necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *Sherette.funncoleman@hhs.gov*, or call the Reports Clearance Office on (202) 690–5683. Send written comments and recommendations for the proposed information collections within 30 days of this notice directly to the OMB Desk Officer; via fax to 202–395–6974.

Proposed Project: Evaluation of HIV Prevention Program for Women—OMB No. 0990–New—Office on Women's Health (OWH).

Abstract: OWH is seeking a new clearance to conduct data collection activities associated with the three-year evaluation of HIV/AIDS prevention programs at minority institutions for college women. The evaluation will assess the effectiveness of gender-centered HIV prevention models employed to address the needs of college women attending six (6) Historically Black Colleges and Universities, four (4) Hispanic Serving Institutions, and two (2) Tribal Colleges and Universities. Evaluation data will assess the extent to which gender sensitive interventions employed by these institutions impact HIV/AIDS related knowledge, attitudes and risk behaviors. Results of the evaluation will identify the prevention outcomes for college women and the larger campus population. The program goals are to (1) identify effective methods to educate and increase awareness for prevention of HIV/AIDS and STD's; (2) develop capacity for young, minority women to address prevention education on campus; (3) establish partnerships and student organizations to increase health education, risk reduction, counseling, HIV/STD testing and (4) to ensure that the education is culturally and linguistically appropriate for young, minority women. HIV intervention recipients will complete a survey at pretest, post-test and follow-up that assesses their HIV/AIDS knowledge, risk behaviors and gender specific attitudes. Student data will be submitted on a quarterly basis. Peer facilitators/health educators will participate in a yearly interview and project implementation staff will participate in individual interviews twice during each program year that assesses their experiences with the program.

ESTIMATED ANNUALIZED BURDEN TABLE

Forms (if necessary)	Type of respondent	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Prevention Education: Pre-test Questionnaire	Student	660	1	20/60	220
Prevention Education: Post-test Questionnaire	Student	660	1	20/60	220
Prevention Education: Follow-up	Student	660	1	20/60	220
Process Interview: Program Directors	Program Directors	14	2	1.5	42
Process Interview: Program Staff	Program Staff	12	2	45/60	18
	Program Staff	12	55	15/60	165
Process Interview: Peer Educators	Peer Educators	50	1	30/60	25
Total					910

Seleda Perryman,

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

[FR Doc. E8–21119 Filed 9–10–08; 8:45 am]

BILLING CODE 4150–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Communities Empowering Youth (CEY) Program Evaluation.

OMB No.: 0970–0335.

Description: This proposed information collection activity is to obtain information from Communities Empowering Youth (CEY) grantee agencies and the faith-based and community organizations working in partnership with them. The CEY evaluation is an important opportunity to examine the outcomes achieved through this component of the Compassion Capital Fund in meeting its objective of improving the capacity of faith-based and community organizations and the partnerships they form to increase positive youth development and address youth violence, gang involvement, and child abuse/neglect. The evaluation will be designed to assess changes and improvements in the structure and functioning of the partnership and the organizational capacity of each participating organization. The purpose of this request will be to revise the approved baseline instruments for follow-up data collection.

Respondents: CEY grantees and the faith-based and community organizations that are a part of the partnership approved under the CEY grant.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response (minutes)	Total burden hours
2009 Follow-up Survey.	464	1	45	348
2010 Follow-up Survey	143	1	45	107.25

Estimated Total Annual Burden Hours: 455.25

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447. Attn: ACF Reports Clearance Officer. E-mail address: *OPREinfocollection@acf.hhs.gov*. All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: September 4, 2008.

Brendan C. Kelly,

OPRE Reports Clearance Officer.

[FR Doc. E8–21007 Filed 9–10–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–N–0132]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; State Petitions for Exemption from Preemption

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''State Petitions for Exemption from Preemption'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

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

SUPPLEMENTARY INFORMATION: In the **Federal Register** of June 16, 2008 (73 FR 34024), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0277. The approval expires on August 31, 2011. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: September 4, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–21020 Filed 9–9–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2006–N–0178] (formerly Docket No. 2006N–0362)

Draft Guidance for Industry and Food and Drug Administration Staff; Class II Special Controls Guidance Document: Absorbable Hemostatic Device; Availability; Reopening of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; reopening of comment period.

SUMMARY: The Food and Drug Administration (FDA) is reopening until October 14, 2008, the comment period for a draft guidance entitled ''Class II Special Controls Guidance Document: Absorbable Hemostatic Device.'' The draft guidance describes a means by which the absorbable hemostatic device may comply with the requirements of special controls if it is reclassified. FDA is reopening the comment period to update comments and to receive any new information. Elsewhere in this issue of the **Federal Register**, FDA is reopening the comment period on a proposed rule to reclassify the absorbable hemostatic device from class III (premarket approval) into class II (special controls).

DATES Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that the agency considers your comments on this draft guidance before it begins work on the final version of the guidance, submit written or electronic comments on the draft guidance October 14, 2008.

ADDRESS: Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.regulations.gov*. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.