comprehensive reference to assist with Regional and State implementation of the rules. The draft guidance has been developed based on input from an Environmental Protection Agency (EPA) Headquarters and Regional staff workgroup, several State-EPA training meetings, and State review of a previous version of the guidance. Along with summaries of each rule, the document contains guidance for preparing State primacy revision applications, and a thorough list of questions and answers compiled during Regional and State training meetings. The implementation guidance covers special primacy requirements for States, information on compliance determinations, Safe Drinking Water Information System (SDWIS) reporting and definitions for significant non-compliance.

DATES: Comments must be submitted on or before July 23, 1999.

ADDRESSES: Address all comments concerning this notice to Nicole Foley (Mailcode 4606), U.S. EPA Headquarters, 401 M Street SW, Washington, DC 20460. See Supplementary Information section for information to request a copy of the draft guidance and electronic addresses.

FOR FURTHER INFORMATION CONTACT: For general information related to IESWTR and Stage 1 DBPR, please contact: Doug McKenna of EPA's Office of Ground Water and Drinking Water at (202) 260–5760 or by sending electronic mail (e-mail) at mckenna.doughlas@epamail.epa.gov, or Nicole Foley at (202) 260–0875 or e-mail at foley.nicole@epamail.epa.gov.

SUPPLEMENTARY INFORMATION: To request a copy of the draft guidance, please contact Nicole Foley of EPA's Office of Ground Water and Drinking Water at (202) 260–0875. You may request a copy of the document or submit comments e-mail to: foley.nicole@epamail.epa.gov.

Cynthia C. Dougherty,

Director, Office of Ground Water and Drinking Water.

[FR Doc. 99–15980 Filed 6–22–99; 8:45 am]

BILLING CODE 6560–50–M

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 12:00 noon, Monday, June 28, 1999.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, NW, Washington, DC 20551.

STATUS: Closed.

MATTERS TO BE CONSIDERED:

1. Personnel actions (appointments, promotions, assignments, reassignments, and salary actions) involving individual Federal Reserve System employees.

2. Any matters carried forward from a previously announced meeting.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.

Dated: June 21, 1999.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 99–16075 Filed 6–21–99; 11:30 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSYTEM

Sunshine Act Meeting

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

FEDERAL REGISTER CITATION OF PREVIOUS ANNOUNCEMENT: 64 FR 32878, June 18, 1999.

PREVIOUSLY ANNOUNCED TIME AND DATE OF THE MEETING: 10:00 a.m., Wednesday, June 23, 1999.

CHANGES IN THE MEETING: Change in the time of the open meeting to 9:00 a.m., Wednesday, June 23, 1999.

CONTACT PERSON FOR MORE INFORMATION: Lynn S. Fox, Assistant to the Board; 202–452–3204.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 for a recorded announcement of this meeting; or you may contact the Board's Web site at http://www.federalreserve.gov for an electronic announcement. (The Web site also includes procedural and other information about the open meeting.)

Dated: June 21, 1999.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 99–16076 Filed 6–21–99; 11:30 am]

BILLING CODE 6210–01–P

GENERAL SERVICES ADMINISTRATION

Availability of Environmental Impact Study Record of Decision

AGENCY: General Services Administration, National Capital Region.

ACTION: Notice.

SUMMARY: The General Services Administration (GSA) National Capital Region (NCR) announces the Record of Decision (ROD) for the Environmental Impact Study (EIS) undertaken for the U.S. Patent and Trademark Office (PTO) consolidation project. The project is for the lease acquisition of 2.4 million rentable square feet with a 20-year term on three possible sites in northern Virginia: Crystal City, Eisenhower Avenue, and Carlyle. The ROD as well as EIS is available at *http://ncr.gsa.gov/pto*.

DATES: The Rod was issued on June 14, 1999 and the final EIS availability was published in the **Federal Register** on January 29, 1999.

FOR FURTHER INFORMATION CONTACT: Carl W. Winters, General Services Administration, Capital Development Division (WPC), 7th & D Streets, S.W., Washington, DC 20407, (202) 401–1025. E-mail *carl.winters@gsa.gov*.

Dated: June 17, 1999.

Jeffrey Hysen,

Assistant Regional Counsel (WL).

[FR Doc. 99–15878 Filed 6–22–99; 8:45 am]

BILLING CODE 6820–BR–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99F–1866]

Goldschmidt Chemical Corp.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Goldschmidt Chemical Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of silicone acrylate resins produced by addition of ω-hydroxyalkenes and/or propenyloxy-2,3-dihydroxypropane, mono- or diester with acrylic acid, acetic acid or other saturated monocarboxylic acid, to dimethyl polysiloxane, methylhydrogen polysiloxane, or dimethyl-methylhydrogen polysiloxane as

coatings or components of coatings on polymers and on paper and paperboard intended for contact with food. The following optional adjuvants may also be required in the manufacture of silicone acrylate resins: 2-hydroxy-2-methyl-1-phenyl-1-propanone and/or oligomeric 2-hydroxy-2-methyl-1-[4-(1-methylvinyl)phenyl]-1-propanone.

