DATES: The public meeting will be held on Thursday, December 7, 1995, from 8:30 a.m. to 3:30 p.m.

ADDRESSES: The public meeting will be held at the Sheraton Denver West Hotel, 360 Union Blvd., Lakewood, CO.

FOR FURTHER INFORMATION CONTACT: Virlie M. Walker, FDA Denver District, Bldg. 20, Entrance W–10, Denver Federal Center, Sixth and Kipling Sts., Denver, CO 80225–0087, 303–236–3018, FAX 303–236–3551.

SUPPLEMENTARY INFORMATION: In the Federal Register of April 20, 1995 (60 FR 19753), FDA announced that a series of Grassroots Regulatory Partnership Meetings would be held. This document announces a followup meeting to the one held on April 24, 1995, in Dallas, TX. Those persons interested in attending this public meeting should FAX their registration including name(s), affiliation, address, telephone and FAX numbers, and any specific questions about the workshop to Virlie M. Walker (address above), 303–236–3551. There is no registration fee for this meeting. However due to space limitations, early registration is required. The goal of this meeting is to listen to concerns and ideas, and to identify next steps for the agency.

Dated: November 22, 1995.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 95–29130 Filed 11–28–95; 8:45 am]

BILLING CODE 4160–01–F

Grassroots Regulatory Partnership Meeting; Atlanta and Florida District Offices; Medical Device Industry

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of a public meeting.

SUMMARY: The Food and Drug Administration (FDA) (Office of Regulatory Affairs, Southeast Region/Atlanta and Florida District Offices) is announcing a free public meeting as a followup to a meeting held in April 1995. FDA Atlanta and Florida District Offices will meet with interested persons in Georgia, Florida, North Carolina, and South Carolina to address specific issues related to the medical device industry, Atlanta and Florida Districts, and FDA. The agency is holding this meeting to promote the President's initiative for a partnership approach with front-line regulators and the people affected by the work of this agency, and to create local partnerships.

DATES: The public meeting will be held on Thursday, December 7, 1995, from 8 a.m. to 3:30 p.m.

ADDRESSES: The public meeting will be held at the Sheraton Colony Square Hotel, Peachtree at 14th St., Atlanta, GA.

FOR FURTHER INFORMATION CONTACT:

Sheila Bayne-Lisby, FDA Atlanta District, 60 Eighth St. NE., Atlanta, GA 30309, 404–347–7355, or FAX 404–347–1912, or

Lynne Isaacs, FDA Florida District, 7200 Lake Ellenor Dr., suite 120, Orlando, FL 32809, 407–648–6922 ext. 202, or FAX 407–648–6881.

SUPPLEMENTARY INFORMATION: In the Federal Register of April 20, 1995 (60 FR 19753), FDA announced that a series of Grassroots Regulatory Partnership meetings would be held. This document announces a followup to the one held on April 25, 1995, in Atlanta, GA. Those persons interested in attending this meeting should FAX their comments and registration including name, firm/organization name, address, and telephone number to 404–347–1912. There is no registration fee for this meeting. However, due to space limitations advance registration is required, and all interested parties are encouraged to register early. The goal of this meeting is to ''listen'' to concerns and ideas, and to identify next-steps for the agency.

Dated: November 22, 1995.

William B. Schultz,

Deputy Commissioner for Policy.

[FR Doc. 95–29131 Filed 11–28–95; 8:45 am]

BILLING CODE 4160–01–F

Health Care Financing Administration

Public Information Collection Requirements Submitted for Public Comment and Recommendations

AGENCY: Health Care Financing Administration, Department of Health and Human Services.

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 summaries 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.

1. Type of Information Collection Request: Revision of a currently approved collection; *Title of Information Collection:* Medicare Uniform Institutional Provider Bill; *Form No.:* HCFA–1450; *Use:* Medicare reimbursement of claims. This form is the standardized form used in the Medicare/Medicaid program to apply for reimbursement for covered services by all providers that accept Medicare/Medicaid assigned claims. It will reduce cost and administrative burdens associated with claims since only one coding system is used and maintained. *Frequency:* On occasion; *Affected Public:* Business or other for-profit, not-for-profit institutions, Federal Government, and State, local or tribal government; *Number of Respondents:* 123,432,041; *Total Annual Hours Requested:* 1,890,490.

2. Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* End Stage Renal Disease Medical Evidence Report Medicare Entitlement and/or Patient Registration; *Form No.:* HCFA–2728; *Use:* This form captures the necessary medical information required to determine Medicare eligibility of an end stage renal disease claimant. It also captures the specific medical data required for research and policy decisions on this population as required by law. *Frequency:* Annually; *Affected Public:* Individuals or households, business or other for-profit, not-for-profit institutions; *Number of Respondents:* 60,000; *Total Annual Hours Requested:* 25,200.

To request copies of the proposed paperwork collections referenced above, call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections should be sent within 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Financial and Human Resources, Management Planning and Analysis Staff, Attention: Louis Blank, Room C2–26–17, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.