

collection requirement concerning the Make-or-Buy Program.

DATES: Submit comments on or before May 9, 2016.

ADDRESSES: Submit comments identified by Information Collection 9000–0078, Make-or-Buy Program, by any of the following methods:

- *Regulations.gov:* <http://www.regulations.gov>. Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link “Submit a Comment” that corresponds with “Information Collection 9000–0078, Make-or-Buy Program”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0078, Make-or-Buy Program” on your attached document.

- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0078, Make-or-Buy Program.

Instructions: Please submit comments only and cite Information Collection 9000–0078, Make-or-Buy Program, in all correspondence related to this collection. Comments received generally will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check www.regulations.gov, approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

FOR FURTHER INFORMATION CONTACT: Mr. Edward Loeb, Procurement Analyst, Office of Acquisition Policy, GSA, 202–501–0650 or via email at edward.loeb@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose

Price, performance, and/or implementation of socio-economic policies may be affected by make-or-buy decisions under certain Government prime contracts. Accordingly, FAR 15.407–2, Make-or-Buy Programs:

(i) Sets forth circumstances under which a Government contractor must submit for approval by the contracting officer a make-or-buy program, *i.e.*, a written plan identifying major items to be produced or work efforts to be performed in the prime contractor’s facilities and those to be subcontracted;

(ii) Provides guidance to contracting officers concerning the review and

approval of the make-or-buy programs; and

(iii) Prescribes the contract clause at FAR 52.215–9, Changes or Additions to Make-or-Buy Programs, which specifies the circumstances under which the contractor is required to submit for the contracting officer’s advance approval a notification and justification of any proposed change in the approved make-or-buy program.

The information is used to assure the lowest overall cost to the Government for required supplies and services.

B. Annual Reporting Burden

Respondents: 150.

Responses Per Respondent: 3.

Total Responses: 450.

Hours per Response: 8.

Total Burden Hours: 3,600.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the Federal Acquisition Regulation (FAR), and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0078, Make-or-Buy Program, in all correspondence.

Dated: March 3, 2016.

Lorin S. Curit,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2016–05204 Filed 3–8–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–16CM]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.*, permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to omb@cdc.gov. Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

The Cooperative Re-engagement Controlled Trial (CoRECT)—New—National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC), National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), Division of HIV/AIDS Prevention (DHAP) requests a new three year OMB approval for information collection for a new research study entitled “The Cooperative Re-engagement Controlled Trial (CoRECT)”. The purpose of the study is to evaluate a combined health department and clinic intervention to improve engagement in HIV care. Increasing the number of people living with HIV who access HIV care and achieve viral load suppression addresses one of the priorities of the National HIV/AIDS Strategy. The data collection is authorized under the Section 301 of the Public Health Service Act (42 U.S.C. 241).

The CoRECT Study data collection is comprised of six core components: 1. Electronic clinic data abstraction (Electronic Medical Record (EMR) abstraction will be conducted by project clinic staff at each project clinic to develop the clinic-based “Out of Care” list); 2. electronic surveillance data abstraction (Electronic surveillance data abstraction will be conducted by project health department staff at each health department to develop the health department based “Out of Care” list); 3. a “Barriers to Care” survey (These surveys will provide information regarding barriers to accessing healthcare (e.g., transportation, financial assistance, housing, substance abuse services, etc.)); 4. a “Standard of Care” survey (Investigators will administer this survey to clinic managers, at baseline and every six months during

the study period to assess how the delivery of health services has evolved over time); 5. a Participant Eligibility Disposition form (a listing of potential out-of-care patients will be reviewed to determine those who appear to be out-of-care, as determined by study eligibility, versus those who meet criteria for exclusion); and 6. a Case Conference form (project health department staff will determine if potentially eligible patients met criteria for inclusion in the study and if so randomization will occur). Prospective data collection will provide information about participant’s baseline characteristics including sex, race/ethnicity, HIV exposure risk category, CD4 and viral load test results, date of first clinic visit, and insurance status.

HIV antiretroviral therapy (ART) can durably suppress the plasma HIV viral load, which improves individual survival and dramatically reduces further HIV transmission. Increasing the number of people living with HIV who access HIV care and achieve viral load suppression is a priority of the National HIV/AIDS Strategy. Within the continuum of HIV care in the United States, improvements in linkage to and retention in effective care provide the greatest opportunity to improve rates of HIV viral suppression. It is estimated that of the 1.2 million persons living with HIV in 2011, only 40% were engaged in HIV medical care and only 30% achieved viral suppression.

