

preparedness to respond to disasters in their communities. The MRC unit data collection has been refined to eliminate

duplication and streamline data collection tools.

ANNUALIZED BURDEN TABLE

Forms (if necessary)	Type of respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Unit Profile	MRC Unit Leader	748	4	15/60	748
Capability Assessment	MRC Unit Leader	748	1	30/60	374
Factors for Success	MRC Unit Leader	748	1	30/60	374
Unit Activity Reporting	MRC Unit Leader	748	4	15/60	748
Total	10	2,244

Sherrette A. Funn,

*Paperwork Reduction Act Reports Clearance
Officer, Office of the Secretary.*

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BILLING CODE 4150-47-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Cancer Therapy Evaluation Program (CTEP) Branch and Support Contracts Forms and Surveys (NCI)

AGENCY: National Institutes of Health,
HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Michael Montello, Cancer Therapy Evaluation Program—DCTD, National Cancer Institute, 9609 Medical Center Drive, Rockville, Maryland, 20850 or call non-toll-free number (240) 276-6080 or email your request, including your address to: montellom@mail.nih.gov. Formal requests for

additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Proposed Collection Title: Cancer Therapy Evaluation Program (CTEP) Branch and Support Contracts Forms and Surveys (NCI), 0925-0753, Expiration Date 05/31/2024, REVISION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This is a request for OMB to approve the revised information collection, Cancer Therapy Evaluation Program (CTEP) Support Contracts Forms and Survey. This revision removes one form (A17 CTSU System Access Request Form), adds one new form (A22 CLASS Course Setup Request Form), revises three forms (A18 CTSU Open Rave Request Form; B41 Annual Principal Investigator Worksheet about Local Context; B47 CIRB Waiver of

