Additionally submit a copy to GSA 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 9000–0077, Quality Assurance Requirements”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0077, Quality Assurance Requirements” on your attached document.
- Mail: General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0077, Quality Assurance Requirements.

Instructions: Please submit comments only and cite Information Collection 9000–0077, Quality Assurance Requirements, 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. Curtis E. Glover, Sr., Procurement Analyst, Contract Policy Division, at 202–501–1448 or email curtis.glover@gsa.gov.

SUPPLEMENTARY INFORMATION:

A. Purpose
Supplies and services acquired under Government contracts must conform to the contract’s quality and quantity requirements. FAR Part 46 prescribes inspection, acceptance, warranty, and other measures associated with quality requirements. Standard clauses related to inspection require the contractor to provide and maintain an inspection system that is acceptable to the Government; gives the Government the right to make inspections and test while work is in process; and requires the contractor to keep complete, and make available to the Government, records of its inspection work. A notice was published in the Federal Register at 81 FR 11794 on March 7, 2016. No comments were received.

B. Annual Reporting Burden
Respondents: 138,292.

Responses per Respondent: 1.03226.
Total Responses: 142,753.
Hours per Response:.83511.
Total Burden hours: 119,214.
Affected Public: Businesses or other for-profit and not-for-profit institutions.
Frequency: On occasion.

C. Public Comments
Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the 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. 9000–0077, Quality Assurance Requirements, in all correspondence.

Dated: May 17, 2016.
Lorin S. Curit,
Director, Federal Acquisition Policy, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[CFR Doc. 2016–12002 Filed 5–20–16; 8:45 am]
BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–16BX]

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 ombr@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

Monitoring and Reporting System (MRS) for Rape Prevention and Education (RPE) Program Awardees—New—National Center for Injury Prevention and Control NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Sexual violence (SV) is a major public health problem, but it is preventable. According to CDC’s National Intimate Partner and Sexual Violence Survey (NISVS), nearly 1 in 5 women and 1 in 71 men in the U.S. have been raped during their lifetime, and nearly 1 in 2 women and 1 in 5 men have experienced severe SV victimization other than rape at some point in their lives. The majority of victimization starts early in life with approximately 80% of female victims experiencing their first rape before the age of 25, and almost half experiencing their first rape before the age of 18.

CDC’s RPE Program is a national initiative that addresses SV through cooperative agreement funding and technical assistance to health
The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the proposed National Health and Nutrition Examination Survey (NHANES) Longitudinal Study—Feasibility Component. This project will provide a logistical test of proposed survey procedures along with contact, interview, and examination rates for a sample of previously examined NHANES participants. The information obtained will be used to determine the feasibility of conducting future follow-up surveys.

DATES: Written comments must be received on or before July 22, 2016.

APPLICATION: You may submit comments, identified by Docket No. CDC–2016–0043 by any of the following methods:

- Federal eRulemaking Portal: Regulations.gov. Follow the instructions for submitting comments.
- Mail: Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

FOR FURTHER INFORMATION CONTACT: Leroy A. Richardson, the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: omb@cdc.gov.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also