

Action: Revision.

Respondents: Businesses or other for-profit (including small businesses).

Frequency of Response: On occasion reporting and other: Reports are to be filed 6, 9, 15 and 21 months after 12/21/90.

Estimated Annual Burden: 1,200 responses, 10 hours average burden per respondent, 12,000 hours total annual burden.

Needs and Uses: Operator service providers are required to file reports to the FCC on rates, complaints and cost information. The information will be used by the Commission for monitoring purposes and to prepare reports to Congress as required by the Telephone Operator Consumer Services Improvement Act of 1990 (TOCSIA).

Federal Communications Commission.

Donna R. Searcy.

Secretary.

[FR Doc. 91-14178 Filed 6-13-91; 8:45 am]

BILLING CODE 6712-01-M

FEDERAL MARITIME COMMISSION**Notice of Agreement(s) Filed**

The Federal Maritime Commission hereby gives notice of the filing of the following agreement(s) pursuant to section 5 of the Shipping Act of 1984.

Interested parties may inspect and obtain a copy of each agreement at the Washington, DC Office of the Federal Maritime Commission, 1100 L Street, NW., room 10325. Interested parties may submit comments on each agreement to the Secretary, Federal Maritime Commission, Washington, DC 20573, within 10 days after the date of the *Federal Register* in which this notice appears. The requirements of comments are found in § 572.603 of title 46 of the Code of Federal Regulations. Interested persons should consult this section before communicating with the Commission regarding a pending agreement.

Agreement No.: 203-009857-004.

Title: Florida-Caribbean Cruise Association.

Parties:

Carnival Cruise Line
Chandris, Inc.
Commodore Cruise Line
Costa Cruise Lines, N.V.
Cunard/NAC
Dolphin Cruise Line
Kloster Cruise limited
Premier Cruise Line
Princess Cruises
Royal Caribbean Cruises, Ltd.

Synopsis: The proposed amendment would expand the geographic scope to

include all North American ports and ports in Central America and South America. It would also make other nonsubstantive changes.

Agreement No.: 203-011038-010.

Title: Southeastern Caribbean Discussion Agreement.

Parties:

United States Atlantic and Gulf/
Southeastern Caribbean Conference
"Independent Carrier Parties"
Seaboard Marine, Ltd.
Tecmarine Line, Inc.
Bernuth Lines
North American Caribbean Line Ltd.
Blue Caribe Line.

Synopsis: The proposed amendment would add West Indies Shipping Corporation as an Independent Carrier Party to the Agreement. The parties have requested a shortened review period.

Agreement No.: 232-011335.

Title: TMM/ACS Space Charter and Sailing Agreement.

Parties:

Transportation Maritima Mexicana
S.A. de C.V.
Star Shipping A/S.

Synopsis: The proposed Agreement would permit the parties to charter space on each other's vessels, coordinate sailings, utilize common terminals, jointly negotiate terminal and stevedoring arrangements and interchange equipment in the trade between U.S. Atlantic and Gulf ports and U.S. points via such ports and Northern European ports and points via such ports. The parties have requested a shortened review period.

Dated: June 10, 1991.

By Order of the Federal Maritime Commission

Joseph C. Polking,

Secretary

[FR Doc. 14122 Filed 6-13-91; 8:45 am]

BILLING CODE 6730-01-M

Notice of Agreement(s) Filed

The Federal Maritime Commission hereby gives notice of the filing of the following agreement(s) pursuant to section 5 of the Shipping Act of 1984.

Interested parties may inspect and obtain a copy of each agreement at the Washington, DC Office of the Federal Maritime Commission, 1100 L Street, NW., Room 10220. Interested parties may submit comments on each agreement to the Secretary, Federal Maritime Commission, Washington, DC 20573, within 10 days after the date of the *Federal Register* in which this notice appears. The requirements for

comments are found in § 572.603 of Title 46 of the Code of Federal Regulations. Interested persons should consult this section before communicating with the Commission regarding a pending agreement.