Consent Request Supplemental Form), and includes an updated Privacy Impact Assessment. The National Cancer Institute (NCI) Cancer Therapy Evaluation Program (CTEP) and the Division of Cancer Prevention (DCP) fund an extensive national program of cancer research, sponsoring clinical trials in cancer prevention, symptom management and treatment for qualified clinical investigators. As part of this effort, CTEP implements programs to register clinical site investigators and clinical site staff, and to oversee the conduct of research at the clinical sites. CTEP and DCP also oversee two support programs, the NCI Central Institutional Review Board (CIRB) and the Cancer Trial Support Unit (CTSU). The combined systems and processes for initiating and managing clinical trials is termed the Clinical Oncology Research Enterprise (CORE) and represents an integrated set of information systems and processes which support investigator registration, trial oversight, patient enrollment, and clinical data collection. The information collected is required to ensure compliance with applicable federal regulations governing the conduct of human subject's research (45 CFR 46 and 21 CFR 50), and when CTEP acts as the Investigational New Drug (IND) holder (Food and Drug Administration (FDA) regulations pertaining to the sponsor of clinical trials and the selection of qualified investigators under 21 CFR 312.53). Survey collections assess satisfaction and provide feedback to guide improvements with processes and technology.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 151,769 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
CTSU IRB/Regulatory Approval Transmittal Form (Attachment A01).	Health Care Practitioner	2,444	12	2/60	978
CTSU IRB Certification Form (Attachment A02) ..	Health Care Practitioner	2,444	12	10/60	4,888
Withdrawal from Protocol Participation Form (Attachment A03).	Health Care Practitioner	279	1	10/60	47
Site Addition Form (Attachment A04)	Health Care Practitioner	80	12	10/60	160
CTSU Request for Clinical Brochure (Attachment A06).	Health Care Practitioner	360	1	10/60	60
CTSU Supply Request Form (Attachment A07) ..	Health Care Practitioner	90	12	10/60	180
RTOG 0834 CTSU Data Transmittal Form (Attachment A10).	Health Care Practitioner	12	76	10/60	152
CTSU Patient Enrollment Transmittal Form (Attachment A15).	Health Care Practitioner	12	12	10/60	24
CTSU Transfer Form (Attachment A16)	Health Care Practitioner	360	2	10/60	120
CTSU OPEN Rave Request Form (Attachment A18).	Health Care Practitioner	30	21	10/60	105
CTSU LPO Form Creation (Attachment A19)	Health Care Practitioner	5	2	120/60	20
CTSU Site Form Creation (Attachment A20)	Health Care Practitioner	400	10	30/60	2,000
CTSU Electronic Signature Form (Attachment A21).	Health Care Practitioner	400	10	10/60	667
CTSU CLASS Course Setup Form (Attachment A22).	Health Care Practitioner	10	2	20/60	7
NCI CIRB AA & DOR between the NCI CIRB and Signatory Institution (Attachment B01).	Participants	50	1	15/60	13
NCI CIRB Signatory Enrollment Form (Attachment B02).	Participants	50	1	15/60	13
CIRB Board Member Application (Attachment B03).	Board Member	100	1	30/60	50
CIRB Member COI Screening Worksheet (Attachment B08).	Board Members	100	1	15/60	25
CIRB COI Screening for CIRB meetings (Attachment B09).	Board Members	72	1	15/60	18
CIRB IR Application (Attachment B10)	Health Care Practitioner	80	1	60/60	80
CIRB IR Application for Exempt Studies (Attachment B11).	Health Care Practitioner	4	1	30/60	2
CIRB Amendment Review Application (Attachment B12).	Health Care Practitioner	400	1	15/60	100
CIRB Ancillary Studies Application (Attachment B13).	Health Care Practitioner	1	1	60/60	1
CIRB Continuing Review Application (Attachment B14).	Health Care Practitioner	400	1	15/60	100
Adult IR of Cooperative Group Protocol (Attachment B15).	Board Members	65	1	180/60	195
Pediatric IR of Cooperative Group Protocol (Attachment B16).	Board Members	15	1	180/60	45
Adult Continuing Review of Cooperative Group Protocol (Attachment B17).	Board Members	275	1	60/60	275
Adult Amendment of Cooperative Group Protocol (Attachment B19).	Board Members	40	1	120/60	80
Pediatric Amendment of Cooperative Group Protocol (Attachment B20).	Board Members	25	1	120/60	50
Pharmacist's Review of a Cooperative Group Study (Attachment B21).	Board Members	50	1	120/60	100
Adult Expedited Amendment Review (Attachment B23).	Board Members	348	1	30/60	174
Pediatric Expedited Amendment Review (Attachment B24).	Board Members	140	1	30/60	70
Adult Expedited Continuing Review (Attachment B25).	Board Members	140	1	30/60	70
Pediatric Expedited Continuing Review (Attachment B26).	Board Members	36	1	30/60	18
Adult Cooperative Group Response to CIRB Review (Attachment B27).	Health Care Practitioner	30	1	60/60	30
Pediatric Cooperative Group Response to CIRB Review (Attachment B28).	Health Care Practitioner	5	1	60/60	5
Adult Expedited Study Chair Response to Required Modifications (Attachment B29).	Board Members	40	1	30/60	20
Reviewer Worksheet—Determination of UP or SCN (Attachment B31).	Board Members	400	1	10/60	67

