otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act. Additional information on all bank holding companies may be obtained from the National Information Center website at *www.ffiec.gov/nic/*.

Unless otherwise noted, comments regarding the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than February 20, 2001.

A. Federal Reserve Bank of Chicago (Phillip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 606901–1413:

1. AmericaUnited Bancorp, Inc., Schaumburg, Illinois; to engage *de novo* in extending credit and servicing loans, pursuant to § 225.28(b)(1) of Regulation Y.

B. Federal Reserve Bank of San Francisco (Maria Villanueva, Consumer Regulation Group) 101 Market Street, San Francisco, California 94105–1579:

1. Mitsubishi Tokyo Financial Group, Inc., and The Bank of Tokyo–Mitsubishi, Ltd., both of Tokyo, Japan; to acquire KOKUSAI America Incorporated, New York, New York, and thereby engage in providing financial and investment advisory services, pursuant to § 225.28(b)(6) of Regulation Y, providing certain agency transactional services for customer investments, pursuant to § 225.28(b)(7) of Regulation Y, and engaging in investment transactions as principal, pursuant to § 225.28(b)(8) of Regulation Y.

Board of Governors of the Federal Reserve System, January 31, 2001.

Robert deV. Frierson

Associate Secretary of the Board.

[FR Doc. 01–3005 Filed 2–5–01; 8:45 am]

BILLING CODE 6210–01–S

FEDERAL RESERVE SYSTEM

Sunshine Act Meeting

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

TIME AND DATE: 11:00 a.m., Monday, February 12, 2001.

PLACE: Marriner S. Eccles Federal Reserve Board Building, 20th and C Streets, N.W., Washington, D.C. 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 items 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 indicated procedural and other information about the meeting.

February 2, 2001.

Robert deV. Frierson,

Associate Secretary of the Board.

[FR Doc. 01–3204 Filed 1–2–01; 4:09 pm.

BILLING CODE 6210–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Guide to Community Preventive Services Task Force Meeting; Notice

Office of the Director, Centers for Disease Control and Prevention (CDC), announces the following meeting:

Name: Guide to Community Preventive Services (GCPS) Task Force Meeting.

Times and Dates: 9 a.m.–6 p.m., February 7, 2001. 9 a.m.–4 p.m., February 8, 2001.

Place: The Westin Peachtree Plaza, 210 Peachtree Street, Atlanta, Georgia 30303–1745, telephone (404) 659–1400.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 40 people.

Purpose: The mission of the Task Force is to develop and publish a Guide to Community Preventive Services, which is based on the best available scientific evidence and current expertise regarding essential public health services and what works in the delivery of those services.

Matters To Be Discussed: Agenda items include: presentation of recommendations for approval for the following chapters: Cancer, Motor Vehicle Occupant Injury, Physical Activity, and Sociocultural Environment; presentation of the dissemination/implementation/evaluation plan, discussions on the expansion and update of the Vaccine Preventable Disease Chapter; and general updates on the following information: Methods, Clinical Guide, and Alcohol, Diabetes, Mental Health, Sexual Behavior, and Violence Prevention Chapters.

Agenda items are subject to change as priorities dictate.

CONTACT PERSON FOR ADDITIONAL INFORMATION: Stephanie Zaza, M.D., M.P.H., Chief, Community Guide Branch, Division of Prevention Research and Analytic Methods, Epidemiology Program Office, CDC, 4770 Buford Highway, M/S K–73, Atlanta, Georgia 30341, telephone 770/488–8189.

Persons interested in reserving a space for this meeting should call 770/488–8189 by close of business on February 6, 2001.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: January 31, 2001.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 01–3022 Filed 2–5–01; 8:45am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01F–0047]

The National Fisheries Institute; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the National Fisheries Institute has filed a petition proposing that the food additive regulations be amended to provide for the safe use of ionizing radiation for control of foodborne pathogens in crustaceans and processed crustaceans.

FOR FURTHER INFORMATION CONTACT: Lane A. Highbarger, Center for Food Safety and Applied Nutrition (HFS–

206), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3032.

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 1M4727) has been filed by the National Fisheries Institute, 1901 North Fort Myer Dr., Arlington, VA 22209. The petition proposes to amend the food additive regulations in Part 179 *Irradiation in the Production, Processing and Handling of Food* (21 CFR part 179) to provide for the safe use of ionizing radiation for control of foodborne pathogens in raw-, frozen-, cooked-, partially cooked-, shelled-, or dried-crustaceans, or cooked- or ready-to-cook crustaceans processed with batter, breading, spices, or small amounts of other food ingredients.

The agency has determined under 21 CFR 25.32(j) that this action is of a 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: January 11, 2001.