Agreement No.: 224-004070-004.

Title: San Francisco Port Commission/Stevedoring Services of America Terminal Agreement.

Parties:

San Francisco Port Commission
Stevedoring Services of America
(SSA).

Synopsis: The Agreement, filed June 3, 1991, amends the basic agreement to: (1) Increase the service area at Pier 94 & 96; (2) provide rent credits for certain improvements; and (3) clarify that the Agreement cannot be extended beyond October 31, 1992.

Dated: June 10, 1991.

By Order of the Federal Maritime Commission.

Joseph C. Polking,

Secretary.

[FR Doc. 91-14123 Filed 6-13-91; 8:45 am]

BILLING CODE 6730-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control**

[Announcement Number 123]

National Institute for Occupational Safety and Health; Grants for Education Programs in Occupational Safety and Health**Introduction**

The National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control (CDC), announces the availability of funds in Fiscal Year 1992 for training grants in occupational safety and health. The Public Health Service (PHS) is committed to achieving the health promotion and disease prevention objectives of Healthy People 2000, a PHS-led national activity to reduce morbidity and mortality and improve the quality of life. This announcement is related to the priority area of Occupational Safety and Health. (For ordering a copy of Healthy People 2000, see the Section Where to obtain additional information.)

Authority

This program is authorized under section 21(a)(1) of the Occupational Safety and Health Act of 1970 (29 U.S.C. 670(a)(1)). Regulations applicable to this program are in 42 CFR part 86, "Grants

for Education Programs in Occupational Safety and Health."

Eligible Applicants

Any public or private educational or training agency or institution that has demonstrated competency in the occupational safety and health field and is located in a State, the District of Columbia, or U.S. Territory is eligible to apply for a training grant.

Availability of Funds and Recipient Activities

Funds in the total amount of \$11,072,000 will be available in Fiscal Year 1992.

Approximately \$10,225,000 of the total funds available will be utilized as follows:

1. To award approximately 5 non-competing continuation and 9 competing Educational Resource Center (ERC) training grants totaling approximately \$8,200,000 and ranging from approximately \$400,000 to \$800,000 with the average award being approximately \$600,000 (for specific activities refer to **Federal Register** Announcement, 51 FR 32963, September 17, 1986).

2. To award approximately 24 non-competing continuation and 6 competing long-term training project grants totaling \$2,000,000 and ranging from approximately \$10,000 to \$500,000, with the average award being \$50,000, to support academic programs in the fields of industrial hygiene, occupational health nursing, occupational/industrial medicine, and occupational safety (for specific activities refer to **Federal Register** Announcement, 52 FR 3172, February 2, 1987); and

3. To conduct the peer review and evaluations of all new, competing continuation, and supplemental applications received.

Awards will be made for a 1- to 5-year project period with an annual budget period. Funding estimates may vary and are subject to change. Non-competing continuation awards within the approved project periods will be made on the basis of satisfactory progress and the availability of funds.

In addition, approximately \$600,000 of the total funds available will be awarded to ERCs to support the development and presentation of continuing education and short courses for professionals engaged in the management of hazardous substances. These funds were provided to NIOSH through an Interagency Agreement with the National Institute for Environmental Health Sciences as authorized by section 209(b) of the Superfund Amendments and Reauthorization Act (SARA) of 1986 (100 Stat. 1708-1710).

The hazardous substance training funds are being used to supplement previous hazardous substance continuing education grant support provided to the ERC's in FY 1984 and 1985 under the authority of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) of 1980 for the ERC continuing education program (for specific activities refer to the **Federal Register** Announcement, 51 FR 32963, September 17, 1986). Program support is available for faculty and staff salaries, and other costs to provide occupational safety and health training to practicing professionals in State and local health and environmental agencies and other professional personnel engaged in the evaluation, management, and handling of hazardous substances. The policies regarding project periods also apply to these activities.

