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 a trial to "Submit Study Subject Accrual Information" and submits subject level accrual information on that registered trial – see screenshots, pages 4 – 6

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National Cancer Institute at the National Institutes of Health	POLICIES Accessibility Disclaimer FOIA Privacy & Security HHS Vulnerability Disclosure	HELP AND SUPPORT ctrp_support@mail.nih.gov LAST BUILD 04-19-22 10:51 API Build Details

Nation

CTRP Accrual Burden Statement

cer.gov

MCI CTRP Accrual

NCI CLINICAL TRIALS REPORTING PROGRAM (CTRP) SYSTEM

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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

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