

FEDERAL RESERVE SYSTEM**Change in Bank Control Notices; Acquisition of Shares of Bank or Bank Holding Companies**

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the office of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than April 9, 2001.

A. Federal Reserve Bank of Chicago (Phillip Jackson, Applications Officer) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. *Ann Manning*, Minneapolis, Minnesota; *Elizabeth Edwards Manning*, Bettendorf, Iowa; *Albert Manning* and *Kim Manning*, both of Holt, Michigan; *David Manning* and *Janet Manning*, both of West Des Moines, Iowa; *John Manning*, Lynnette Manning, George Manning, Ona Manning, Matthew Manning and *Judith Manning*, all of Keosauqua, Iowa; to retain ownership of Van Buren Bancorporation, Keosauqua, Iowa, and thereby indirectly retain additional voting shares of Community First Bank, Keosauqua, Iowa.

Board of Governors of the Federal Reserve System, March 20, 2001.

Robert deV. Frierson

Associate Secretary of the Board.

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BILLING CODE 6210-01-S

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Office of the Secretary****Meeting of the National Human Research Protections Advisory Committee**

AGENCY: Office of Public Health and Science, Office for Human Research Protections, DHHS.

ACTION: Notice of second meeting.

SUMMARY: Pursuant to section 10(d) of the Federal Advisory committee Act, as amended (5 U.S.C. appendix 2), notice

is hereby given of a meeting of the National Human Research Protections Advisory Committee.

The meeting will be open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the contact person listed below. Individuals planning on attending the meeting and who want to ask questions must submit their requests in writing in advance of the meeting to the contact person listed below.

DATES: The Committee will hold its next meeting on April 9-10, 2001. The meeting will convene from 8:30 a.m. to its recess at 5:30 p.m. on April 9 and resume at 8:30 a.m. to 5 p.m. EST on April 10.

ADDRESSES: Bethesda Marriott—Pooks Hill Road, Bethesda, Maryland 20814, (301) 897-9400.

FOR FURTHER INFORMATION CONTACT: Ms. Kate-Louise Gottfried, Executive Director, National Human Research Protections Advisory Committee, Office for Human Research Protections, 6100 Executive Boulevard, Room 310B (MSC 7507), Rockville, Maryland 20892-7507, (301) 496-7005. The electronic mail address is: kg123a@nih.gov.

SUPPLEMENTARY INFORMATION: The National Human Research Protections Advisory Committee was established on June 6, 2000 to provide expert advice and recommendations to the Secretary of HHS, Assistant Secretary for Health, the Director, Office for Human Research Protections, and other departmental officials on a broad range of issues and topic pertaining to or associated with the protection of human research subjects.

The draft meeting agenda for the second meeting of this committee is below. Updates to this agenda will be posted on the NHRPAC website at: <http://ohrp.osophs.dhhs.gov/nhrpac/nhrpac.htm>.

Draft Agenda

Monday April 9, 2001

8:30 a.m.-9:15 a.m. Welcome: (45 minutes)

Introduction of New Members (brief comments about yourself; areas of particular concern) (5 minutes each)
Mary Faith Marshall, Ph.D.,
Chairperson NHRPAC

Office for Human Research Protection Updates (10 minutes)

Greg Koski, Ph.D., M.D., Director of OHRP, Executive Secretary, NHRPAC, Chairman, HSRS

Clarification of NHRPAC Roles (10

minutes)

Mary Faith Marshall, Ph.D.

9:15 a.m.-9:30 a.m. The Department's Commitment to Protection of Human Subjects (15 minutes)

The Honorable David Satcher, M.D., Ph.D., Surgeon General (Invited)

9:30 a.m.-11 a.m. Update: Financial Relationships (1 hour, 30 minutes)
Mark Barnes, J.D., Chair, Working Group

Stuart L. Nightingale, M.D., Senior Medical Advisor to the Assistant Secretary for Planning & Evaluation

10:30 a.m.-10:45 a.m. Break (15 minutes)

11 a.m.-11:45 a.m. Update: Declaration of Helsinki (45 minutes)

Greg Koski, Ph.D., M.D.

