and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Label Comprehension Study (U.S.C. 393(d)(2)(C))

FDA issued the ''Draft Guidance for Industry and FDA Staff: Class II Special Controls Guidance Document: Labeling for Male Condoms Made of Natural Rubber Latex'' on November 14, 2005 (70 FR 69156). Section 21 U.S.C. 393(d)(2)(C) of the Federal Food, Drug and Cosmetic Act (the act) states that the Secretary, through the Commissioner, shall be responsible to conduct research relating to * * * devices in carrying out this chapter. In order to evaluate the understandability of the condom labeling language currently on the market and the labeling language proposed in this draft guidance, as well as a future revised version of the labeling, FDA plans to evaluate readers' comprehension of three versions of condom labeling through a label comprehension study.

The proposed label comprehension study will measure current and potential condom consumers' understanding of the current market labeling and the proposed condom labeling in the draft guidance of the retail package, foil and package insert of condom labeling, as well as a future revised version of the labeling. The label comprehension study will follow a sequential design, first testing both the current market labeling (Part A) and the draft labeling in the guidance (Part B) in Stage 1, and then a revised version of the labeling in Stage 2.

FDA will conduct a label comprehension study via a mall intercept/central location intercept methodology with pre-screened participants. The FDA will administer a screening instrument, the Rapid Estimate of Adult Literacy in Medicine (REALM) test, an informed consent, and a questionnaire with approximately 20 questions related to the condom labeling language to a total of 1,200 participants: 400 participants for Part A of Stage 1, 400 participants for Part B of Stage 1, and 400 participants for Stage 2 of the study. Results of the study will be considered in FDA's condom labeling recommendations to provide important risk/benefit and use information associated with condoms in an easily understood language.

In the **Federal Register** of February 16, 2007 (72 FR 7661), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

FDA estimates the burden for this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Activity	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Screening Tool	3,300	1	3,300	.05	165
Stage 1: Part A - REALM test; Informed Consent; Read Labeling; Questionnaire	400	1	400	.45	180
Stage 1: Part B - REALM test; Informed Consent; Read Labeling; Questionnaire	400	1	400	.45	180
Stage 2 - REALM test; Informed Consent; Read Labeling; Questionnaire	400	1	400	.45	180
Total					705

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

This was based on similar types of FDA studies conducted in the past. FDA has conducted both focus group studies and label comprehension studies, where similar participant activities, such as reading the labeling, taking the REALM test, signing the informed consent, and answering questions on a self-administered questionnaire took place. In order to achieve the 1,200 participants for the condom label comprehension study, FDA estimates screening 3,300 to achieve 1,200 interviews.

Dated: June 8, 2007.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E7–11528 Filed 6–14–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub.

L. 104–13), the Health Resources and Services Administration (HRSA) will publish periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans, call the HRSA Reports Clearance Officer on (301) 443–1129.

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 of other forms of information technology.

Proposed Project: The Nursing Education Loan Repayment Program Application (OMB No. 0915–0140)—Revision

This is a request for revision of the Nursing Education Loan Repayment Program (NELRP) application and participant monitoring forms. The NELRP was originally authorized by 42 U.S.C. 297b(h) (section 836(h) of the Public Health Service Act) as amended by Public Law 100–607, November 4, 1988. The NELRP is currently authorized by 42 U.S.C. 297n (section 846 of the Public Health Service Act) as amended by Public Law 107–205, August 1, 2002.

Under the NELRP, registered nurses are offered the opportunity to enter into a contractual agreement with the Secretary to receive loan repayment for up to 85 percent of their qualifying educational loan balance as follows: 30 percent each year for the first 2 years and 25 percent for the third year. In exchange, the nurses agree to serve full-time as a registered nurse for 2 or 3 years at a health care facility with a critical shortage of nurses.

NELRP requires the following information:

1. Applicants must provide information on their nursing education, employment, and proposed service site;

2. Applicants must provide information on their outstanding nursing educational loans;

3. Applicants must provide banking information from their financial institution; and

4. Employers must provide information on the health care facility and on the employment status of applicants and participants.

Form	Number of respondents	Responses per respondents	Total responses	Hours per response	Total burden hours
Estimates of Annualized Hour Burden are as Follows for Applicants:					
NELRP Application	5,000	1	5,000	1.5	7,500
Loan Verification Form	5,000	3	15,000	1	15,000
Applicant Employment Verification Form	5,000	1	5,000	.5	2,500
Payment Information Form	5,000	1	5,000	1	5,000
Application Checklist	5,000	1	5,000	.5	2,500
Pre-Award Confirmation Checklist	600	1	600	.25	150
Total	5,000	.....	35,600	.....	32,650
Estimates of Annualized Hour Burden are as Follows for Participants:					
Participant Semi-Annual Employment Verification Form	1,300	2	2,600	.5	1,300
Total	1,300	2	2,600	.5	1,300

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 10–33 Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Written comments should be received with 60 days of this notice.

Dated: June 11, 2007.

Caroline Lewis,

Associate Administrator for Management.

[FR Doc. E7–11557 Filed 6–14–07; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Commission of Childhood Vaccines; Request for Nominations for Voting Members

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: The Health Resources and Services Administration (HRSA) is requesting nominations to fill three vacancies on the Advisory Commission on Childhood Vaccines (ACCV). The ACCV was established by Title XXI of the Public Health Service Act (the Act), as enacted by Public Law (Pub. L.) 99–660 and as subsequently amended, and advises the Secretary of Health and Human Services (the Secretary) on issues related to implementation of the National Vaccine Injury Compensation Program (VICP).

DATES: The agency must receive nominations on or before July 16, 2007.

ADDRESSES: All nominations are to be submitted to the Director, Division of Vaccine Injury Compensation, Healthcare Systems Bureau (HSB), HRSA, Parklawn Building, Room 11C–26, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: Ms. Delia Jones, Principal Staff Liaison, Policy Analysis Branch, Division of Vaccine Injury Compensation, HSB, HRSA at (301) 443–6593 or e-mail: *djones2@hrsa.gov.*

SUPPLEMENTARY INFORMATION: Under the authorities that established the ACCV, the Federal Advisory Committee Act of October 6, 1972 (Pub. L. 92–463) and