

Dated: April 13, 1995.

Stephen L. Johnson,
*Director, Registration Division, Office of
Pesticide Programs.*

[FR Doc. 95-10255 Filed 4-25-95; 8:45 am]

BILLING CODE 6560-50-F

FEDERAL COMMUNICATIONS COMMISSION

[Report No. 2067]

Petition for Reconsideration of Actions in Rulemaking Proceedings

April 24, 1995.

Petition for reconsideration have been filed in the Commission rulemaking proceedings listed in this Public Notice and published pursuant to 47 CFR 1.429(e). The full text of this document are available for viewing and copying in Room 239, 1919 M Street, N.W., Washington, D.C. or may be purchased from the Commission's copy contractor ITS, Inc. (202) 857-3800. Opposition to this petition must be filed May 11, 1995. See § 1.4(b)(1) of the Commission's rules (47 CFR 1.4(b)(1)). Replies to an opposition must be filed within 10 days after the time for filing oppositions has expired.

Subject: Review of the Pioneer's
Preference Rules. (ET Docket No.
93-266)

Number of Petition Filed: 1.

Federal Communications Commission.

William F. Caton,

Acting Secretary.

[FR Doc. 95-10211 Filed 4-25-95; 8:45 am]

BILLING CODE 6712-01-M

FEDERAL MARITIME COMMISSION

[Petition No. P2-95]

Household Goods Forwarders Association of America, Inc.; Petition for Exemption From Tariff and Bonding Requirements in Regard to Household Goods and Personal Effects for the Account of the General Services Administration; Notice of Filing

Notice is hereby given that the Household Goods Forwarders Association of America, Inc. ("Petitioner") has petitioned for an exemption pursuant to Section 16 of the Shipping Act of 1984 [46 U.S.C. app. 1715] and Section 35 of the Shipping Act, 1916 [46 U.S.C. app. 833a]. Petitioner seeks an exemption for non-vessel operating common carriers by water from the tariff filing requirements of Part 514 and the bonding requirement

of Part 583 of Title 46 CFR, to the extent they engage in the transportation of used household goods and personal effects of personnel of federal civilian executive agencies in the domestic and foreign commerce of the United States, pursuant to a solicitation issued and administered by the General Services Administration of the United States.

In order for the Commission to make a thorough evaluation of the petition for exemption, and the proposed CFR amendments suggested therein, interested persons are requested to submit views or arguments in reply to the petition no later than May 25, 1995. Replies shall be directed to the Secretary, Federal Maritime Commission, Washington, DC 20573-0001 in an original and 15 copies.

Replies shall also be served on Alan F. Wohlstetter, Denning & Wohlstetter, 1700 K Street NW., Washington, DC 20006.

Copies of the petition are available for examination at the Washington, DC office of the Commission, 800 N. Capitol St NW., Room 1046.

Joseph C. Polking,
Secretary.

[FR Doc. 95-10173 Filed 4-25-95; 8:45 am]

BILLING CODE 6730-01-M

FEDERAL RESERVE SYSTEM

Dalrymple Family Limited Partnership, L.P., et al.; Change in Bank Control Notices; Acquisitions of Shares of Banks 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. Once the notices have been accepted for processing, they will also be available for inspection at the offices 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 May 10, 1995.

A. Federal Reserve Bank of Atlanta
(Zane R. Kelley, Vice President) 104 Marietta Street, N.W., Atlanta, Georgia 30303:

1. *Dalrymple Family Limited Partnership, L.P. and 2105 South Broadway Associates, L.P.*, both of

Elmira, New York; each to acquire 4.92 percent of the voting shares of Chemung Financial Corporation, Elmira, New York, and thereby indirectly acquire Chemung Canal Trust Company, Elmira, New York.

Board of Governors of the Federal Reserve System, April 20, 1995.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 95-10228 Filed 4-25-95; 8:45 am]

BILLING CODE 6210-01-F

Republic Security Financial Corporation; Formation of, Acquisition by, or Merger of Bank Holding Companies

The company listed in this notice has applied for the Board's approval under section 3 of the Bank Holding Company Act (12 U.S.C. 1842) and § 225.14 of the Board's Regulation Y (12 CFR 225.14) to become a bank holding company or to acquire a bank or bank holding company. The factors that are considered in acting on the applications are set forth in section 3(c) of the Act (12 U.S.C. 1842(c)).