Approximately \$247,000 of the total funds available will be awarded to ERCs to support the development of specialized educational programs in agricultural safety and health within the existing core disciplines of industrial hygiene, occupational medicine, occupational health nursing, and occupational safety (for specific activities refer to **Federal Register** Announcement, 51 FR 32963, September 17, 1986). Program support is available for faculty and staff salaries, trainee costs, and other costs to educate professionals in agricultural safety and health.

Purpose

The objective of this grant program is to award funds to eligible institutions or agencies to assist in providing an adequate supply of qualified professional and para-professional occupational safety and health personnel to carry out the purposes of the Act.

Review and Evaluation Criteria

In reviewing ERC grant applications, consideration will be given to:

1. Needs assessment directed to the overall contribution of the training program toward meeting the job market, especially within the applicant's region, for qualified personnel to carry out the purposes of the Occupational Safety and Health Act of 1970. The needs assessment should consider the regional requirements for outreach, continuing education, information dissemination, and special industrial or community training needs that may be peculiar to the region.

2. Plan to satisfy the regional needs for training in the areas outlined by the application, including projected

enrollment, recruitment and current workforce populations. The need for supporting students in allied disciplines must be specifically justified in terms of user community requirements.

3. The extent to which arrangements for day-to-day management, allocation of funds and cooperative arrangements are designed to effectively achieve Characteristics of an Educational Resource Center (**Federal Register** Announcement, 51 FR 32963, September 17, 1986).

4. The extent to which curriculum content and design includes formalized training objectives, minimal course content to achieve certificate or degree, course descriptions, course sequence, additional related courses open to occupational safety and health students, time devoted to lecture, laboratory and field experience, and the nature of specific field and clinical experiences including their relationships with didactic programs in the educational process.

5. Academic training including the number of full-time and part-time students and graduates for each core program, the placement of graduates, employment history, and their current location by type of institution (academic, industry, labor, etc.). Previous continuing education training in each discipline and outreach activity and assistance to groups within the ERC region.

6. Methods in use or proposed methods for evaluating the effectiveness of training and services including the use of placement services and feedback mechanisms from graduates as well as employers, critiques from continuing education courses, and reports from consultations and cooperative activities with other universities, professional associations, and other outside agencies.

7. The competence, experience and training of the Center Director, the Deputy Center Director, the Program Directors and other professional staff in relation to the type and scope of training and education involved.

8. Institutional commitment to Center goals.

9. Academic and physical environment in which the training will be conducted, including access to appropriate occupational settings.

10. Appropriateness of the budget required to support each academic component of the ERC program, including a separate budget for the academic staff's time and effort in continuing education and outreach.

11. Evidence of a plan describing the research and research training the Center proposes. This shall include

goals, elements of the program, research faculty and amount of effort, support faculty, facilities and equipment available and needed, and methods for implementing and evaluating the program.

12. Evidence of success in attaining outside support to supplement the ERC grant funds including other federal grants, support from states and other public agencies, and support from the private sector including grants from foundations and corporate endowments, chairs, and gifts.

In reviewing long-term training project grant applications, consideration will be given to:

1. The need for training in the program area outlined by the application. This should include documentation of ability and a plan for student recruitment, projected enrollment, job opportunities, regional/national need both in quality and quantity, and similar programs, if any within the geographic area.

2. The potential contribution of the project toward meeting the needs for graduate or specialized training in occupational safety and health.

3. Curriculum content and design which should include formalized program objectives, minimal course content to achieve certificate or degree, course sequence, related courses open to students, time devoted to lecture, laboratory and field experience, nature and the interrelationship of these educational approaches.

4. Previous records of training in this or related areas, including placement of graduates.

5. Methods proposed to evaluate effectiveness of the training.

6. Degree of institutional commitment: Is grant support necessary for program initiation or continuation? Will support gradually be assumed? Is there related instruction that will go on with or without the grant?

