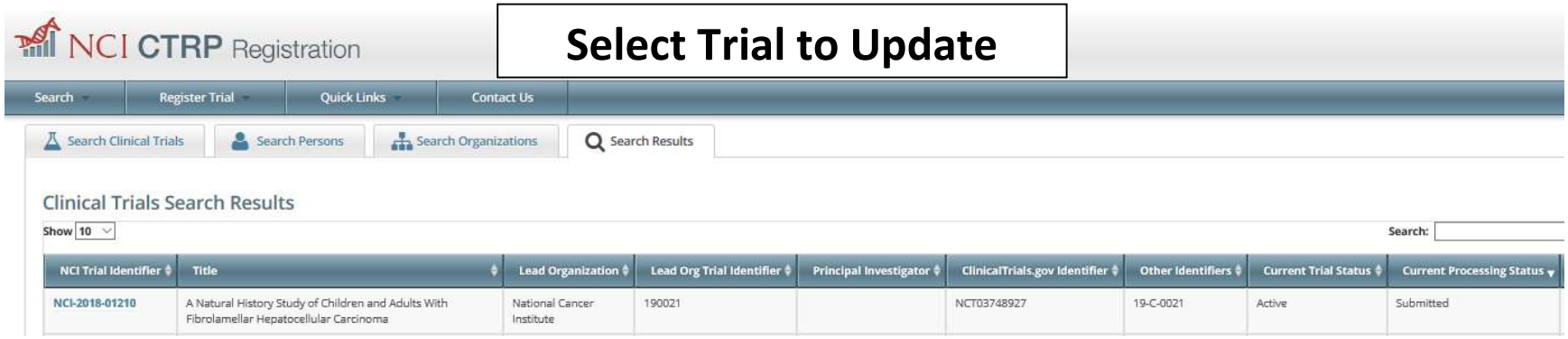
CTRP Burden Statement



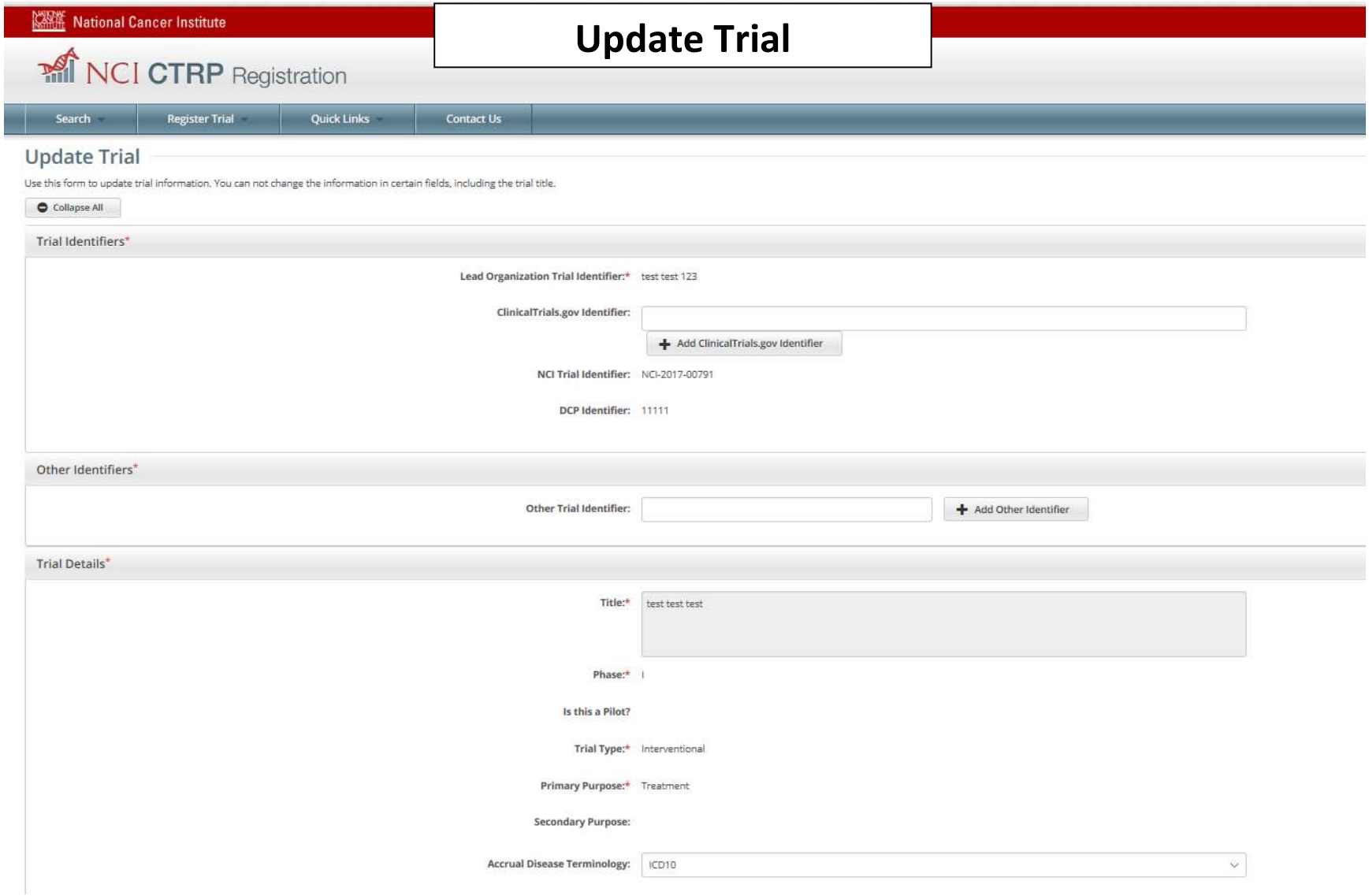
Search Submitted Clinical Trials



Select Trial to Update



Update Trial



Lead Organization/Principal Investigator*

Lead Organization:* Dana-Farber Cancer Institute

Principal Investigator:* Izbicki, Michael

Sponsor / Responsible Party

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

Sponsor: Dana-Farber Cancer Institute

Responsible Party: Sponsor

Data Table 4 Information*

Data Table 4 Funding Sponsor Type:* National

Data Table 4 Funding Sponsor: * Dana-Farber Cancer Institute

Program Code: *Select program code*

NIH Grant Information (for NIH funded Trials)*

Funding Mechanism	Institute Code	Serial Number	NCI Division/Program	
--Select--	--Select--		--Select--	+ Add Grant

Funding Mechanism	Institute Code	Serial Number	NCI Division/Program Code
P30	CA	90687	TRP

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mm/dd/yyyy	--Select--	1000 characters left *Administratively Complete, Withdrawn and Temporarily Closed statuses only*	+ Add Status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 Status History

Show 10 entries

Status Date	Status	Comments	Validation Messages	Actions
06/26/2017	In Review			
09/24/2018	Approved			
09/24/2018	Administratively Complete			
09/24/2018	Temporarily Closed to Accrual			