The application is available for immediate inspection at the Federal Reserve Bank indicated. Once the application has been accepted for processing, it will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that application or to the offices of the Board of Governors. Any comment on an application that requests a hearing must include a statement of why a written presentation would not suffice in lieu of a hearing, identifying specifically any questions of fact that are in dispute and summarizing the evidence that would be presented at a hearing.

Comments regarding this application must be received not later than May 19, 1995.

A. Federal Reserve Bank of Atlanta
(Zane R. Kelley, Vice President) 104 Marietta Street, N.W., Atlanta, Georgia 30303:

1. *Republic Security Financial Corporation*, West Palm Beach, Florida; to become a bank holding company by acquiring 100 percent of the voting shares of Republic Security Bank, West Palm Beach, Florida.

Board of Governors of the Federal Reserve System, April 20, 1995.

Jennifer J. Johnson,

Deputy Secretary of the Board.

[FR Doc. 95-10229 Filed 4-25-95; 8:45 am]

BILLING CODE 6210-01-F

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

[Announcement 533]

National Institute for Occupational Safety and Health; Cooperative Agreement for Model Program for Occupational Respiratory Disease Evaluation and Rehabilitation**Introduction**

The Centers for Disease Control and Prevention (CDC) announces the availability of fiscal year (FY) 1995 funds for a cooperative agreement program for Occupational Respiratory Disease Evaluation and Rehabilitation through the National Institute for Occupational Safety and Health (NIOSH). 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 Healthy People 2000 see the Section WHERE TO OBTAIN ADDITIONAL INFORMATION.)

Authority

This program is authorized under Sections 20(a) and 21(a) of the Occupational Safety and Health Act of 1970 and Section 501(a) of the Federal Mine Safety and Health Act (29 U.S.C. 669(a) and 670(a); 30 U.S.C. 951(a)).

Smoke-Free Workplace

PHS strongly encourages all grant recipients to provide a smoke-free workplace and promote the non-use of all tobacco products, and Public Law 103-227, the Pro-Children Act of 1994, prohibits smoking in certain facilities that receive Federal funds in which education, library, day care, health care, and early childhood development services are provided to children.

Eligible Applicants

Applications may be submitted by public and private, non-profit and for-profit organizations, and governments and their agencies. Thus, universities, colleges, research institutions, hospitals, other public and private organizations, State and local health departments or their bona fide agents, federally recognized Indian tribal governments, Indian tribes or Indian tribal organizations, and small, minority- and/or women-owned businesses are eligible to apply.

Availability of Funds

Approximately \$275,000 will be available in FY 1995 to fund one to two awards. The award(s) is expected to begin on or about September 30, 1995, for a 12-month budget period within a project period of three to five years. Funding estimates may vary and are subject to change. Continuation awards within the project period will be made on the basis of satisfactory progress and the availability of funds.

Purpose

The purpose of this occupational respiratory disease cooperative agreement is to assist in the development, implementation, and maintenance of a model program for the diagnosis, evaluation, and rehabilitation of individuals with occupational respiratory disease. This program may build on existing expertise of an institution or provide assistance in initiating a new program. This program will report and disseminate findings, as well as relevant health and safety education and training information, to State health officials, health-care providers, workers, managers, unions, and employers. This program will include an evaluation of current standard and innovative interventions for early identification of occupational respiratory diseases which results in recommendations for, or a plan for the development of, new methods and techniques to improve the early recognition, rehabilitation and therapy of these diseases. The evaluation component built into each project should include carefully developed, realistic and appropriate evaluation tools. The evaluation results will be used to modify and improve ongoing program plans.