7. Adequacy of facilities (classrooms, laboratories, library services, books, and journal holdings relevant to the program, and access to appropriate occupational settings).

8. The competence, experience, training, time commitment to the program and availability of faculty to advise students, faculty/student ratio, and teaching loads of the program director and teaching faculty in relation to the type and scope of training involved.

9. Admission Requirements: Student selection standards and procedures, student performance standards and student counseling services.

10. Advisory Committee (if established): Membership, industries and labor groups represented; how often

they meet; who they advise, role in designing curriculum and establishing program need.

Funding Allocation Criteria

For Educational Resource Center grants, the following criteria will be considered in determining funding allocations.

1. Academic core programs

a. Budget to support programs primarily for personnel and other personnel-related costs. Advanced (doctoral and post-doctoral) and specialty (master's) programs will be considered.

b. Budget to support programs based on program quality and need. Factors considered include faculty commitment/breadth, faculty reputation/strength, national/regional manpower needs, unique program contribution, interdisciplinary interaction, and technical merit.

c. Budget to support students based on the program level and the number of students supported.

2. Center administration

Budget to support Center administration to assure coordination and promotion of academic programs.

3. Continuing education/outreach program

Budget to support outreach and continuing education activities to prepare, distribute, and conduct short courses, seminars, and workshops.

4. Research training programs

Budget to support research training programs to establish a research base within the core disciplines and for the training of researchers in occupational safety and health.

5. Hazardous substance training programs

Budget to support the development and presentation of continuing education courses for professionals engaged in the management of hazardous substances.

6. Agricultural safety and health academic programs

Budget to support the development and presentation of specialized academic programs and continuing education courses in agricultural safety and health.

For Long-Term Training Project grants, the following factors will be considered in determining funding allocations.

1. Academic core programs

a. Budget to support programs primarily for personnel and other personnel-related costs. Advanced (doctoral and post-doctoral), specialty (master's), and baccalaureate/associate programs will be considered.

b. Budget to support programs based on program quality and need. Factors considered include faculty commitment/breadth, faculty reputation/strength, national/regional manpower needs, unique program contribution, interdisciplinary interaction, and technical merit.

c. Budget to support students based on the program level and the number of students supported.

Catalog of Federal Domestic Assistance Number (CFDA)

This program is described in the Catalog of Federal Domestic Assistance Program No. 93.263, Occupational Safety and Health Training Grants.

Executive Order 12372 Review

Applications are not subject to review as governed by Executive Order 12372, Intergovernmental Review of Federal Programs.

Application Submission and Deadline

Applications should be clearly identified as an application for an Occupational Safety and Health Long-Term Training Project Grant or ERC Training Grant. The submission schedule is as follows:

New, Competing Continuation and Supplemental Receipt Date: July 1, 1991.

An original and two copies of new, competing continuation and supplemental applications (Form CDC 2.145A ERC or TPG) should be submitted to: Henry S. Cassell, III, Grants Management Officer, Procurement and Grants Office, Centers for Disease Control, 255 East Paces Ferry Road, NE, room 300, Atlanta, GA 30305.

1. Deadline: Applications shall be considered as meeting the deadline if they are either:

a. Received on/or before the deadline date, or

b. Sent on or before the deadline date and received in time for submission to the independent review group. (Applicants must request a legibly dated U.S. Postal Service postmark or obtain a legibly dated receipt from a commercial carrier or the U.S. Postal Service. Private metered postmarks shall not be acceptable as proof of timely mailing.)

2. Late Applications: Applications which do not meet the criteria in 1.a. or b. above are considered late

applications. Late applications will not be considered in the current competition and will be returned to the applicant.