Stuart L. Nightingale, M.D.

11:45 a.m.-12:15 p.m. Public Comment (30 minutes)

12:15 p.m.-1:30 p.m. Lunch—On your own

1:30 p.m.-4:30 p.m. Genetics

1:30 p.m.-2:15 p.m. Genetic Research: An overview (45 minutes)

Francis Collins, M.D., Ph.D., director, National Human Genome Research Institute

2:15 p.m.-4:30 p.m. Panel Discussion Moderator, Francis Collins, M.D., Ph.D.

Family Members:

Should Family Members of Survey Subjects, Themselves become Subjects of a Protocol—if so, Must Informed Consent be Obtained for Investigator to Retain Private Information on these Individuals?

2:15 p.m.-3:15 p.m. Guest Panel: (1 hour)

Jeff Botkin, M.D., M.P.H., Department of Pediatrics, The University of Utah Medical Center (15 minutes)

Terry Arledge, Ph.D.,

GalaxoSmithKline (15 minutes)

Sharon Terry, Genetic Alliance (15 minutes)

Terry Seargent (15 minutes)

3:15 p.m.-3:30 p.m. Break (15 minutes)

3:30 p.m.-4:30 p.m. Discussion (45 minutes)

4:30 p.m.-5:15 p.m. Public Comment (1 hour)

5:15 p.m.-5:30 p.m. Closing Comments/Adjourn (15 minutes)

Tuesday, April 10, 2001

8:30 a.m.-8:45 a.m. Brief Recap of Day One (15 minutes)

Questions/Clarifications

Mary Faith Marshall, Ph.D.

8:45 a.m.-9:00 a.m. The National

Institutes of Health and Human

Subject Protections (15 minutes)

Ruth Kirschstein, M.D., Acting

Director, National Institutes of Health

9:00 a.m.–12:00 p.m. Children (3 hours)
 9:00 a.m.–9:45 a.m. Discussion of
 Current Definitions and their
 Interpretation (45 minutes)
 NHRPAC Committee
 9:45 a.m.–10:15 a.m. Children's
 Workgroup (30 minutes)
 Alan Fleischman, M.D., Senior Vice
 President, NY Academy of
 Medicine, Clinical Professor of
 Pediatrics and Clinical Professor of
 Epidemiology & Social Medicine,
 Albert Einstein College, New York
 10:15 a.m.–10:30 a.m. Break (15
 minutes)
 10:30 a.m.–11:45 a.m. Committee
 Discussion (1 hour, 15 minutes)
 11:45 a.m.–12:00 p.m. The National
 Science Foundation and Human
 Subject Protections (15 minutes)
 Rita Colwell, Ph.D., Director, National
 Science Foundation
 12:00 p.m.–1:30 p.m. Lunch—on your
 own (1 hour, 30 minutes)
 1:30 p.m.–3:00 p.m. Update: Social
 Science (1 hour, 30 minutes)
 Felice Levine, Ph.D., Executive
 Officer, American Sociological
 Association (30 minutes)
 Jeff Cohen, Ph.D., Director, Education,
 OHRP
 Discussion (1 hour)
 3:00 p.m.–3:15 p.m. Break (15 minutes)
 3:15 p.m.–4:15 p.m. Public Comment (1
 hour)
 4:15 p.m.–5:00 p.m. Meeting Recap (45
 minutes)
 Review Recommendations:
 Financial Relationships
 Declaration of Helsinki
 Genetics
 Secondary Subjects
 Children
 Social Science
 Mary Faith Marshall, Ph.D.
 5:00 p.m. Thank You—Adjourn
 Dated: March 20, 2001.

Greg Koski,

*Director, Office for Human Research
 Protections.*

[FR Doc. 01-7443 Filed 3-23-01; 8:45 am]

BILLING CODE 4160-17-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Program Announcement 01041]

Exposure to Tremolite Asbestos in Vermiculite Ore Site-Specific Health Activities; Notice of Availability of Funds

A. Purpose

The Agency for Toxic Substances and
 Disease Registry (ATSDR) announces

the availability of fiscal year (FY) 2001
 funds for a cooperative agreement
 program to conduct site-specific health
 activities related to human exposure to
 contaminated vermiculite ore at sites
 around the United States that received
 and/or processed ore from the mine in
 Libby, Montana.