The objectives of the occupational respiratory disease evaluation and rehabilitation program are as follows:

1. To assist an institution in the development or refinement of a program for evaluation and rehabilitation of occupational respiratory disease.
2. To provide the opportunity for an institution to evaluate the effectiveness of a model program for evaluation and rehabilitation of occupational respiratory disease.
3. To provide a collaborative focus for occupational health expertise in occupational respiratory disease.
4. Contribute to a better understanding of occupational respiratory diseases.
5. Ultimately reduce the morbidity, mortality, and social and economic burden of occupational respiratory diseases in the United States.

Program Requirements

In conducting activities to achieve the purpose of this program, the recipient shall be responsible for conducting activities under A. (Recipient Activities) below, and CDC/NIOSH will be responsible for conducting activities under B. (CDC/NIOSH Activities) below:

A. Recipient Activities

1. Identify a director for the program (or each program component).
2. Develop a targeted list of occupational respiratory diseases to be evaluated such as (but not limited to) silicosis, coal workers' pneumoconiosis, asbestosis, occupational asthma, hypersensitivity pneumonitis, organic dust diseases, and acute toxic respiratory injuries
3. Develop and conduct a model program for the early recognition, evaluation, diagnosis, rehabilitation, and therapy of occupational respiratory diseases.
4. Report and disseminate information on the organization, activities, and findings of the model program, as well as relevant health and safety education and training information to State and Federal health officials, health-care providers, workers, managers, unions, and employers.
5. Work with State and Federal disability compensation programs to identify and enroll workers who could be offered diagnosis, evaluation and rehabilitation of occupational respiratory diseases.
6. Develop a protocol(s) for the evaluation and rehabilitation of occupational respiratory diseases. Obtain peer review of the protocol(s); revise and finalize, as required, for final approval; evaluate the effectiveness of the protocol(s). Disseminate the results of these efforts to other health-care institutions evaluating, diagnosing, and rehabilitating workers with occupational respiratory diseases.
7. Develop new methods and techniques that improve the early recognition and rehabilitation of workers with occupational respiratory diseases.

B. CDC/NIOSH Activities

1. Provide technical assistance through site visits and correspondence for the development and implementation of the model program.
2. Provide scientific collaboration for the model program.
3. Provide limited professional assistance during the conduct of the program including, but not limited to, physicians, nurses, epidemiologists, statisticians, industrial hygienists and laboratory scientists.

4. Participate in peer review of the project protocol(s).

5. Provide technical assistance in all phases of development, implementation, and maintenance of the cooperative agreement and collaborative project activities.

6. Assist in reporting and disseminating findings as well as relevant health and safety education and training information to state health officials, health-care providers, workers, managers, unions, and employers.

Evaluation Criteria

Applications will be reviewed and evaluated according to the following criteria:

1. The applicant's understanding of the objectives of the proposed initiative. (5%)
2. Proposed schedule for initiating and accomplishing the activities of the cooperative agreement. (10%)
3. Responsiveness of the proposal to the scope and objectives described in the Announcement. (25%)
4. Technical merit and originality of the program proposal. (30%)
5. Training and experience of the proposed Program Director(s) and staff. The Program Director(s) must be a recognized scientist and technical expert, and must assume and provide assurances of major time commitment to the program. (10%)
6. Suitability of the facilities to conduct the program. (15%)
7. Evidence of plans for creative collaboration and coordination with local resources which could facilitate identification, evaluation or rehabilitation of workers with occupational respiratory diseases, including establishment of working relationships with State and Federal disability programs. (5%)
8. Extent to which the budget is reasonable, clearly justified, and consistent with the use of funds. (Not Scored)

Executive Order 12372 Review

Applications are subject to Intergovernmental Review of Federal Programs as governed by Executive Order (E.O.) 12372. E.O. 12372 sets up a system for State and local government review of proposed Federal assistance applications. Applicants (other than federally recognized Indian tribal governments) should contact their State Single Point of Contact (SPOC) as early as possible to alert them to the prospective applications and receive any necessary instructions on the State process. For proposed projects serving more than one State, the applicant is advised to contact the SPOC for each

affected State. Indian tribes are strongly encouraged to request tribal government review of the proposed application. A current list of SPOCs is included in the application kit.