Non-Competing Continuation Receipt Date: November 15, 1991 An original and two copies of non-competing continuation applications (Form CDC 2.145B ERC or TPG) should be submitted to: Henry S. Cassell, III, Grants Management Officer, Procurement and Grants Office, Centers for Disease Control, Procurement and Grants Office, 255 East Paces Ferry Road, NE, room 300, Atlanta, GA 30305.

Where to Obtain Additional Information

To receive additional written information call (404) 332-4561. You will be asked to leave your name, address and phone number, and will need to refer to Announcement Number 123. You will receive a complete program description, information on application procedures, and an application package containing address and phone numbers for the contact personnel. Lisa Tamaroff will provide the business management technical assistance.

Programmatic technical assistance may be obtained from John T. Talty, Chief, Educational Resource Development Branch, Division of Training and Manpower Development, National Institute for Occupational Safety and Health, CDC, 4676 Columbia Parkway, Cincinnati, Ohio 45226.

Potential applicants may obtain a copy of Healthy People 2000 (Full Report, Stock No. 017-001-00474-0) or Healthy People 2000 (Summary Report, Stock No. 017-001-00473-1) through the Superintendent of Documents, Government Printing Office, Washington, DC 20402-9325 (Telephone (202) 783-3238).

Dated: June 10, 1991.

Larry W. Sparks,

Acting Director, National Institute for Occupational Safety and Health.

[FR Doc. 91-14175 Filed 6-13-91; 8:45 am]

BILLING CODE 4160-19-M

Family Support Administration

Forms Submitted to the Office of Management and Budget for Clearance

The Administration for Children and Families will publish on Fridays information collection packages submitted to the Office of Management and Budget (OMB) for clearance, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). Following is the package submitted to OMB since the last publication.

(For a copy of the package, call the FSA, Report Clearance Officer 202-401-5604.)

Quarterly Report of Expenditures and Estimates—OCSE-131—0970-0014—

This information and data is needed to compute quarterly grant awards to estimate incentive payments to States, for required recordkeeping, prepare appropriation requests and to prepare an annual report to Congress. *Respondents:* State and local governments; *Number of Respondents:* 54; *Frequency of Response:* Quarterly; *Estimated Average Burden per Response:* 3.5; *Estimated Annual Burden:* 1,134 hours. *OMB Desk Officer:* Laura Oliven.

Written comments and recommendations for the proposed information collection should be sent directly to the OMB Desk Officer designated above at the following address: OMB Reports Management Branch, New Executive Office Building, Room 3201, 725 17th Street, NW., Washington, DC 20503.

Dated: June 10, 1991.

Naomi B. Marr,

Associate Administrator, Office of Management and Information Systems

[FR Doc. 91-14140 Filed 6-13-91; 8:45 am]

BILLING CODE 4150-04-M

Food and Drug Administration

[Docket No. 91C-0189]

Sola/Barnes-Hind; Filing of Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Sola/Barnes-Hind, has filed a petition proposing that the color additive regulations be amended to provide for the safe use of 1,4-bis(2-methacryloxyethylamino)-9,10-anthraquinone to color contact lenses prepared with glyceryl methacrylate/methyl methacrylate/ethylene glycol dimethacrylate copolymer and N,N-dimethyl acrylamide/methyl methacrylate/ethylene glycol dimethacrylate copolymer.

FOR FURTHER INFORMATION CONTACT: Sandra L. Varner, Center for Food and Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 706(d)(1) (21 U.S.C. 376(d)(1))), notice is given that a petition (CAP

OC0226), has been filed by Sola/Barnes-Hind, 810 Kifer Rd., Sunnyvale, CA 94086-5200, proposing that 21 CFR part 73 of the color additive regulations be amended to provide for the safe use of 1,4-bis(2-methacryloxyethylamino)-9,10-anthraquinone to color contact lenses prepared with glyceryl methacrylate/methyl methacrylate/ethylene glycol dimethacrylate copolymer and N,N-dimethyl acrylamide/methyl methacrylate/ethylene glycol dimethacrylate copolymer.