This program addresses the "Healthy
 People 2010" focus area of
 Environmental Health. The purpose of
 the program is to assist public health
 agencies in conducting site-specific
 health activities related to human
 exposure to contaminated vermiculite
 ore at sites identified by the
 Environmental Protection Agency (EPA)
 as receiving and/or processing ore.

B. Eligible Applicants

Assistance will be provided only to
 public health agencies of States or their
 bona fide agents or instrumentalities.
 State organizations, including State
 universities, must establish that they
 meet their respective State legislature's
 definition of a State entity or political
 subdivision to be considered an eligible
 applicant.

Note: Public Law 104-65 states that an
 organization described in section 501(c)(4) of
 the Internal Revenue Code of 1986 that
 engages in lobbying activities is not eligible
 to receive Federal funds constituting an
 award, grant, cooperative agreement,
 contract, loan, or any other form.

C. Availability of Funds

Approximately \$1,000,000 is available
 in FY 2001 to fund approximately 10
 awards. It is expected that the average
 award will range from a maximum of
 \$10,000 for the conduct of health
 statistics reviews to a maximum of
 \$500,000 for epidemiologic
 investigations. It is expected that the
 awards will begin on or about either
 July 1, 2001 and will be made for a 12-
 month budget period within a project
 period of up to 3 years. Funding
 estimates may change.

Continuation awards within an
 approved project period will be made
 on the basis of satisfactory progress as
 evidenced by required reports and the
 availability of funds.

Use of Funds

Funds may be expended for
 reasonable program purposes, such as
 personnel, travel, supplies, and services.
 Funds for contractual services may be
 requested; however, the grantee, as the
 direct and primary recipient of PHS
 grant funds, must perform a substantive
 role in carrying out project activities
 and not merely serve as a conduit for an
 award to another party or provide funds

to an ineligible party. Funds may not be
 used to purchase equipment.

D. Program Requirements

In conducting activities to achieve the
 purpose of this program, the recipient
 will be responsible for activities under
 1. Recipient Activities, and ATSDR will
 be responsible for the activities listed
 under 2. ATSDR Activities.

1. Recipient Activities

a. For health statistics reviews:
 Analyze existing health outcome data of
 select asbestos-related diseases.
 Mortality data will be the most readily
 available data for asbestos-related
 diseases such as mesothelioma, lung
 cancer, and asbestosis, although cancer
 registry data should be utilized where
 available.

b. For epidemiologic investigations:
 Develop a protocol and conduct the
 recommended investigation. This
 protocol will undergo scientific peer
 review as required by ATSDR.

c. Provide proof by citing a State code
 or regulation or other State
 pronouncement under authority of law,
 that medical information obtained
 pursuant to the agreement will be
 protected from disclosure when the
 consent of the individual to release
 identifying information is not obtained.

d. Develop a mechanism for ongoing
 interaction with, and education of
 affected community.

2. ATSDR Activities

a. For the health statistics review:
 Make available to states both technical
 assistance and a standard protocol to
 use to analyze existing health outcome
 data of select asbestos-related diseases.

b. For epidemiologic investigations:
 Provide consultation and assist in
 monitoring the data; participate if
 requested in the study analysis and
 collaborate, if requested, in interpreting
 the study findings.

c. Conduct technical and peer review.

E. Application Content

In a narrative form, the application
 should include a discussion of areas
 under the "Evaluation Criteria" section
 of this announcement as they relate to
 the proposed program. These criteria
 serve as the basis for evaluating the
 application; therefore, omissions or
 incomplete information may affect the
 rating of the application. This program
 does not require in-kind support or
 matching funds, however, the applicant
 should describe any in-kind support in
 the application.

The narrative should be no more than
 30 pages, double-spaced, printed on
 one-side, with 1" margins, and