If SPOCs or tribal governments have any State process recommendations on applications submitted to CDC, they should send them to Henry S. Cassell, III, Grants Management Officer, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Atlanta, GA 30305, no later than 60 days after the application deadline date. The Program Announcement number and Program Title should be referenced on the document. The granting agency does not guarantee to "accommodate or explain" State process recommendations it receives after that date.

Public Health System Reporting Requirements

This program is not subject to the Public Health System Reporting Requirements.

Catalog of Federal Domestic Assistance Numbers

The Catalog of Federal Domestic Assistance Number for this program is 93.262.

Other Requirements

Paperwork Reduction Act

Projects that involve the collection of information from 10 or more individuals and funded by this cooperative agreement will be subject to approval by the Office of Management and Budget (OMB) under the Paperwork Reduction Act.

Human Subjects

This program involves research on human subjects; therefore, all applicants must comply with the Department of Health and Human Services Regulations, 45 CFR part 46, regarding the protection of human subjects. Assurance must be provided which demonstrates the project or activity will be subject to initial and continuing review by an appropriate institutional review committee.

In addition to other applicable committees, Indian Health Service (IHS) institutional review committees also must review the project if any component of IHS will be involved or will support the research. If any American Indian community is involved, its tribal government must also approve that portion of the project applicable to it.

The applicant will be responsible for providing assurances in accordance

with the appropriate guidelines and forms provided in the application kit.

Application Submission and Deadline

The original and two copies of the application PHS Form 5161-1 (OMB Number 0937-0189) must be submitted to Henry S. Cassell III, Grants Management Officer, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 300, Mailstop E13, Atlanta, GA 30305, on or before June 22, 1995.

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 objective review group. (Applicants must request a legibly dated U.S. Postal Service postmark or obtain a legibly dated receipt from a commercial carrier or U.S. Postal Service. Private metered postmarks shall not be acceptable as proof of timely mailings.)

2. Late Applications: Applications which do not meet the criteria in 1.(a) or 1.(b) above are considered late applications. Late applications will not be considered in the current competition and will be returned to the applicant.

Where To Obtain Additional Information

To receive additional written information call (404) 332-4561. You will be asked to leave your name, address, telephone number and will need to refer to Announcement 533.

You will receive a complete program description, information on application procedures, and application forms.

If you have questions after reviewing the contents of all the documents, business management technical assistance may be obtained from Oppie M. Byrd, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 300, Mailstop E-13, Atlanta, GA 30305, telephone (404) 842-6546. Programmatic technical assistance may be obtained from John E. Parker, M.D., National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC), 1095 Willowdale Road, Morgantown, WV 26505-2888, telephone (304) 285-5724.

Please refer to Announcement Number 533 when requesting information and submitting an application.

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) referenced in the Introduction through the Superintendent of Documents, Government Printing Office, Washington, DC 20402-9325, telephone (202) 512-1800.

Dated: April 20, 1995.

Diane D. Porter,

Acting Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC).

[FR Doc. 95-10197 Filed 4-25-95; 8:45 am]

BILLING CODE 4163-19-P

National Committee on Vital and Health Statistics (NCVHS) Executive Subcommittee: Meeting

Pursuant to Pub. L. 92-463, the National Center for Health Statistics (NCHS), Centers for Disease Control and Prevention (CDC), announces the following committee meeting.

Name: NCVHS Executive Subcommittee.
Time and Date: 8:30 a.m.-5 p.m., May 24, 1995.

Place: Suite 200 East, Conference Room 002-003, 1100 New York Avenue, NW., Washington, DC 20005.

Status: Open.

Purpose: The purpose of this meeting is for the Executive Subcommittee to review accomplishments, logistics, needs and work plans of NCVHS and individual subcommittees.

Contact Person for More Information: Substantive program information as well as summaries of the meeting and a roster of committee members may be obtained from Gail F. Fisher, Ph.D., Executive Secretary, NCVHS, NCHS, CDC, Room 1100, Presidential Building, 6525 Belcrest Road, Hyattsville, Maryland 20782, telephone 301/436-7050.

Dated: April 20, 1995.