The potential environmental impact of this action is being reviewed. If the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: June 7, 1991.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 91-14238 Filed 6-13-91; 8:45 am]

BILLING CODE 4160-01-M

Advisory Committees; Meetings

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: This notice announces forthcoming meetings of public advisory committees of the Food and Drug Administration (FDA). This notice also summarizes the procedures for the meetings and methods by which interested persons may participate in open public hearings before FDA's advisory committees.

Meetings: The following advisory committee meetings are announced:

Oncologic Drugs Advisory Committee

Date, time, and place. July 1 and 2, 1991, 9 a.m., Conference Rms. D and E, Parklawn Bldg., 5600 Fishers Lane, Rockville, MD.

Type of meeting and contact person. Open committee discussion, July 1, 1991, 9 a.m. to 4:30 p.m.; open public hearing, 4:30 p.m. to 5:30 p.m., unless public participation does not last that long; open committee discussion, July 2, 1991, 8:30 a.m. to 5 p.m.; Adele S. Seifried, Center for Drug Evaluation and Research (HFD-9), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4695.

General function of the committee. The committee reviews and evaluates

data on the safety and effectiveness of marketed and investigational human drugs for use in the treatment of cancer.

Agenda—Open public hearing.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before June 21, 1991, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. On July 1, 1991, the committee will discuss: (1) New drug application (NDA) NDA 20-119, VeeM-26 (tenoposide, Bristol-Myers Squibb), for multiple indications for childhood acute lymphocytic leukemia; (2) supplemental NDA 19-880, Paraplatin (carboplatin, Bristol-Myers Squibb), for first line combination chemotherapy stages III and IV ovarian cancer; and (3) NDA 20-141, Isovorin (1-leucovorin, Lederle Labs), for rescue after high dose methotrexate for osteosarcoma, patients with impaired methotrexate elimination, and inadvertent methotrexate overdose. On July 2, 1991, the committee will discuss: (1) NDA 20-122, Oncopent (pentostatin, Parke-Davis), for alpha interferon refractory hairy cell leukemia; and (2) the NSABP tamoxifen breast cancer prevention protocol.

Endocrinologic and Metabolic Drugs Advisory Committee

Date, time, and place. July 24 and 25, 1991, 9 a.m., Holiday Inn Crowne Plaza, Regency Rm., 1750 Rockville Pike, Rockville, MD.

Type of meeting and contact person. Open public hearing, July 24, 1991, 9 a.m. to 10 a.m., unless public participation does not last that long; open committee discussion, 10 a.m. to 5 p.m.; open public hearing, July 25, 1991, 9 a.m. to 10 a.m., unless public participation does not last that long; open committee discussion, 10 a.m. to 5 p.m.; John L. Gueriguan, Center for Drug Evaluation and Research (HFD-510), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-3490, fax 301-443-9282.

General function of the committee. The committee reviews and evaluates data on the safety and effectiveness of marketed and investigational human drugs for use in endocrine and metabolic disorders.

Agenda—Open public hearing.

Interested persons may present data, information, or views, orally or in writing, on issues pending before the

committee. Those desiring to make formal presentations should notify the contact person before June 28, 1991, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. On July 24, 1991, the committee will discuss the osteoporosis indication of injectable salmon calcitonin (Calcimar, Rhone-Poulenc Rorer) in light of recently acquired efficacy data. On July 25, 1991, the committee will discuss safety and efficacy for NDA 20-086, acarbose (Glucobay), submitted by Miles, Inc.

Gastrointestinal Drugs Advisory Committee

Date, time, and place. July 25 and 26, 1991, 9 a.m., Conference Rms. D and E, Parklawn Bldg., 5600 Fishers Lane, Rockville, MD.