FOR FURTHER INFORMATION CONTACT: Hortense S. Macon, Center for Food Safety and Applied Nutrition (HFS–206), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3086.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5)), notice is given that a food additive petition (FAP 9B4658) has been filed by Goldschmidt Chemical Corp., 914 East Randolph Rd., Hopewell, VA 23860. The petition proposes both to amend the food additive regulations in part 177 (21 CFR part 177) by adding a new section and to amend § 176.170 *Components of paper and paperboard in contact with aqueous and fatty foods* (21 CFR 176.170) to provide for the safe use of silicone acrylate resins produced by addition of ω-hydroxyalkenes and/or propenyloxy-2,3-dihydroxypropane, mono- or diester with acrylic acid, acetic acid or other saturated monocarboxylic acid, to dimethyl polysiloxane, methylhydrogen polysiloxane, or dimethyl-methylhydrogen polysiloxane as coatings or components of coatings on polymers and on paper and paperboard intended for contact with food. The following optional adjuvants may also be required in the manufacture of silicone acrylate resins: 2-hydroxy-2-methyl-1-phenyl-1-propanone and/or oligomeric 2-hydroxy-2-methyl-1-[4-(1-methylvinyl)phenyl]-1-propanone.

The agency has determined under 21 CFR 25.32(i) that this action is of the type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: June 2, 1999.

Alan M. Rulis,

Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 99–15877 Filed 6–22–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

The FDA Review Process for New Product Applications: An Interactive Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of workshop.

The Food and Drug Administration (FDA), Los Angeles District, in cosponsorship with the Orange County Regulatory Affairs Discussion Group (OCRA) is announcing the following workshop: The FDA Review Process for New Product Applications: An Interactive Workshop, which is intended to give the medical products industry (drugs, biologics, and medical devices) an opportunity to learn and discuss the process by which the centers and district offices review new product applications. Reviewing staff from the Centers for Biologics, Devices, and Drugs will make presentations regarding the elements of submissions that make the review process more efficient.

Date and Time: The workshop will be held on July 12 and 13, 1999, from 7:30 a.m. to 5 p.m.

Location: The workshop will be held at the Irvine Marriott, 18000 Von Karman Ave., Irvine, CA, 949–553–0100.

Contact: Sandi Velez, Los Angeles District Office, Food and Drug Administration, 19900 MacArthur Blvd., Irvine, CA 92612–2445, 949–798–7748 or FAX 949–798–7715, for further information including a registration form.

Registration: Space is limited. Preregistration and confirmation are required. Registration forms can be obtained at the OCRA web site ''http://www.ocra-dg.org'' or from Sandi Velez at the numbers given previously. There is a $250 registration fee if postmarked by June 30, 1999 ($275 after July 1, 1999) payable to OCRA. The registration fee and form should be sent to PeriAnn DiRocco at OCRA Submissions Conference, 5405 Alton Pkwy., suite 5A–624, Irvine, CA 92604, FAX and voice 949–348–9141, and received no later than July 7, 1999. The registration fee will cover actual expenses incurred by OCRA including refreshments, lunch, materials, parking fees, and speaker expenses.

If you need special accommodations due to disability, please contact Sandi Velez at least 7 days in advance.

Dated: June 17, 1999.

Margaret M. Dotzel,

Acting Associate Commissioner for Policy Coordination.

[FR Doc. 99–16091 Filed 6–21–99; 2:29 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–R–263]

Agency Information Collection Activities: Submission for OMB Review; 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, has submitted to the Office of Management and Budget (OMB) the following proposal for the collection of information. 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:* On Site Inspection for Durable Medical Equipment (DME) Supplier Location & Supporting Regulations in 42 CFR, 424.57; *Form Nos.:* HCFA–R–263 (OMB# 0938–0749);

Use: To identify and implement measures to prevent fraud and abuse in the Medicare program. Controlling the entry of suppliers of durable medical equipment, prosthetics, orthotics, or supplies (DMEPOS) to Medicare has been identified as one of the most effective ways to prevent fraud and abuse. To meet this challenge, HCFA is moving forward with a plan to improve the quality of the process for enrolling and reenrolling DMEPOS suppliers into the Medicare program by enhancing procedures for verifying supplier information collected on the Form HCFA 855S (DMEPOS Supplier Enrollment Application, OMB Approval No. 0938–0685). This form will be used