Showing 1 to 4 of 4 entries

Previous 1 Next

Trial Dates*

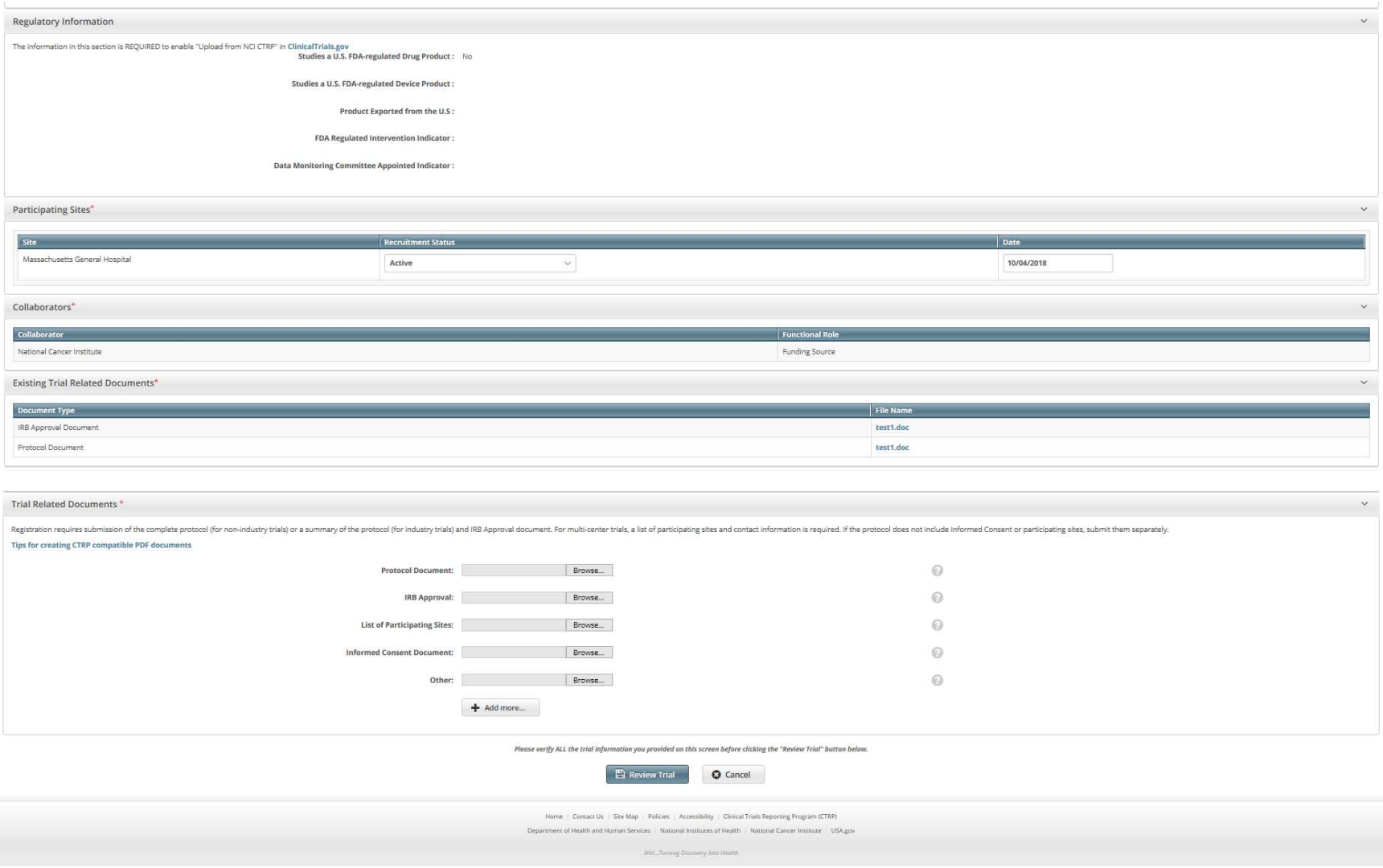
Trial Start Date:* 07/04/2017 ○ Actual ○ Anticipated

Primary Completion Date: 09/24/2018 ○ Actual ◉ Anticipated ○ N/A

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*


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 : No
Studies a U.S. FDA-regulated Device Product :
Product Exported from the U.S :
FDA Regulated Intervention Indicator :
Data Monitoring Committee Appointed Indicator :
Participating Sites*
Site
Recruitment Status
Date
Massachusetts General Hospital
Active
10/04/2018
Collaborators*
Collaborator
Functional Role
National Cancer Institute
Funding Source
Existing Trial Related Documents*
Document Type
File Name
IRB Approval Document
test1.doc
Protocol Document
test1.doc
Trial Related Documents *
Registration requires submission of the complete protocol (for non-industry trials) or a summary of the protocol (for industry trials) and IRB Approval document. For multi-center trials, a list of participating sites and contact information is required. If the protocol does not include Informed Consent or participating sites, submit them separately.
Tips for creating CTRP compatible PDF documents
Protocol Document:
Browse...
IRB Approval:
Browse...
List of Participating Sites:
Browse...
Informed Consent Document:
Browse...
Other:
Browse...
Add more...
Please verify ALL the trial information you provided on this screen before clicking the "Review Trial" button below.
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