Alan M. Rulis,

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

[FR Doc. 01–3095 Filed 2–5–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00N–1672]

Ashford Blood Bank, Inc.; Opportunity for Hearing on a Proposal to Revoke U.S. License No. 0740–001

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for a hearing on a proposal to revoke the establishment license (U.S. License No. 0740–001) and product licenses issued to Ashford Blood Bank, Inc., for the manufacture of Whole Blood and Red Blood Cells. The proposed revocation is based on the fact that authorized FDA employees have been unable to gain access to either of the establishment's locations for the purpose of carrying out a required inspection of the facility and that the manufacturing of products has been discontinued to an extent that a meaningful inspection or evaluation cannot be made.

DATES: The firm may submit written requests for a hearing by March 8, 2001, and any data and information justifying a hearing by April 9, 2001. Other interested persons may submit written comments on the proposed revocation by April 9, 2001.

ADDRESSES: Submit written requests for a hearing, any data and information justifying a hearing, and any written comments on the proposed revocation to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Joseph L. Okrasinski, Center for Biologics Evaluation and Research (HFM–17), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–827–6210.

SUPPLEMENTARY INFORMATION: FDA is initiating proceedings to revoke the establishment license (U.S. License No. 0740–001) and product licenses issued to Ashford Blood Bank, Inc., Ashford Medical Center, suite 401–402, Santurce, PR 00907, for the manufacture of Whole Blood and Red Blood Cells. Proceedings to revoke the licenses are being initiated because: (1) Authorized FDA employees have been unable to gain access to either of the establishment's locations for the purpose of carrying out a required inspection of the facility, and (2) manufacturing of products has been discontinued to an extent that a meaningful inspection or evaluation cannot be made.

In a certified return-receipt letter dated October 28, 1997, FDA notified an authorized official of the firm that FDA had suspended the firm's establishment and product licenses for the manufacture of Whole Blood and Red Blood Cells at its facilities at Santurce, PR, and Bayamon, PR. This action was based on the fact that significant deviations from the regulations were noted by FDA's San Juan district office during inspections of the facilities conducted August 19, 1997, through September 17, 1997, and September 9, 1997, through September 17, 1997, respectively. FDA's San Juan district office attempted to conduct additional inspections of the two Ashford facilities. On May 1, 1998, FDA investigators attempted to inspect the satellite collection facility at Bayamon, PR, but found that the facility was no longer in operation, and the manufacturing of Whole Blood and Red Blood Cells had been discontinued. On November 23, 1999, FDA investigators attempted to inspect the main facility in Santurce, PR, but found that the facility was no longer in operation and the manufacturing of Whole Blood and Red Blood Cells had been discontinued.

In certified, return-receipt letters dated April 13, 2000, sent to the firm's facility at Santurce, PR, and also to the Ashford Blood Bank, Inc., P.O. Box 195034, San Juan, PR, 00919, FDA notified an authorized official of the firm that FDA's attempts to conduct inspections of the two facilities at Santurce, PR and Bayamon, PR were unsuccessful because the facilities were no longer in operation and the manufacture of Whole Blood and Red Blood Cells had been discontinued. The letter also advised the authorized official that, under 21 CFR 601.5(b)(1) and (b)(2) (now codified as 21 CFR 601.5(b)(1)(i) and (b)(1)(ii)), when FDA finds that authorized employees have been unable to gain access to an establishment for the purpose of carrying out an inspection under 21 CFR 600.21, or the manufacturing of products or of a product has been discontinued to an extent that a meaningful inspection cannot be made, the Commissioner of Food and Drugs (the Commissioner) shall institute proceedings for license revocation. In the same letter, FDA stated that a meaningful inspection could not be made at the establishment and notified the firm of FDA's intent to revoke U.S. License No. 0740–001 and its intent to offer an opportunity for a hearing.

Because FDA has made reasonable efforts to notify the firm of the proposed revocation and has not received any response from the firm to the revocation letter, FDA is proceeding under 21 CFR 12.21(b) and publishing this notice of opportunity for a hearing on a proposal to revoke the licenses of the previously mentioned firm.

FDA has placed copies of the documents relevant to the proposed revocation on file with the Dockets Management Branch (address above) under the docket number found in brackets in the heading of this notice. These documents include: (1) Summary of Findings, May 1, 1998; (2) memorandum regarding FDA visit to Santurce location, November 23, 1999; and (3) FDA letters to the authorized official dated October 28, 1997, and April 13, 2000. These documents are available for public examination in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

Ashford Blood Bank, Inc., may submit a written request for a hearing to the Dockets Management Branch by March