organizations representing State child support programs.

Agenda

The agenda for these meeting includes a discussion of the issues to be contained in the MCSWG's report to the Secretaries containing recommendations for appropriate measures to address the impediments to the effective enforcement of medical child support as listed above. At the May, 1999, meeting, the MCSWG formed four (4) sub-committees to discuss barriers, issues, options, and recommendations in the interim between full MCSWG meetings. At the next three meetings (August, 1999, October, 1999, and November, 1999), the sub-committees presented their draft recommendations to the full MCSWG for further discussion and consideration. At the January, 2000, meeting, the MCSWG will discuss the recommendations contained in their report to the Secretaries.

Public Participation

Members of the public wishing to present oral statements to the MSCWG should forward their requests to Samara Weinstein, MCSWG Executive Director, as soon as possible and at least four days before the meeting. Such request should be made by telephone, fax machine, or mail, as shown above. Time permitting, the Chairs of the MCSWG will attempt to accommodate all such requests by reserving time for presentations. The order of persons making such presentations will be assigned in the order in which the requests are received. Members of the public are encouraged to limit oral statements to 5 minutes, but extended written statements may be submitted for the record. Members of the public also may submit written statements for distribution to the MCSWG membership and inclusion in the public record without presenting oral statements. Such written statements should be sent to the MCSWG Executive Director, as shown above, by mail or fax at least 5 business days before the meeting.

Minutes of all public meetings and other documents made available to the MCSWG will be available for public inspection and copying at both the DOL and DHHS. At DHHS, these documents will be available at the MCSWG Executive Director's Office, Office of child Support Enforcement (OCSE), Administraton for Children and Families, U.S. Department of Health and Human Services, Aerospace Building, Fourth Floor-East, 370 L'Enfant Promenade, SW, Washington, DC from 8:30 a.m. to 5:30 p.m. Questions regarding the availability of documents from DHHS should be directed to Andrew J. Hagan, OCSE (telephone (202) 401–5375). This is not a toll-free number. Any written comments on the minutes should be directed to Ms. Samara Weinstein, Executive Director of the Working Group, as shown above.

Dated: December 3, 1999.

David Gray Ross,

Commissioner, Office of Child Support Enforcement.

[FR Doc. 99–32095 Filed 12–9–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–0407]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Reclassification Petitions for Medical Devices

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.

DATES: Submit written comments on the collection of information by January 10, 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: Wendy Taylor, 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 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Reclassification Petitions for Medical Devices—21 CFR 860.123 (OMB Control Number 0910–0138—Extension)

FDA has the responsibility, under sections 513(e) and (f), 514(b), 515(b), and 520(l) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360c(e) and (f), 360d(b), 360e(b), and 360j(l)) and part 860 (21 CFR part 860) subpart C, to collect data and information contained in reclassification petitions. The reclassification provisions of the act allow any person to petition for reclassification of a medical device from any one of three classes (I, II, and III) to another class. The reclassification procedures (§ 860.123) requires the submission of sufficient, valid scientific evidence demonstrating that the proposed classification will provide a reasonable assurance of safety and effectiveness of the device for its intended use. The reclassification provisions of the act serve primarily as a vehicle for manufacturers to seek reclassification from a higher to a lower class, thereby reducing the regulatory requirements applicable to a particular device. The reclassification petitions requesting classification from class III to class II or class I, if approved, provide an alternative route to the market in lieu of premarket approval for class III devices.

In the **Federal Register** of September 17, 1999 (64 FR 50516), the agency requested comments on the proposed collections of information. No significant comments were received.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
860.123	11	1	11	500	5,500

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

Based on current trends, FDA anticipates that 11 petitions will be submitted each year. The time required to prepare and submit a reclassification petition, including the time needed to assemble supporting data, averages 500 hours per petition. This average is based upon estimates by FDA administrative and technical staff who are familiar with the requirements for submission of a reclassification petition, have consulted and advised manufacturers on these requirements, and have reviewed the documentation submitted.

In addition, FDA is correcting a document that appeared in the **Federal Register** of Friday, September 17, 1999 (64 FR 50516). On page 50517, in Table 1 of the document, ''860.133'' is corrected to read ''860.123''.

Dated: December 3, 1999.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 99–32000 Filed 12–9–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–2553]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Citizen Petition

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.

DATES: Submit written comments on the collection of information by January 10, 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: Wendy Taylor, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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.

Citizen Petition—21 CFR 10.30 (OMB Control Number 0910–0183)—Extension

The Administrative Procedures Act (5 U.S.C. 553(e)) provides that every agency shall accord any interested person the right to petition for issuance, amendment, or repeal of a rule. Section 10.30 provides that any person may submit to the agency a citizen petition requesting the Commissioner of Food and Drugs to issue, amend, or revoke a regulation or order, or to take or refrain from taking any other form of administrative action. The information is used by the agency to determine the need or desirability of the requested action and also to determine if the submitted information is sufficient to support the action. FDA determines if the submitted information is sufficient to support the action. FDA determines whether or not to grant the petition based on the information submitted. The affected respondents are individuals or households, State or local governments, nonprofit institutions and businesses or other for-profit institutions or groups.

In the **Federal Register** of June 9, 1998 (63 FR 31502), the agency requested comments on the proposed collections of information. No significant comments were received.

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

Dated: December 6, 1999.

William K. Hubbard,

Senior Associate Commissioner for Policy, Planning, and Legislation.

[FR Doc. 99–32026 Filed 12–9–99; 8:45 am]

BILLING CODE 4160–01–F

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
10.30	120	21	120	12	1,440

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

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–0605]

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, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding the 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.

Type of Information Collection Request: Revision of a currently approved collection; *Title of Information Collection:* Hospital Provider of Extender Care Services (Swing-Beds) in the Medicare and Medicaid Programs, 42 CFR 447.280 and 482.66; *Form No.:* HCFA–605 (OMB# 0938–0624); *Use:* This is a facility identification and screening form. It will be completed by a hospital that is requesting approval. It initiates the process of determining the hospital's eligibility and also requests approval for its bed count category. *Frequency:* Other (one time); *Affected Public:* Business or other for profit, and Not for profit institutions; *Number of Respondents:* 50; *Total Annual Responses:* 50; *Total Annual Hours:* 12.5.