Governmentwide Policy, General Services Administration, Washington, DC 20405; telephone (202) 501–3846; email martha.caswell@gsa.gov.

GSA Bulletin FPMR H-79—Utilization and Disposal

To: Heads of Federal agencies Subject: Reporting requirements for firearms

- 1. What is the purpose of this bulletin? This bulletin announces an additional reporting requirement for firearms
- 2. When does this bulletin expire? This bulletin will remain in effect until specifically cancelled.
- 3. What is the background? The Federal Property Management Regulations (FPMR) were amended at 41 CFR 101–43.4801(c) to add new reporting requirements for firearms. On July 23, 1999, GSA Bulletin FPMR H–75 was issued, requiring that each firearm be reported as a single item per report to include serial number, make, and model.
- 4. What must I do as a result of this bulletin? When reporting firearms to GSA, your report must identify each firearm that was acquired through abandonment or forfeiture.
- 5. Whom should I contact for further information? Martha Caswell, Director, Personal Property Management Policy Division (MTP), Office of Governmentwide Policy, General Services Administration, Washington, DC 20405; telephone, (202) 501–3846; email, martha.caswell@gsa.gov.

Dated: December 2, 1999

G. Martin Wager,

Associate Administrator for Governmentwide Policy.

[FR Doc. 99–32073 Filed 12–9–99; 8:45 am] BILLING CODE 6820–24–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Medical Child Support Working Group

AGENCY: Administration for Children and Families, DHHS.

ACTION: Notice of open meeting.

SUMMARY: Pursuant to section 10(a)(2) of the Federal Advisory Committee Act (FACA), notice is given of the revised date for the seventh meeting of the Medical Child Support Working Group (MCSWG). The Medical Child Support Working Group was jointly established by the Secretaries of the Department of

Labor (DOL) and the Department of Health and Human Services (DHHS) under section 401(a) of the Child Support Performance and Incentive Act of 1998. The purpose of the MCSWG is to identify the impediments to the effective enforcement of medical support by State child support enforcement agencies, and to submit to the Secretaries of DOL and DHHS a report containing recommendations for appropriate measures to address those impediments. The Federal Register had previously indicated that the meeting would be December 13 and 14, 1999 (See 64 FR 58858, Nov. 1, 1999 for the DOL Notice and 64 FR 59183, Nov. 2, 1999 for the DHHS Notice).

DATES: The seventh meeting of the MCSWG will be held on Monday, January 10, 2000 and on Tuesday January 11, 2000, from 8:30 a.m. to approximately 6 p.m.

ADDRESSES: The meeting will be held at the Wyndham Bristol Hotel, 2430 Pennsylvania Avenue, NW, Washington, DC 20037, 202–955–6400. All interested parties are invited to attend this public meeting. Seating may be limited and will be available on a first-come, first-serve basis. Persons needing special assistance, such as sign language interpretation or other special accommodation, should contact the Executive Director of the Medical Child Support Working Group, Office of Child Support Enforcement, at the address listed below.

FOR FURTHER INFORMATION CONTACT: Ms. Samara Weinstein, Executive Director, Medical Child Support Working Group, Office of Child Support Enforcement, Fourth Floor East, 370 L'Enfant Promenade, SW, Washington, DC 20447 (telephone (202) 401–6953; fax (202) 401–5559; e-mail:

sweinstein@cf.dhhs.gov). These are not toll-free numbers. The date, location and time for subsequent MCSWG meetings will be announced in advance in the **Federal Register**. However, it is expected this will be the last meeting.

SUPPLEMENTARY INFORMATION: Pursuant to section 10(a)(2) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2) (FACA), notice is given of meetings of the Medical Child Support Working Group (MCSWG). The Medical Child Support Working Group was jointly established by the Secretaries of the Department of Labor (DOL) and the Department of Health and Human Services (DHHS) under section 401(a) of the Child Support Performance and Incentive Act of 1998 (Pub. L. 105–

The purpose of the MCSWG is to identify the impediments to the

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effective enforcement of medical support by State child support enforcement agencies, and to submit to the Secretaries of DOL and DHHS a report containing recommendations for appropriate measures to address those impediments. This report will include: (1) Recommendations based on assessments of the form and content of the National Medical Support Notice, as issued under proposed regulation; (2) appropriate measures that establish the priority of withholding of child support obligations, medical support obligations, arrearages in such obligations, and in the case of a medical support obligation, the employee's portion of any health care coverage premium, by such State agencies in light of the restrictions on garnishment provided under title III of the Consumer Credit Protection Act (15 U.S.C. 1671– 1677); (3) appropriate procedures for coordinating the provision, enforcement, and transition of health care coverage under the State programs for child support, Medicaid and the Child Health Insurance Program; (4) appropriate measures to improve the availability of alternate types of medical support that are aside from health care coverage offered through the noncustodial parent's health plan, and unrelated to the noncustodial parent's employer, including measures that establish a noncustodial parent's responsibility to share the cost of premiums, co-payments, deductibles, or payments for services not covered under a child's existing health coverage; (5) recommendations on whether reasonable cost should remain a consideration under section 452(f) of the Social Security Act; and (6) appropriate measures for eliminating any other impediments to the effective enforcement of medical support orders that the MCSWG deems necessary.

The membership of the MCSWG was jointly appointed by the Secretaries of DOL and DHHS, and includes representatives of: (1) DOL; (2) DHHS; (3) State Child Support Enforcement Directors; (4) State Medicaid Directors; (5) employers, including owners of small businesses and their trade and industry representatives and certified human resource and payroll profession as; (6) plan administrators and plan sponsors of group health plans (as defined in section 607(1) of the Employee Retirement Income Security Act of 1974 (29 U.S.C. 1167(1)); (7) children potentially eligible for medical support, such as child advocacy organizations; (8) State medical child support organizations; and (9)

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) subcommittees 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 BURDEN¹

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

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