Type of meeting and contact person. Open public hearing, July 25, 1991, 9 a.m. to 10 a.m., unless public participation does not last that long; open committee discussion, 10 a.m. to 5 p.m.; open committee discussion, July 26, 1991, 9 a.m. to 5 p.m.; Joan C. Standaert, Center for Drug Evaluation and Research (HFD-180), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 419-259-6211, or the Advisors and Consultants Staff, 301-443-4695.

General function of the committee. The committee reviews and evaluates data on the safety and effectiveness of marketed and investigational human drugs for use in gastrointestinal diseases.

Agenda—Open public hearing. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before July 10, 1991, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time required to make their comments.

Open committee discussion. On July 25, 1991, the committee will discuss NDA 17-939, Tagamet (cimetidine), continuous intravenous injection for prevention of stress ulcer and a draft guidelines for the study of Crohn's disease. On July 26, 1991, the committee will discuss the use of bismuth subsalicylate (Pepto-Bismol) as a Category I over-the-counter (OTC) antidiarrheal.

The committee discussion and conclusions regarding the use of bismuth subsalicylate may be considered by the agency in its preparation of a final monograph for OTC antidiarrheal drug products. This monograph is being developed as part of the OTC drug review. The advance notice of proposed rulemaking for OTC antidiarrheal drug products was published in the **Federal Register** of April 30, 1986 (51 FR 16138). Bismuth subsalicylate has not been proposed as an active ingredient for inclusion in this monograph.

FDA public advisory committee meetings may have as many as four separable portions: (1) An open public hearing, (2) an open committee discussion, (3) a closed presentation of data, and (4) a closed committee deliberation. Every advisory committee meeting shall have an open public hearing portion. Whether or not it also includes any of the other three portions will depend upon the specific meeting involved. There are no closed portions for the meetings announced in this notice. The dates and times reserved for the open portions of each committee meeting are listed above.

The open public hearing portion of each meeting shall be at least 1 hour long unless public participation does not last that long. It is emphasized, however, that the 1 hour time limit for an open public hearing represents a minimum rather than a maximum time for public participation, and an open public hearing may last for whatever longer period the committee chairperson determines will facilitate the committee's work.

Public hearings are subject to FDA's guidelines (Subpart C of 21 CFR part 10) concerning the policy and procedures for electronic media coverage of FDA's public administrative proceedings, including hearings before public advisory committees under 21 CFR part 14. Under 21 CFR 10.205, representatives of the electronic media may be permitted, subject to certain limitations, to videotape, film, or otherwise record FDA's public administrative proceedings, including presentations by participants.

Meetings of advisory committees shall be conducted, insofar as is practical, in accordance with the agenda published in this **Federal Register** notice. Changes in the agenda will be announced at the beginning of the open portion of a meeting.

Any interested person who wishes to be assured of the right to make an oral presentation at the open public hearing portion of a meeting shall inform the contact person listed above, either

orally or in writing, prior to the meeting. Any person attending the hearing who does not in advance of the meeting request an opportunity to speak will be allowed to make an oral presentation at the hearing's conclusion, if time permits, at the chairperson's discretion.

Persons interested in specific agenda items to be discussed in open session may ascertain from the contact person the approximate time of discussion.

Details on the agenda, questions to be addressed by the committee, and a current list of committee members are available from the contact person before and after the meeting. Transcripts of the open portion of the meeting will be available from the Freedom of Information Office (HFI-35), Food and Drug Administration, rm. 12A-16, 5600 Fishers Lane, Rockville, MD 20857, approximately 15 working days after the meeting, at a cost of 10 cents per page. The transcript may be viewed at the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857, approximately 15 working days after the meeting, between the hours of 9 a.m. and 4 p.m., Monday through Friday. Summary minutes of the open portion of the meeting will be available from the Freedom of Information Office (address above) beginning approximately 90 days after the meeting.

This notice is issued under section 10(a)(1) and (2) of the Federal Advisory Committee Act (5 U.S.C. app. 2), and FDA's regulations (21 CFR part 14) on advisory committees.