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Reviewer Worksheet—CIRB Statistical Reviewer Form (Attachment B32).	Board Members	100	1	15/60	25
CIRB Application for Translated Documents (Attachment B33).	Health Care Practitioner	100	1	30/60	50
Reviewer Worksheet of Translated Documents (Attachment B34).	Board Members	100	1	15/60	25
Reviewer Worksheet of Recruitment Material (Attachment B35).	Board Members	20	1	15/60	5
Reviewer Worksheet Expedited Study Closure Review (Attachment B36).	Board Members	20	1	15/60	5
Reviewer Worksheet of Expedited IR (Attachment B38).	Board Members	5	1	30/60	3
Annual Signatory Institution Worksheet About Local Context (Attachment B40).	Health Care Practitioner	400	1	40/60	267
Annual Principal Investigator Worksheet About Local Context (Attachment B41).	Health Care Practitioner	1,800	1	20/60	600
Study-Specific Worksheet About Local Context (Attachment B42).	Health Care Practitioner	4,800	1	15/60	1,200
Study Closure or Transfer of Study Review Responsibility (Attachment B43).	Health Care Practitioner	1,680	1	15/60	420
Unanticipated Problem or Serious or Continuing Noncompliance Reporting Form (Attachment B44).	Health Care Practitioner	360	1	20/60	120
Change of Signatory Institution PI Form (Attachment B45).	Health Care Practitioner	120	1	20/60	40
Request Waiver of Assent Form (Attachment B46).	Health Care Practitioner	35	1	20/60	12
CIRB Waiver of Consent Request Supplemental Form (Attachment B47).	Health Care Practitioner	20	1	15/60	5
Review Worksheet CIRB Review for Inclusion of Incarcerated Participants (Attachment B48).	Board Members	20	1	60/60	20
Notification of Incarcerated Participant Form (B49).	Health Care Practitioner	20	1	20/60	7
CTSU OPEN Survey (Attachment C03)	Health Care Practitioner	10	1	15/60	3
CIRB Customer Satisfaction Survey (Attachment C04).	Participants	600	1	15/60	150
Follow-up Survey (Communication Audit) (Attachment C05).	Participants/Board Members.	300	1	15/60	75
CIRB Board Member Annual Assessment Survey (Attachment C07).	Board Members	60	1	15/60	15
PIO Customer Satisfaction Survey (Attachment C08).	Health Care Practitioner	60	1	5/60	5
Audit Scheduling Form (Attachment D01)	Health Care Practitioner	152	5	21/60	266
Preliminary Audit Finding Form (Attachment D02)	Health Care Practitioner	152	5	10/60	127
Audit Maintenance Form (Attachment D03)	Health Care Practitioner	152	5	9/60	114
Final Audit finding Report Form (Attachment D04).	Health Care Practitioner	75	11	1,098/60	15,098
Follow-up Form (Attachment D05)	Health Care Practitioner	75	7	27/60	236
Roster Maintenance Form (Attachment D06)	Health Care Practitioner	5	1	18/60	2
Final Report and CAPA Request Form (Attachment D07).	Health Care Practitioner	12	9	1,800/60	3,240
NCI/DCTD/CTEP FDA Form 1572 for Annual Submission (Attachment E01).	Physician	26,500	1	15/60	6,625
NCI/DCTD/CTE Biosketch (Attachment E02)	Physician; Health Care Practitioner.	48,000	1	120/60	96,000
NCI/DCTD/CTEP Financial Disclosure Form (Attachment E03).	Physician; Health Care Practitioner.	48,000	1	15/60	12,000
NCI/DCTD/CTEP Agent Shipment Form (ASF) (Attachment E04).	Physician	24,000	1	10/60	4,000
Totals	167,545	235,510	151,769

Dated: May 24, 2022.

Diane Kreinbrink,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2022-11510 Filed 5-27-22; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Generic Clearance for Application Information for Fellowships, Internships, Training Programs, and Specialty Positions, National Cancer Institute (NCI)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open

for Public Comments" or by using the search function.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Diane Kreinbrink, Office of Management Policy and Compliance, National Cancer Institute, 9609 Medical Center Drive, Rockville, Maryland 20892 or call non-toll-free number (240) 276-7283 or email your request, including your address to: diane.kreinbrink@nih.gov. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on March 17, 2022 (Vol. 87, No. 52, Page 15257) and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute (NCI), National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

Proposed Collection: Generic Clearance for Application Information from Fellows, Interns, and Trainees, 0925-0761, Exp., Date 07/31/2022, EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This generic information collection request is to support the science and research in a multidisciplinary environment at the National Cancer Institute (NCI), a part of the National Institutes of Health. Applicants may possess a variety of degrees including, but not limited to, high school, post-baccalaureate, graduate, postdoctoral, Registered Nurse, and Doctor of Medicine (MD). Potential applicants may apply for cancer-related positions by submitting applications, resumes, curriculum vitae (CV), reference letters, letters of intent and interest, and other related documentation directly to the Divisions, Offices, and Centers. This information is necessary to evaluate the eligibility, merits, and quality of potential candidates and will also assist in matching potential candidates to various training and internship programs, and specialty positions. The information is for internal use to make decisions about candidates invited to visit and attend NCI fellowships, internships, training opportunities, and apply for specialized staff and faculty positions.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden is 7,500 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hours
Individuals (Applicants)	3,000	1	60/60	3,000
Individuals (Professional References)	9,000	1	30/60	4,500
Totals	12,000	7,500

Dated: May 24, 2022.

Diane Kreinbrink,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2022-11506 Filed 5-27-22; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which