Dated: June 27, 1996.

Joseph R. Carter

Acting Associate Director for Management and Operations, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–16946 Filed 7–02–96; 8:45 am]

BILLING CODE 4163–18–P

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Annual Survey of Refugees (ORR–9).

OMB No.: 0970–0033.

Description: The Annual Survey of Refugees collects information on the economic circumstances of a random sample of refugees, Amerasians, and entrants who arrived in the U.S. during the previous five years. The survey focuses on their training, labor force participation, and welfare utilization rates. Data are segmented by region of origin, State of resettlement, and number of months since arrival. From their responses, ORR reports on the economic adjustment of refugees annual deliberations of refugee admissions and funding and by program managers in formulating policies for the direction of the Refugee Resettlement Program.

Respondents: State governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per respondent	Total burden hours
ORR–9	1,800	1	.75	1,350

Estimated Total Annual Burden Hours: 1,350.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the Federal Register. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

Dated: June 26, 1996.

Larry Guerrero,

Director, Office of Information Management Services.

[FR Doc. 96–16969 Filed 7–2–96; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0158]

Agency Information Collection Activities; Proposed Collection; Comment Request

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, Federal agencies are required to publish a notice in the Federal Register concerning each collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a survey of hormone replacement therapy (HRT) among women with a previous diagnosis of endometrial cancer.

DATES: Submit written comments on the collection of information by September 3, 1996.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 (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. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c). To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (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.

Survey of HRT Among Women With A Previous Diagnosis of Endometrial Cancer

Under FDA's statutory authority to conduct and sponsor research (21 U.S.C.

393(b)(2)(C) and 42 U.S.C. 300u–1), FDA and the Fred Hutchinson Cancer Research Center in Seattle, WA, have jointly designed a study involving a written survey to be completed by women with a previous diagnosis of endometrial cancer. The study will evaluate the occurrence of menopausal vasomotor symptoms (''hot flashes'') among these women, and the extent to which they have used HRT, either as therapy for menopausal symptoms or for other reasons. (''Hormone replacement therapy'' means treatment with estrogen alone or with a combination of estrogen and progestogen.)

FDA estimates the burden of this survey as follows:

ESTIMATED ANNUAL REPORTING BURDEN

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
575	1	575	.5	288

There are no capital costs or operating and maintenance costs associated with this collection.

Dated: June 26, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–16884 Filed 7–2–96; 8:45 am]

BILLING CODE 4160–01–F

[Docket No. 96N–0186]

Agency Information Collection Activities: Proposed Collection; Comment Request

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, Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a proposed survey of mammography facilities to assess the impact of proposed regulations required by the Mammography Quality Standards Act of 1992 (the MQSA).

DATES: Submit written comments on the collection of information by September 3, 1996.

ADDRESSES: Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Charity B. Smith, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, rm. 16B–19, Rockville, MD 20857, 301–827–1686.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 (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. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c). To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (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.

FDA Survey of Mammography Facilities

The MQSA of 1992 (Pub. L. 102–539) has resulted in regulatory efforts by FDA to ensure high quality standards for mammography in the United States. In the Federal Register of April 3, 1996 (61 FR 14856), FDA proposed final regulations to implement these standards. Interim regulations are codified at 21 CFR 900. In connection with this rulemaking, FDA proposes to conduct a survey of facilities to determine current operating costs and procedures. The information to be collected from the proposed survey includes general provider characteristics, equipment characteristics, facility operating procedures, personnel qualifications, and costs of compliance with current quality standards. This information is necessary in order to ensure that costs to affected facilities are minimized to the extent consistent with maintenance of high quality mammography services, and that patient access to mammography services is not diminished as a result of agency action.

The proposed survey will be a one-time data collection effort. Surveys will be mailed to 1,000 randomly selected facilities. Facilities will be contacted by telephone in order to respond to any specific issues, or questions that may arise. Responses will not be mandatory, and no facility will be required to respond. All responses will be kept confidential, although a compilation of data that does not reveal facility-specific information will be made available upon request to participants and to the public. A toll-free telephone number will be installed to allow respondents the opportunity to call if specific issues arise.

FDA estimates the burden of this one-time survey as follows:

ESTIMATED ANNUAL REPORTING BURDEN

Burden Element	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours	Total Costs
Initial Contact by Telephone	1,000	1	1,000	0.083	83	$1,385
Compile Requested Information	702	1	702	1.0	702	$11,716