Alan L. Hoeting,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 91-14371 Filed 6-12-91; 2:18 p.m.]

BILLING CODE 4160-01-M

Health Resources and Services Administration

Infant Mortality Advisory Council; Establishment

Pursuant to the Federal Advisory Committee Act, Public Law 92-463 (5 U.S.C. appendix 2), the Secretary, Department of Health and Human Services announces the establishment of the following advisory committee.

Designation: Advisory Committee on Infant Mortality

Purpose: Advises the Secretary on Department programs related to infant mortality, including implementation of the Healthy Start Initiative and other recommendations of the Domestic Policy Council, and infant mortality objectives from Healthy People: 2000: National

Health Promotion and Disease Prevention Objectives.

Structure: The Committee consists of the Assistant Secretary for Health, the Assistant Secretary for Children and Families, the Administrator, Health Care Financing Administration and the Assistant Secretary for Food and Consumer Services, USDA, as ex-officio members, and 21 members, including the chair, selected by the Secretary.

Members shall be representatives of both the public and private sectors. Members from the private sector will represent corporations and foundations, the media, the clergy, health and other professional organizations. Members from the public sector will include elected officials, and health and human services administrators from State and local levels including representatives of minority, rural and urban interests.

Authority for this Committee will terminate in two years unless the Secretary, HHS, formally determines that continuance is in the public interest.

Dated: June 11, 1991.

Jackie E. Baum,

Advisory Committee Management Officer, HRSA.

[FR Doc. 91-14239 Filed 6-13-91; 8:45 am]

BILLING CODE 4160-15-M

Social Security Administration

Agency Forms Submitted to the Office of Management and Budget for Clearance

Each Friday the Social Security Administration publishes a list of information collection packages that have been submitted to the Office of Management and Budget (OMB) for clearance in compliance with Public Law 96-511, The Paperwork Reduction Act. The following clearance packages have been submitted to OMB since the last list was published in the Federal Register on May 17, 1991.

(Call Reports Clearance Officer on (301) 965-4149 for copies of package.)

1. Compliance With Pass-Along—0960-0240—The authority granted by the final rule F-20-416.2099 allows the Social Security Administration (SSA) to request cost-of-living adjustment pass-along information from those States which administer their own supplementary payments programs. Without this authority, SSA would be unable to determine that those States have complied with the requirement to meet certain supplementary expenditures. Compliance failure could

result in a state losing Federal financial participation in the title XIX, Medicaid program. The affected public consists of States which administer their own supplementary payments program.

Number of Respondents: 26

Frequency of Response: 15(4) ²/11(1) ¹

Average Burden Per Response: 1 hour

Estimated Annual Burden: 71 hours

2. Employment Relationship Questionnaire—0960-0040—The information collected on the form SSA-7160 is used by the Social Security Administration to determine the status of certain employees in various occupational fields. The affected public is comprised of employers of individuals applying for Social Security benefits whose employee status is in doubt.

Number of Respondents: 47,500

Frequency of Response: 1

Average Burden Per Response: 25 minutes

Estimated Annual Burden: 19,792 hours

3. Statement Regarding Contributions—0960-0020—The information collected on the form SSA-783 is used by the Social Security Administration to make determination of one-half support which is an eligibility factor for certain claimants. The affected public is comprised of claimants who need to prove one-half support by the wage earner or other persons having knowledge of the situation.

Number of Respondents: 30,000

Frequency of Response: 1

Average Burden Per Response: 15 minutes

Estimated Annual Burden: 7,500 hours

OMB Desk Officer: Laura Oliven.

Written comments and recommendations regarding these information collections should be sent directly to the appropriate OMB Desk Officer designated above at the following address: OMB Reports Management Branch, New Executive Office Building, room 3208, Washington, DC 20503.

Dated: June 10, 1991.

Ron Compston,

Social Security Administration, Reports Clearance Officer.

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¹ 15 States report four times annually

² 11 States report once annually