## **NCI CTRP Attachment 3a**

### **NCI CTRP Registration Workflow and Screen Shots**

- Step 1: User accesses the NCI Clinical Trials Reporting Program website at http://trials.nci.nih.gov see screenshot, page 2
- Step 2: User enters "Username" and "Password" see screenshot, page 2
- Step 3: User reviews NCI Clinical Trials Reporting Program burden statement see screenshot, page 3
- Step 4: System displays "Search Submitted Clinical Trials" page see screenshot, page 4
- Step 5: User selects to perform "Initial Trial Registration" and completes initial registration see screenshots, pages 5 8



# **CTRP Home and Login page**

## Clinical Trials Reporting Program Login

NIH Login	
OR	
me	
SIGN IN	
	OR —

You are accessing a U.S. Government information system, which includes (1) this computer, (2) this computer network, (3) all computers connected to this network, and (4) all devices and storage media attached to this network or to a computer on this network. This information system is provided for U.S. Government-authorized use only. Unauthorized or improper use of this system may result in disciplinary action, as well as civil and criminal penalties. By using this information system, you understand and consent to the following:

You have no reasonable expectation of privacy regarding any communications or data transiting or stored on this information system. At any time, and for any lawful Government purpose, the government may monitor, intercept, record, and search and seize any communication or data transiting or stored on this information system.

Any communication or data transiting or stored on this information system may be disclosed or used for any lawful Government purpose.

### **National Cancer Institute**

at the National Institutes of Health

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HHS Vulnerability Disclosure

HELP AND SUPPORT ctrp\_support@mail.nih.gov

LAST BUILD

04-19-22 10:51

API Build Details

U.S. Department of Health and Human Services | National Institutes of Health | National Cancer Institute | USA.gov



### **CTRP Burden Statement**



### **Clinical Trials Reporting Program Registration**

#### NCI CLINICAL TRIALS REPORTING PROGRAM (CTRP) SYSTEM

### Warning Notice

- This warning banner provides privacy and security notices consistent with applicable federal laws, directives, and other federal guidance for accessing this Government system, which includes (1) this computer network, (2) all computers connected to this network, and (3) all devices and storage media attached to this network or to a computer on this network.
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- Unauthorized or improper use of this system is prohibited and may result in disciplinary action and/or civil and criminal penalties.
- Personal use of social media and networking sites on this system is limited as to not interfere with
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    intercept, and search and seize any communication or data transiting or stored on this system.
  - Any communication or data transiting or stored on this system may be disclosed or used for any lawful Government purpose.

#### NOTIFICATION TO RESPONDENT OF ESTIMATED BURDEN

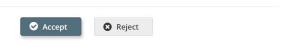
OMB#: 0925-0600 EXP. DATE: 10/31/22

Public reporting burden for this collection of information is estimated to average sixty (60) minutes for this questionnaire, including the time to review instructions, search existing data sources, gather and maintain the data needed, and complete and review the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a current, valid OMB control number.

Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden to

NIH, Project Clearance Branch, 6705 Rockledge Drive, MSC 7974, Bethesda, MD 20892-7974, ATTN: PRA (0925-0600).

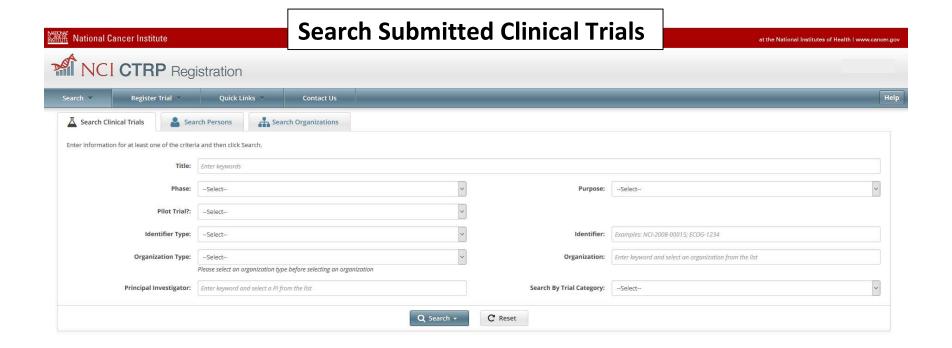
Do not return the completed form to this address.



