Denning, Office of Governmentwide Policy, Office of Travel, Transportation, and Asset Management, at (202) 208–7642, or by e-mail at *http://www.gsa.gov/perdiemquestions.* Please cite Notice of Per Diem Bulletin 10–01.

SUPPLEMENTARY INFORMATION:

A. Background

After an analysis of current data, GSA has determined that current lodging rates for certain localities do not adequately reflect the lodging economics in those areas. GSA used the same lodging rate setting methodology for establishing the FY 2010 per diem rates as when establishing the FY 2009 rates.

B. Change in Standard Procedure

GSA issues/publishes the CONUS per diem rates, formerly published in Appendix A to 41 CFR Chapter 301, solely on the Internet at *http://www.gsa.gov/perdiem.* This process, implemented in 2003, ensures more timely changes in per diem rates established by GSA for Federal employees on official travel within CONUS. Notices published periodically in the **Federal Register**, such as this one, now constitute the only notification of revisions in CONUS per diem rates to agencies.

Dated: August 20, 2009.

Becky Rhodes,

Deputy Associate Administrator.

[FR Doc. E9–20504 Filed 8–21–09; 4:15 pm]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-09–0762]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Formative Research to Inform an HIV Testing Social Marketing Campaign for African American Men Who Have Sex with Men (MSM), formally known as Formative Research to Inform an HIV Testing Social Marketing Campaign for African American Heterosexual Men [OMB No. 0920–0762] [exp. 01/31/2011]—Revision—National Center for HIV/AIDS, Viral Hepatitis, Sexually Transmitted Diseases, and Tuberculosis Elimination Programs (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of the proposed revised study is to conduct formative research for the development of an HIV testing social marketing campaign for African American MSM, a CDC-sponsored social marketing campaign aimed at increasing HIV testing rates among young, African American MSM. The study entails conducting interviews with a sample of African American MSM, ages 18 to 44 to: (1) Explore participants' knowledge, attitudes and beliefs about HIV and HIV testing to inform the development of campaign messages; (2) identify the most motivating approach, supporting data, and key messages for materials development; (3) test creative concepts, potential campaign themes, logos and names; and (4) test creative materials developed based on the findings from the previous phases of the research. Findings from this study will be used by CDC and its partners to inform current and future program activities. Changes to the previous approved data collection consist of a change in the target audience from African American heterosexual men to African American Men who have sex with men. Instead of a combination of interviews and focus groups, now only interviews will be conducted.

A total of 288 participants will be screened for eligibility in 12 cities with high incidence and prevalence of HIV. Of the participants screened, 144 men will complete individual interviews and a short paper and pencil survey. Appropriate consent processes will be used to obtain verbal consent at the screening and interview phases of this study. The Institutional Review Board at CDC has approved the revised study. There are no costs to the respondents other than their time. The total annualized burden hours are 228.

ESTIMATE OF ANNUALIZED BURDEN TABLE

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
African American MSM	Screener	288	1	10/60
	Interview	144	1	1
	Paper and Pencil Survey	144	1	15/60

Dated: August 19, 2009.

Maryam I. Daneshvar,

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

[FR Doc. E9–20374 Filed 8–24–09; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; NIH Intramural Research Training Program Applications

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Director (OD), 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. This proposed information collection was previously published in the **Federal Register** on June 16, 2009 (Volume 74, Number 114, pages 28501–28502) 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 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.

Proposed Collection: Title: NIH Intramural Research Training Program Applications. *Type of Information Collection Request:* Extension of a currently approved collection. *Need and Use of Information Collection:* The proposed information collection activity is necessary in order to determine the eligibility and quality of potential awardees for traineeships in ten (10) NIH intramural research training programs. *Frequency of Response:* On occasion. *Affected Public:* Individuals seeking intramural training opportunities and references for these individuals. *Type of Respondents:* Postdoctoral, predoctoral, postbaccalaureate, technical, clinical, and student applicants. The annual reporting burden is as follows: *Estimated Number of Respondents:* 67,631; *Estimated Number of Responses per Respondent:* 1.0506; *Average Burden Hours Per Response:* 0.9545; and *Estimated Total Annual Burden Hours Requested:* 67,825. The annualized cost to respondents is estimated at $2,033,085. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on 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.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, *Attention:* Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Marilyn Tuttleman, M.S., Chief, Project Clearance Branch, Office of Policy for Extramural Research Administration (OPERA), OER, OD, NIH, One Rockledge Center, Room 3509, 6705 Rockledge Drive MSC 7974, Bethesda, MD 20892–7983, or call non-toll-free number 301–594–7949 or e-mail your request, including your address to: *mtuttleman@mail.nih.gov.*

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

Dated: August 17, 2009.

Steven Alves,

Project Officer, OD, OIR, OITE, National Institutes of Health.

[FR Doc. E9–20439 Filed 8–24–09; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0380]

Agency Information Collection Activities; Proposed Collection; Comment Request; Product Jurisdiction: Assignment of Agency Component for Review of Premarket Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction 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 procedure by which an applicant may obtain an assignment or designation determination for combination products.

DATES: Submit written or electronic comments on the collection of information by October 26, 2009.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, *JonnaLynn.Capezzuto@fda.hhs.gov*, 301–796–3794.

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 requests 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, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA'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.