Carolyn J. Russell,

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

[FR Doc. 95-10198 Filed 4-25-95; 8:45 am]

BILLING CODE 4163-18-M

Food and Drug Administration

[Docket No. 95N-0013]

Benton County Ag Center, Inc.; Proposal to Withdraw Approval of Applications for Medicated Animal Feeds; Opportunity for a Hearing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Center for Veterinary Medicine (CVM), Food and Drug Administration (FDA), is providing an opportunity for a hearing on a proposal to withdraw approval of certain medicated feed applications (MFA's) held by Benton County Ag Center, Inc., for animal feeds bearing or containing new animal drugs (NAD's). This action is based on new information showing the firm's methods and controls used for manufacturing, processing, and packing of the medicated feeds are inadequate to assure and preserve the identity, strength, quality, and purity of the NAD's therein, and they were not made adequate within a reasonable time after receipt of written notice from FDA.

DATES: Requests for a hearing and data and information in support of the hearing request are due by May 26, 1995.

ADDRESSES: Requests for a hearing in response to this notice should be identified with Docket No. 95N-0013 and sent to the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Karen A. Kandra, Center for Veterinary Medicine (HFV-246), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-594-1765.

SUPPLEMENTARY INFORMATION: CVM is providing an opportunity for a hearing on a proposal to withdraw approval of 11 MFA's held by the firm doing business as Benton County Ag Center, Inc., 312 Railroad St., P.O. Box 308, Keystone, IA 52249-0308, for the manufacture of animal feeds bearing or containing Category II NAD's. Benton County Ag Center, Inc., is a feed mill that manufactures both medicated and nonmedicated animal feeds. The 11 MFA's, held by Benton County Ag Center, Inc., were approved under section 512(m) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360b(m)) and are identified as follows:

MFA Number	Drug/Combination	Species
1. F 93-642.	Carbadox	Swine
2. F 127-333.	Tylosin/Sulfamethazine.	Swine
3. F 131-878.	Carbadox	Swine
4. F 139-280.	Levamisole Hydrochloride.	Cattle and swine
5. F 141-603.	Carbadox/Pyrantel tartrate.	Swine
6. F 141-604.	Pyrantel tartrate	Swine

MFA Number	Drug/Combination	Species
7. F 141-757.	Lincomycin/Pyrantel tartrate.	Swine
8. F 144-054.	Sulfamethazine/Chlortetracycline (CTC)/Penicillin.	Swine
9. F 147-607.	Sulfamethazine/CTC.	Cattle
10. F 147-617.	Arsanilic acid	Chickens, turkeys, and swine
11. F 147-641.	Oxytetracycline/Neomycin.	Chickens, turkeys, swine, cattle, and mink

To manufacture a Type B or C animal feed bearing or containing a Category II NAD (i.e., Type A medicated article) a firm must file an MFA (Form FDA 1900) with FDA and obtain its approval. FDA does not approve such an application unless, among other things, the firm agrees to comply with the agency's regulations for current good manufacturing practice (CGMP) for medicated feeds (21 CFR part 225), which are intended to help assure that feed bearing or containing an NAD meets the requirements of the act pertaining to identity, strength, quality, and purity. The agency determines whether the firm's manufacture of medicated feed is in compliance with the CGMP regulations by inspecting the facilities and controls used for, and the methods used in, the manufacture, processing, and packing of the feed by the firm.

On December 22, 1992, the Iowa Department of Agriculture (Medicated Feed Bureau), under contract with FDA pursuant to section 702(a) of the act (21 U.S.C. 372(a)), inspected Benton County Ag Center, Inc. The inspection revealed significant deviations from the CGMP's for medicated feeds. The investigator noted the deviations on an inspectional observations form (Form FDA 483) (Ref. 1), issued a copy to the firm's General Manager, and discussed in detail the deviations with him. The deviations included the following:

1. Production records did not show when flushing of equipment was performed or the final disposition of flush materials, as required by 21 CFR 225.102(b)(4).

2. Production records did not show the actual quantity of medicated feed produced, as required by 21 CFR 225.102(b)(2)(iv).

3. Production records were not checked by a responsible person to determine if all required production steps had been performed, as required by 21 CFR 225.102(b)(4).