

## **NCI CTRP Attachment 3e**

### **NCI CTRP Aggregate Accrual Portal Workflow and Screen Shots**

Step 1: User accesses the NCI Clinical Trials Reporting Program Accrual website at <http://trials.nci.nih.gov/accrual> – see screenshot, page 2

Step 2: User enters “Username” and “Password” – see screenshot, page 2


Step 3: User reviews NCI Clinical Trials Reporting Program Accrual burden statement – see screenshot, page 3

Step 4: User selects a trial to “Submit Aggregate Study Subject Accrual Information” and submits aggregate accrual information that registered trial – see screenshot, pages 4 – 5

# CTRP Accrual Home and Login page

## Clinical Trials Reporting Program Login

Clinical Trials Reporting Program

 NIH Login

OR

Username

Password

Remember me

**SIGN IN**

Need help signing in?

Don't have an account? [Sign up](#)

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[ctrp\\_support@mail.nih.gov](mailto:ctrp_support@mail.nih.gov)  
**LAST BUILD**  
04-19-22 10:51  
[API Build Details](#)

# CTRP Accrual Burden Statement



## 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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## 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 fifteen (15) 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.

Accept

Reject

# Select Trial to Submit Aggregate Accrual Information

## Trial Search

NCI Trial Identifier  ?  
 ClinicalTrials.gov ID  ?  
 Official Title  ?

## List of Trials

Show 10   << < 1 > >>

NCI Trial Identifier	Official Title	Current Trial Status	Trial Type	Accrual Disease Terminology
NCI-2015-02030	A Phase III Randomized, Open-Label, Multi-Center, Global Study of MEDI4736 in Combination With Tremelimumab Therapy or MEDI4736 Monotherapy Versus Standard of Care Platinum-Based Chemotherapy in First Line Treatment of Patients With Advanced or Metastatic Non Small-Cell Lung Cancer (NSCLC).	Active	Interventional	SDC

Showing 1 to 1 of 1  
 Export options: CSV | Excel

# Submit Aggregate Accrual Information



- Trial Search
- Batch Upload
- Prior Submissions
- Accrual Counts
- Disease Search
- Quick Links
- Contact Us
- Help

NCI-2017-01026: The Natural History of Congenital Trigger Thumbs  
 Lead Organization Trial ID: 32047  
 Lead Organization: Huntsman Cancer Institute/University of Utah

## Participating Site Subject Accrual Count

Message: Record Created.

Show 10

Search:  << < 1 > >>

PO ID	Site Name	# of Subjects Enrolled	Cut-Off Date	Date Last Updated	Actions
149280	Duke University Medical Center	12	12/31/2015	05/02/2019 16:04	+ [Save] [Delete]

Showing 1 to 1 of 1

<< < 1 > >>

Reset