APPENDIX 2: 60-DAY FEDERAL REGISTER NOTICE

Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to the Supplemental Form to the Financial Status Report for all AoA Title III Grantees.

DATES: Submit written or electronic comments on the collection of information by July 23, 2007.

ADDRESSES: Submit electronic comments on the collection of information to: *Stephen.Daniels@aoa.hhs.gov.*

Submit written comments on the collection of information to Administration on Aging, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Stephen Daniels, Director of Grants Management, Administration on Aging, Washington, DC 20201.

SUPPLEMENTARY INFORMATION: Under the 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. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, AoA is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, AoA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of AoA's functions, including whether the information will have practical utility; (2) the accuracy of AoA'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 respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology. The Supplemental form to the Financial Status Report for all AoA Title III Grantees provides an understanding of how projects funded by the Older Americans Act are being administered by grantees, in conformance with legislative requirements, pertinent Federal regulations and other applicable instructions and guidelines issued by Administration on Aging (AoA). This information will be used for Federal oversight of Title III Projects. AoA estimates the burden of this collection of information as follows: 56 State Agencies on Aging respond semiannually which should be an average burden of 1 hour per State agency per submission.

Dated: May 21, 2007.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. E7–10075 Filed 5–23–07; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–07–07BB]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov.*

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Testing of Sexual Violence Definitions and Recommended Data Elements in Three Different Racial/Ethnic Minority Communities -New-National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC). The data collection methodology will be conducted in two phases. The first phase consists of 36 in-person cognitive interviews conducted with women of African American, Hispanic, or American Indian descent. To assess the effectiveness and appropriateness of questions in the sexual violence survey instrument, we will conduct a series of 12 cognitive interviews with adult women from each of these minority groups (for a total of 36 interviews). Cognitive interviewing offers a structured methodology for ascertaining whether the respondent has understood the questions in the way the researchers intend them to be understood, and to assess the ability of the respondents to provide meaningful, accurate, and honest information. A secondary purpose is to make sure that issues pertinent to the research goals are covered adequately.

The second phase of data collection (''main data collection'') will entail 200 in-person interviews with women in each of the minority groups to develop an estimate of sexual violence prevalence within these three communities and describe the characteristics of sexual violence within each community.

Background and Brief Description

This study examines the definitions of sexual violence in three racial/ethnic minority communities: African-American, American Indian, and Hispanic. The purpose of this project is to develop an understanding of sexual violence in these communities. The developed survey will include the following: Projecting estimates of sexual violence; describing the type of sexual violence; and developing a strategy that will increase awareness of sexual violence in minority communities. In addition, this project will establish the groundwork for similar future research.

This research builds on findings from the National Violence against Women Survey (NVAW) (OMB No. 1121–0188; expiration 5/1998), a joint research effort funded by the (CDC) and National Institute of Justice (NIJ) that explored

the occurrence of violence against women through a survey administered to a national sample of adult females and males. The proposed study will expand on this work by clarifying definitions, expanding the categories of sexual violence, and examining the sexual violence event.

This study will focus on women and will occur in two phases: Cognitive and in-person interviews. In each of the three communities, in-depth cognitive interviews will be conducted with 12 adult women, for a total of 36 cognitive interviews. However, a total of 66 individuals will be screened. Respondents will be identified through agencies working with victims of sexual violence. Participants will be interviewed (in either English or Spanish) at the referral agency. The primary purpose of this interview is to assess the questions for the next phase of the study.

In the next phase, researchers will conduct face-to-face interviews with approximately 200 women in each of the three minority communities for a total of 600 women. However, a total of 701 individuals will be screened. Female respondents who are 18 years old will be selected randomly from the communities. Letters will be mailed to each household in the sample. These households will be contacted at a later date in order to collect eligibility information and to randomly select an individual. Participants will complete a 45 minute interview.

There are no costs to respondents except for their time to participate in the interview.

ESTIMATE OF ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Phase One: Screening for Cognitive Interview	66	1	3/60	3
Phase One: Cognitive Interview	36	1	2	72
Phase Two: Screening for Main Survey	701	1	5/60	58
Phase Two: Main Survey	600	1	45/60	450
Total				583

Dated: May 18, 2007.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–10027 Filed 5–23–07; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–07–0199]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Maryam I. Daneshvar, CDC Acting Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to *omb@cdc.gov*.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Written comments should be received within 60 days of this notice.

Proposed Project

Importation of Etiologic Agents, Hosts, and Vectors of Human Disease (42 CFR 71.54)—(OMB Control No. 0920–0199)—Extension—Office of the Director (OD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Foreign Quarantine Regulations (42 CFR part 71) set forth provisions to prevent the introduction, transmission, and spread of communicable disease from foreign countries into the United States. Subpart F—Importations—contains provisions for importation of etiologic agents, hosts, and vectors (42 CFR 71.54), requiring persons that import or distribute after importation these materials to obtain a permit issued by the CDC. This request is for the information collection requirements contained in 42 CFR 71.54 for issuance of permits by CDC to importers or distributors after importation of etiologic agents, hosts, or vectors of human disease.

CDC is requesting continued OMB approval to collect this information through the use of two separate forms. *These forms are:* (1) Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease and (2) Application for Permit to Import or Transport Live Bats.

The Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease will be used by laboratory facilities, such as those operated by government agencies, universities, research institutions, and zoologic exhibitions, and also by importers of nonhuman primate trophy materials, such as hunters or taxidermists, to request permits for the importation and subsequent distribution after importation of etiologic agents, hosts, or vectors of human disease. The Application for Permit to Import or Transport Etiologic Agents, Hosts, or Vectors of Human Disease requests applicant and sender contact information; description of material for importation; facility isolation and containment information; and personnel qualifications. Estimated average time to complete this form is 20 minutes.

The Application for Permit to Import or Transport Live Bats will be used by laboratory facilities such as those operated by government agencies, universities, research institutions, and zoologic exhibitions entities to request importation and subsequent distribution after importation of live bats. The