

NCI CTRP Attachment 3d

NCI CTRP 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 to “Submit Study Subject Accrual Information” and submits subject level accrual information on a registered trial – see screenshots, pages 4 - 6



Welcome to NCI's Clinical Trials Reporting Program

This site allows for the upload of accrual data for trials submitted to CTRP. If trials are NCI-supported CTEP/DCP trials, accrual submission is handled via your normal accrual reporting, and it is not necessary to submit additional accrual data via this site. If you have any questions regarding accrual submission for a specific trial, please contact the Clinical Trials Reporting Office (CTRO) at ncictro@mail.nih.gov.

Want to learn more about the Reporting Program? Visit the [NCI Clinical Trials Reporting Program](#) website. You can email NCI CBIIT Application Support at ncicbiit@mail.nih.gov if you have questions or need assistance.

 Sign In

 Sign Up

 Forgot Password

Username

Password

Sign In 

CTRP Accrual Burden Statement



NCI CLINICAL TRIALS REPORTING PROGRAM (CTRP) SYSTEM

This is a U.S. Government computer system, which may be accessed and used only for authorized Government business by authorized personnel. Unauthorized access or use of this computer system may subject violators to criminal, civil, and/or administrative action.

All information on this computer system may be intercepted, recorded, read, copied, and disclosed by and to authorized personnel for official purposes, including criminal investigations. Such information includes sensitive data encrypted to comply with confidentiality and privacy requirements. Access or use of this computer system by any person, whether authorized or unauthorized, constitutes consent to these terms. There is no right of privacy in this system.

NOTIFICATION TO RESPONDENT OF ESTIMATED BURDEN


OMB#: 0925-0600 EXP. DATE: 5/31/16


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.

National Cancer Institute

NCI CTRP

Accrual

Submit Study Subject Accrual Information

at the National Institutes of Health | www.cancer.gov

Sophia Rarhai

Trial Search

Batch Upload

Prior Submissions

Accrual Counts

Disease Search

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Trial Search

NCI Trial Identifier

NCI-2015-01833

?

ClinicalTrials.gov ID

?

Official Title

?

Search Trials

Reset

List of Trials

Show 10

Search:

Choose columns

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NCI Trial Identifier	Official Title	Current Trial Status	Trial Type	Accrual Disease Terminology
NCI-2015-01833	Phase I Protocol Evaluating a Stereotactic Boost/Treatment for Recurrent or Metastatic Cancer of the Head and Neck	Active	Interventional	ICD9

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Export options: CSV | Excel

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Clinical Trials Reporting Program (CTRP)

Department of Health and Human Services

National Institutes of Health

National Cancer Institute

USA.gov

NIH...Turning Discovery Into Health

Principal Investigator: Schoenfeld, Jonathan

Lead Organization: Dana-Farber Harvard Cancer Center

Study Subject ID: *

Study Subject Birth Date (MM/YYYY): *

mm/yyyyStudy Subject Gender: *--Select--

Study Subject Race: ★

American Indian or Alaska Native
Asian
Black or African American
Native Hawaiian or Other Pacific Islander
Not Reported
Unknown
White

To select multiple races, select one race, and then press and hold the CTRL key as you select the other(s).

Study Subject Ethnicity: *

--Select--Study Subject Country: ★United States

Study Subject Zip Code:

Registration Date: ★

mm/dd/yyyy

Study Subject method of payment:

--Select--


Disease:★

Look Up




Participating Site: ★

--Select-- Save

National Cancer Institute

at the National Institutes of Health | www.cancer.gov

NCI CTRP

Accrual

Sophia Rarhai

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NCI-2015-01833: Phase I Protocol Evaluating a Stereotactic Boost/Treatment for Recurrent or Metastatic Cancer of the Head and Neck

Lead Organization Trial ID: 15-183
Lead Organization: Dana-Farber Harvard Cancer Center

Principal Investigator: Schoenfeld, Jonathan

Search Study Subject

Study Subject ID

Participating Site

--Select--

Study Subject Birth Date (MM/YYYY):

mm/yyyy

Search

+ Add New Study Subject



List of Study Subjects

✓ Message: Record Created.

Show 10

Search:

<< < 1 > >>

Study Subject ID	Registration Date	Participating Site	Last Update Date/Time	Actions
TEST	02/03/2016	Dana-Farber Harvard Cancer Center	02/03/2016 17:58	 

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