Respondents	No. of respondents	No. of responses/respondent	Avg. burden/response (in hrs.)	Total burden (in hrs.)
Antiviral Prophylaxis Rpt	100	1	0.25	25
Total				421

Dated: March 15, 1996.

Wilma G. Johnson,

Acting Associate Director for Policy Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–6792 Filed 3–20–96; 8:45 am]

BILLING CODE 4163–18–P

[30DAY–08]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review, 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 Office on (404) 639–7090.

The following request has been submitted for review since the last publication date on February 6, 1996:

Proposed Project

Hanford Environmental Dose Reconstruction (HEDR) Project Milk Producers Survey—New—OMB approved the information collections for the ''Hanford Thyroid Disease Full Epidemiology Study'' under OMB No. 0920–0296 to determine the health effects to the public from radioactive releases from the Hanford Nuclear Site Operations during the 1940's and 1950's. A primary component of these releases was radioactive iodine. Consumption of fresh milk from cows that have eaten contaminated vegetation and fresh leafy vegetables and eggs from chickens with access to outdoor vegetation are important pathways of radioactive iodine to the human body which adversely affects the thyroid gland. To estimate the doses to the thyroid that individuals and populations could have received, historical milk cow and chicken feeding and distribution practices must be reconstructed for the downwind area. This information is particularly important for use in this ongoing study and its relation to radiation exposures. Researchers from LTG Associates will collect information from a representative sample of individuals who farmed in 7 counties within the study area during the periods of 1945 and 1951.

Respondents	No. of respondents	No. of responses/respondents	Avg. burden/response (in hrs.)
Contact Potential Sources of Names of farmers	50	1	0.16
Initial Contact of Potential Candidates	1,600	1	0.16
Scheduling Interview	400	1	0.08
Telephone Interview	400	1	2

The total annual burden is 1108. Send comments to Allison Eydt; Human Resources and Housing Branch, New Executive Office Building, Room 10235; Washington, DC 20503.

Wilma G. Johnson,

Acting Associate Director for Policy Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 96–6793 Filed 3–20–96; 8:45 am]

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Food and Drug Administration

[Docket No. 96N–0084]

Agency Emergency Processing Request Under OMB Review

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for emergency processing under the Paperwork Reduction Act of 1995. The purpose of the proposed collection of information is to enable FDA to compile lists of U.S. processors that export certain animal-derived foods to the European Community (EC). These lists must be completed by May 1, 1996, to meet EC trade requirements. To meet the EC deadline, FDA is requesting OMB approval by March 28, 1996.

DATES: Submit written comments by March 28, 1996.

ADDRESSES: Submit written comments to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Desk Officer for FDA. 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: FDA has requested emergency processing of this proposed collection of information under section 3507(j) of the Paperwork Reduction Act of 1995 and 5 CFR 1320.13 because the information is needed to meet the May 1, 1996, EC deadline; the information is essential to the agency's mission; and public harm is reasonably likely to result if normal clearance procedures are followed.

With respect to the following collection of information, comments are invited 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 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