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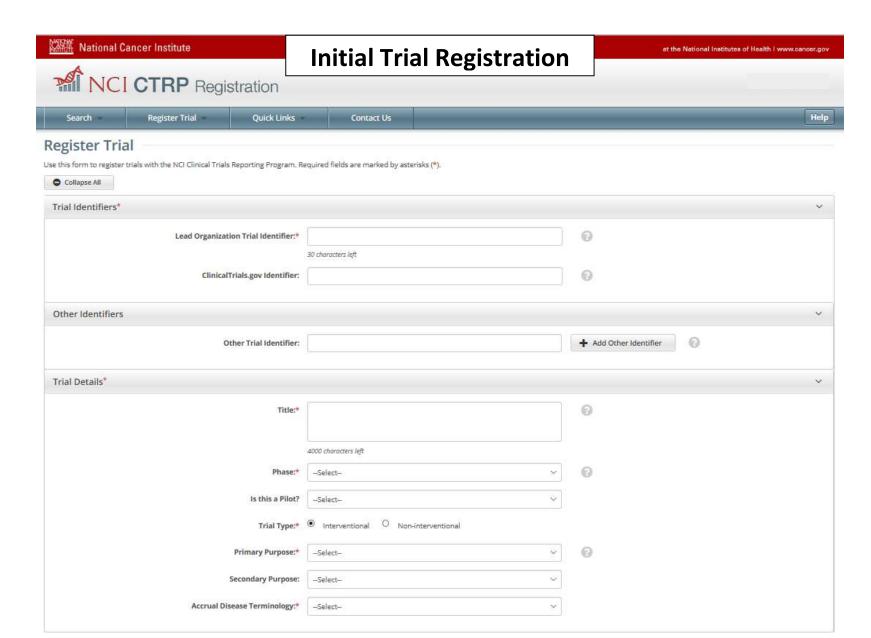
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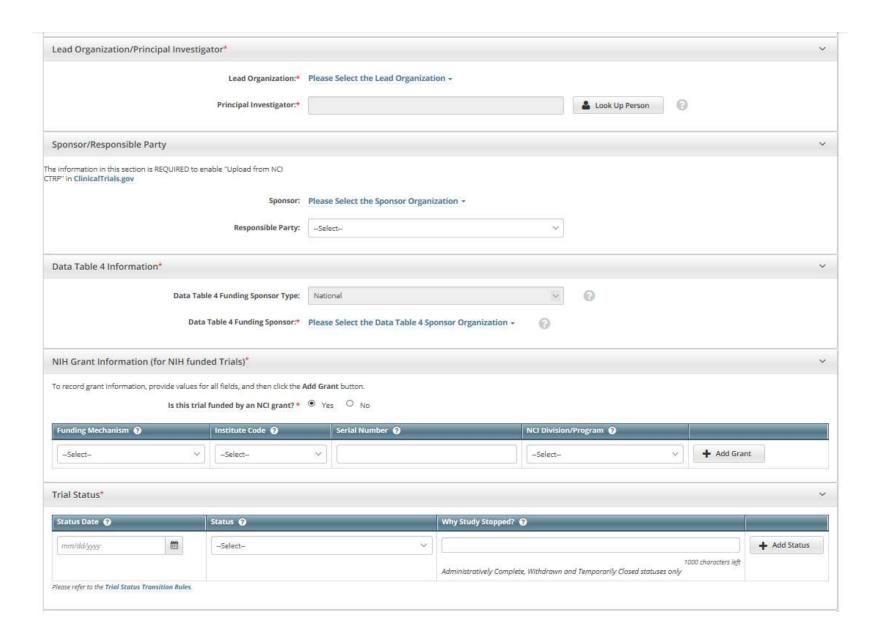


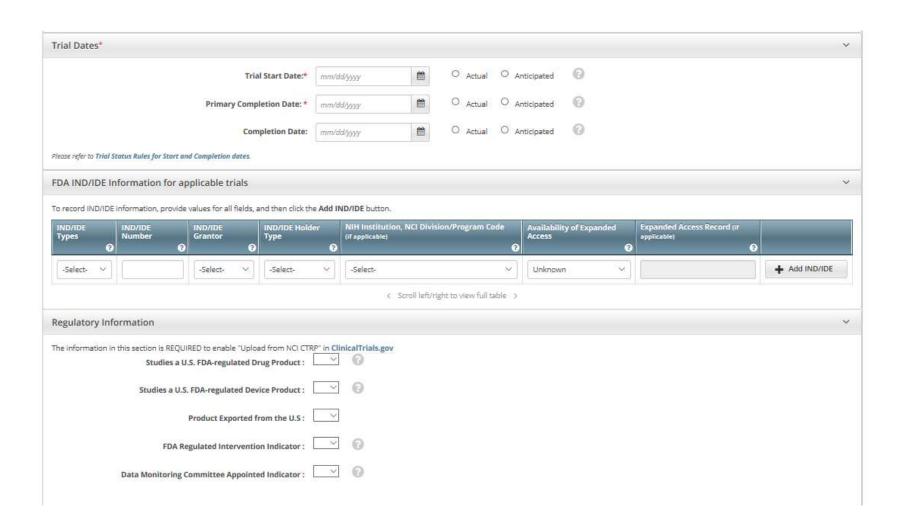
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Trial Related Documents*		¥
list of participating sites separately. You can use the <b>Participating Sites tem</b> CTRP accepts most standard document types. For additional information abo	plate to submit your list of participating sit ut what document types are accepted, plea	use refer to the Help section.
Protocol Document:*	Browse	0
IRB Approval:*	Browse	0
List of Participating Sites:	Browse	0
Informed Consent Document:	Browse	0
Other:	Browse	0
	+ Add more	
	🖺 Save as Draft 📔 Review	vTrial Cancel
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