humans be determined and considered; and (4) under what circumstances might an approved indication for marketing refer to manifestations of disease (biochemical, physiological, anatomic, or pathological processes) common to, or present in, one or more disease states?

Interested parties may want to review section 122 of the FDAMA and a draft regulation for radiopharmaceuticals submitted by the Council on Radionuclides and Radiopharmaceuticals (CORAR). Both the FDAMA and the CORAR proposal have been filed under the docket number found in the heading of this document, and they are available on the Internet.

Electronic Access: Persons with access to the Internet may obtain the FDAMA and the CORAR proposal using the World Wide Web (www) by connecting to ''www.fda.gov/cber/misc.htm''.

Registration and Requests for Oral Presentations: Send registration information (including name, title, firm name, address, telephone, and fax number) and written material and requests to make oral presentations, by February 18, 1998, to Gloria S. Blankenship, Center for Biologics Evaluation and Research (HFM–43), Food and Drug Administration, 1401 Rockville Pike, suite 200N, Rockville, MD 20852–1448, 301–827–1310, FAX 301–827–3079, e-mail ''Blankenship@CBER.FDA.GOV''. Registration at the site will be done on a space available basis on the day of the public meeting beginning at 7:30 a.m.

If you need special accommodations due to a disability, please contact Gloria Blankenship (address above) at least 7 days before the meeting.

Transcripts: Transcripts of the meeting may be requested in writing from the Freedom of Information Office (HFI–35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A–16, Rockville, MD 20857, approximately 15 working days after the meeting at a cost of 10 cents per page.

Dated: January 26, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–2322 Filed 1–30–98; 8:45 am]

BILLING CODE 4160–01–F

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[IA–037–1037b; FRL–5955–3]

Approval and Promulgation of Implementation Plans; State of Iowa

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The EPA proposes to approve the State Implementation Plan (SIP) revision submitted by the state of Iowa for the purpose of updating regulations of the state's two local air pollution control agencies. These agencies are the Polk County Public Works Department and Linn County Health Department.

In the final rules section of the **Federal Register**, the EPA is approving the state's SIP revision as a direct final rule without prior proposal because the Agency views this as a noncontroversial revision amendment and anticipates no adverse comments. The general rationale for the approval is set forth in the direct final rule. If no adverse comments are received in response to this proposed rule, no further activity is contemplated in relation to this rule. If the EPA receives adverse comments, the direct final rule will be withdrawn and all public comments received will be addressed in a subsequent final rule based on this proposed rule. The EPA will not institute a second comment period on this document. Any parties interested in commenting on this document should do so at this time.

DATES: Comments on this proposed rule must be received in writing by March 4, 1998.

ADDRESSES: Comments may be mailed to Christopher D. Hess, Environmental Protection Agency, Air Planning and Development Branch, 726 Minnesota Avenue, Kansas City, Kansas 66101.

FOR FURTHER INFORMATION CONTACT:

Christopher D. Hess at (913) 551–7213.

SUPPLEMENTARY INFORMATION: See the information provided in the direct final rule which is located in the rules section of the **Federal Register** .

Dated: December 30, 1997.

Diane Callier,

Acting Regional Administrator, Region VII.

[FR Doc. 98–2487 Filed 1–30–98; 8:45 am]

BILLING CODE 6560–50–M

DEPARTMENT OF TRANSPORTATION

Research and Special Programs Administration

49 CFR Parts 192, 195

[Docket No. RSPA–98–3347; Notice 1]

Pipeline Safety: Plastic Pipeline Safety Standards

AGENCY: Research and Special Programs Administration, DOT.

ACTION: Notice of public meeting.

SUMMARY: The Research and Special Programs Administration, Office of Pipeline Safety (OPS) invites representatives of the pipeline industry, state and local government, and the public to an open meeting on the Federal gas pipeline safety regulations on plastic pipe system design, construction, maintenance, and rehabilitation in transmission, distribution, and service line applications. The meeting is scheduled to coincide with meetings of the American Gas Association (AGA) Plastic Materials Committee scheduled for the week of March 4, 1998, in Phoenix, Arizona. The purpose of this meeting is to gather information on experience with the current Federal pipeline safety regulations on plastic pipe design, construction, and maintenance and to solicit comments and suggestions to improve these regulations. In particular, OPS seeks comment on whether current regulations should be revised, supplemented, or replaced by references to applicable industry standards and recommended practices.

DATES: The meeting will be held on Wednesday, March 4, 1998, at the Hyatt Regency Phoenix Hotel in Phoenix, Arizona, from 9:00 a.m until all interested persons have been have been afforded an opportunity to speak. Interested persons are invited to attend the meeting and present oral or written statements. Persons wishing to speak at the meeting should notify Jenny Donohue at (202) 366–4046 by the close of business on Friday, February 27, 1998. Please estimate the time that will be required for your presentation. RSPA reserves the right to limit the time of each speaker to ensure that everyone is allowed sufficient time. Other speakers may present statements as time allows.

ADDRESSES: This meeting will be held at the Hyatt Regency Phoenix Hotel, 122 North Second Street, Phoenix, Arizona. The telephone number of the hotel is (602) 252–1234.