HIV clinical trials with enhanced case management have demonstrated that interventions provided by the health department can improve linkage to HIV care and interventions provided by the clinic can improve retention in HIV

care. Although linkage to care has improved in many health department jurisdictions, being linked to care is not enough. There is a need to ensure that: (i) People diagnosed with HIV and linked to care are engaging medical care (i.e., attending their enrollment appointment and returning for follow-up medical appointments); and (ii) people who have disengaged from HIV care (i.e., have missed medical appointments and have not been seen in clinic for more than 6 months) are able to efficiently re-engage in care. There have been no randomized controlled studies using a Data-to-Care approach to identify and re-engage out of care persons. Controlled studies such as the CoRECT study are critical to determine the effectiveness of HIV prevention interventions.

The CoRECT study is a randomized controlled trial that seeks to establish a data-sharing partnership between health departments and HIV care clinical providers to identify HIV-infected persons who are out of care and evaluate an intervention that aims to have randomized participants: (a) Link to an HIV clinic; (b) remain in HIV medical care; (c) achieve HIV viral load suppression within 12 months; and (d) achieve durable HIV viral load suppression over 18 months.

The study is funded by CDC through cooperative agreements with the Connecticut State Department of Public Health (in collaboration with Yale University School of Medicine), the Massachusetts State Department of Public Health, and the Philadelphia Department of Public Health. The total burden hours are 1,731.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
CoRECT Study Coordinator	Electronic transmittal of surveillance variables.	3	4	1
Clinic Data Manager	Electronic transmittal of clinical variables	46	4	1
CoRECT Study Participants	Barriers to Care Survey	1,200	1	30/60
Clinical Nurse Coordinator	Standard of Care Survey	46	2	45/60
Clinic Data Manager	Case Conference Session	46	12	1
CoRECT study Coordinator (health department).	Case Conference Session	3	12	1
CoRECT Study Coordinator (health department).	Participant Eligibility Disposition form	3	12	1
Clinic data manager	Cost analysis form—baseline	46	1	1
CoRECT Study Coordinator	Start-up cost analysis form—Health department.	3	1	1
Clinic Data Manager	Start-up Cost Analysis form—Clinic	46	1	1
CoRECT Study Coordinator	Annual Costs Analysis form—Health department.	3	2	1.5
Clinic Data Manager	Annual Costs Analysis form—Clinic	46	2	1.5

Leroy A. Richardson,
Chief, Information Collection Review Office,
Office of Scientific Integrity, Office of the
Associate Director for Science, Office of the
Director, Centers for Disease Control and
Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces a meeting for the initial review of applications in response to Funding Opportunity Announcement (FOA) DD 16-001, Centers for Autism and Developmental Disabilities Research and Epidemiology (CADDRE): Study to Explore Early Development (SEED) 3.

Time and Date: 10:00 a.m.–6:00 p.m., EDT, April 7, 2016 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to “FOA DD16-001, Centers for Autism and Developmental Disabilities Research and Epidemiology (CADDRE): Study to Explore Early Development (SEED) 3”.

Contact Person for More Information: Jaya Raman Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway, Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488-6511, kva5@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

Catherine Ramadei,
Acting Director, Management Analysis and
Services Office, Centers for Disease Control
and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC) announces a meeting for the initial review of applications in response to Funding Opportunity Announcement (FOA), DP 16-002, Michigan Lupus Epidemiology and Surveillance (MILES) Program Longitudinal Cohort Study.

Time and Date: 11:00 a.m.–6:00 p.m., EDT, April 5, 2016 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to “FOA DP16-002, Michigan Lupus Epidemiology and Surveillance (MILES) Program Longitudinal Cohort Study”.

Contact Person for More Information: Jaya Raman Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway, Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488-6511, kva5@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Catherine Ramadei,
Acting Director, Management Analysis and
Services Office, Centers for Disease Control
and Prevention.

[FR Doc. 2016-05267 Filed 3-8-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, Office of Infectious Diseases (BSC, OID)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC), announces the following meeting of the aforementioned committee:

Time and Date: 2:00–4:00 p.m., EDT, March 28, 2016.

Place: Teleconference.

Status: The meeting is open to the public; the toll free dial in number is 1-877-951-7311 with a pass code of 7972741.

Purpose: The BSC, OID, provides advice and guidance to the Secretary, Department of Health and Human Services; the Director, CDC; the Director, OID; and the Directors of the National Center for Immunization and Respiratory Diseases, the National Center for Emerging and Zoonotic Infectious Diseases, and the National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention, CDC, in the following areas: Strategies, goals, and priorities for programs; research within the national centers; and overall strategic direction and focus of OID and the national centers.

Matters for Discussion: Topics to be discussed during the teleconference include administrative/budget issues, current emerging infectious disease outbreak responses, and reports from recent program meetings.

The agenda and any supplemental material will be available at www.cdc.gov/oid/BSC/meetingschedule.html after March 20.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Robin Moseley, M.A.T., Designated Federal Officer, OID, CDC, 1600 Clifton Road NE., Mailstop D10, Atlanta, Georgia 30333, Telephone: (404) 639-4461.

The Director, Management Analysis and Services Office has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and