The agenda, the questions to be addressed by the committee, and a current list of committee members will be available at the meeting location on the day of the meeting.

Transcripts of the open portion of the meeting may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, approximately 15 working days after the meeting, at a cost of 10 cents per page. The transcript may be viewed at the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, approximately 15 working days after the meeting, between the hours of 9 a.m. and 4 p.m., Monday through Friday. Summary minutes of the open portion of the meeting may be requested in writing from the Freedom of Information Office (address above) beginning approximately 90 days after the meeting.

This notice is issued under section 10(a)(1) and (2) of the Federal Advisory Committee Act (5 U.S.C. app. 2), and FDA's regulations (21 CFR part 14) on advisory committees.

Dated: April 24, 1996.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 96–10780 Filed 4–26–96; 2:36 pm]

BILLING CODE 4160–01–F

[Docket No. 81N–033P]

Background Document for the Dental Drug Products Panel Plaque Subcommittee (Nonprescription Drugs) of the Medical Devices Advisory Committee; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a background document for the meeting of the Dental Drug Products Panel Plaque Subcommittee of the Medical Devices Advisory Committee (the subcommittee). This meeting is announced elsewhere in this issue of the Federal Register, and it is scheduled for June 6 and 7, 1996. This background document is being taken to ensure that all interested parties are aware of the subcommittee's concern regarding the relationship, if any, of alcohol-containing mouthwashes and oral cancer and the development of studies to investigate the relationship. This relationship will be the subject of the subcommittee's discussion on June 6, 1996.

DATES: Written comments or data should be submitted by May 10, 1996, in order to be considered for discussion at the June 6, 1996, subcommittee meeting.

ADDRESSES: Single copies of the background briefing document may be requested in writing from the Freedom of Information Staff (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, at a cost of 10 cents per page. Requests should be identified with the docket number found in brackets in the heading of this document. The background briefing document is available for public examination at the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

Comments and data should be identified with the docket number listed above. Individuals or groups wishing to submit data or comments relevant to alcohol-containing mouthwashes should send them to the Dockets Management Branch (HFA–305), 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. Three copies of written comments should be submitted, except that individuals may submit one copy. The comments and data received are available for public examination at the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

FOR FURTHER INFORMATION CONTACT: Jeanne L. Rippere or Stephanie Mason, Center for Drug Evaluation and Research (HFD–560), Food and Drug Administration, 1600 Rockville Pike, Rockville, MD 20857, 301–827–2244.

SUPPLEMENTARY INFORMATION: Elsewhere in this issue of the Federal Register, FDA announced that a meeting of the Dental Drug Products Panel Plaque Subcommittee will be held on June 6 and 7, 1996. The purpose of the meeting scheduled for June 6, 1996, is to continue the subcommittee's discussion concerning the alcohol content of oral health care mouthwash drug products begun at its meeting of June 28 and 29, 1994. After evaluating the available data, the subcommittee concluded that it should meet in a workshop environment with representatives of the National Cancer Institute, the National Institute of Dental Research, other professional groups, the agency, and industry to address any new information regarding a causal relationship between alcohol-containing mouthwashes and oral cancer. The subcommittee recommended that this workshop should address the development of sound scientific studies to determine the relationship, if any, between alcohol-containing mouthwash products and cancer of the oral cavity.

FDA has established a docket number (81N–033P) as a public record of the comments, views, and other information submitted to the agency from interested persons and organizations regarding alcohol in oral health care mouthwash drug products. After publication of the subcommittee's report, this docket will be the repository of all data and information collected by the agency for the over-the-counter (OTC) antiplaque/antigingivitis drug review, but currently it will contain only those comments and data that are not confidential under the OTC drug review. (See the request for data and information on dental and oral health care drug products for antiplaque use published in the Federal Register of September 19, 1990 (55 FR 38560 at 38562).) Copies of the background briefing documents have been placed in this docket and may be seen in the Dockets Management Branch (address above) or obtained from the agency's Freedom of Information Staff (address above). Copies of the background briefing document will also be available at the committee meeting.

Dated: April 23, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–10781 Filed 4–26–96; 2:36 pm]

BILLING CODE 4160–01–F

Health Care Financing Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, HHS.

In compliance with the Paperwork Reduction Act (44 U.S.C. 3501 *et seq.*), this notice is publishing the following summaries of proposed collections for public comment. The title, description, and respondent description of the information collection are shown below with an estimate of the annual reporting and recordkeeping burden. Included in the estimate is the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The 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.

1. *Type of Information Collection Request:* New; *Title of Information Collection:* National Payer Identifier (PAYER–ID); *Form No.*: HCFA–856; *Use:* The PAYER-ID will allow payers of health care claims to be identified by a unique numeric identifier. PAYER–ID numbers will be assigned, but not limited to the following groups: Medicare, Medicaid, VA, public health service, large employers and unions, HMOs, large insurers, etc.; *Frequency:* One time (Reporting); *Affected Public:* Business or other for profit, Not for profit institutions, Federal Government, State, local or tribal government; *Number of Respondents:* 85,000; *Total Annual Responses:* 8,500; *Total Annual Hours Requested* 85,000.

To request copies of the proposed paperwork collection referenced above, call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections should be sent within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Financial and Human Resources, Management Planning and Analysis Staff, Attention: Zaneta Davis, 7500 Security Boulevard, Room C2–26–17, Baltimore, Maryland 21244–1850.

Dated: April 22, 1996.

Kathleen B. Larson,

Director, Management Planning and Analysis Staff.

[FR Doc. 96–10571 Filed 4–29–96; 8:45 am]

BILLING CODE 4120–03–P

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

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 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 and instruments, 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 or other forms of information technology.

Proposed Projects

Ryan White Comprehensive AIDS Resources and Emergency Act of 1990 Women's Initiatives (WIN)—New

The Health Resources and Services Administration's Maternal and Child Health Bureau proposes to collect information about HIV-related services provided to women of child-bearing age and their children. Information will be collected annually from eight sites funded by HRSA under cooperative agreements and their 320 local service providers that are funded under Sections 2618(a) and 2671 of the Public Health Service Act. The eight funded sites will collect the information by telephone from their providers, and forward the data collection forms to a HRSA contractor. There are no plans to collect or transmit the data electronically.

The purpose is to document current care system characteristics and facilitate planning for services to women with HIV and their children. The information will be used within and outside HRSA to inform the administration and Congress about HIV counseling and testing services for pregnant women, services and referral resources for pregnant women with HIV, antiretroviral therapies, and outreach related to perinatal HIV transmission reduction. Annual burden estimates are as follows:

Type of respondent	Number of respondents	Responses per respondent	Burden hours per response	Total burden hours
Providers	320	1	.33	106
Funded Sites	8	40	1.0	320
Total	328	.....	.....	426

Faculty Loan Repayment Program (FLRP) Application (0915–0150)—Extension and Revision

Under the HRSA FLRP program, disadvantaged graduates from certain health professions schools may enter into a contract under which HRSA with the Department of Health and Human Services will make payments on eligible graduate educational loans in exchange for a minimum of two years of service as a full-time faculty member of a health professions school. Applicants must complete an application and provide information on all eligible education loans. Once HRSA has selected the participants, HRSA will request verification from their lenders of loan balances and terms of their outstanding educational loans.

Estimated annual response burden is as follows:

Type of respondent	Number of respondents	Responses per respondent	Hours per response	Total annual hour burden
Applicants	75	1	1	75