Dated: January 11, 2000.

Barbara M. Williams,

Deputy Standard and Optional, Forms Management Officer.

[FR Doc. 00–1854 Filed 1–25–00; 8:45 am]

BILLING CODE 6820–34–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–00–20]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention is providing opportunity for public comment on proposed data collection projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639–7090.

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 for other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Projects

Continuing Medical Education (CME) Activity Registration Form—(0923–0013)—Extension—The Agency for Toxic Substances and Disease Registry (ATSDR) is mandated pursuant to the 1980 Comprehensive Environmental Response Compensation and Liability Act (CERCLA) and its 1986 Amendments, The Superfund Amendments and Reauthorization Act (SARA), to prevent or mitigate adverse human health effects and diminished quality of life resulting from the exposure to hazardous substances into the environment. As stated in CERCLA, the Administrator of ATSDR is charged to ''assemble, develop as necessary, and distribute to the states, and upon request to medical colleges, physicians, and other health professionals, appropriate educational materials (including short courses) on this topic''.

The development and use of activity registration forms for documenting participation in these activities at these meetings is an integral part of this process. This attendance documentation process is required by the Accreditation Council for Continuing Medical Education (ACCME), the body that authorizes agencies and institutions to award nationally recognized continuing medical education (CME) credit. As a condition of relicensure, physicians in 40 states are required to participate in CME courses. Individual physicians in these states are required to submit the number of hours of CME credit to state boards of professional registration at the time of relicensure. Failure by the physician to provide this information in a timely fashion will result in suspension of professional licensure.

This request is for a 3-year extension of the current OMB approval of uniform CME activity registration forms—one machine entry form and the other manually entered—to serve as the initial step in the development of an attendance documentation system. Other than their time, there will be no cost to the respondents.

Respondents	No. of respondents	No. of responses/ respondent	Avg. burden per response	Total burden
Manual Entry Registration Form	2,000	1	4/60	133
Scantron Registration Form	3,000	1	5/60	250
Total	.....	.....	.....	383

Dated: January 20, 2000.

Nancy Cheal,

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

[FR Doc. 00–1762 Filed 1–25–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0595]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Reporting and Recordkeeping Requirements for Manufacturers, Importers, User Facilities, and Distributors of Medical Devices Under FDAMA

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Submit written comments on the collection of information by February 26, 2000.

ADDRESSES: Submit written comments on the collection of information 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.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has