FEDERAL MARITIME COMMISSION

Notice of Agreements Filed

The Commission hereby gives notice of the filing of the following agreements under the Shipping Act of 1984. Interested parties may submit comments on the agreements to the Secretary, Federal Maritime Commission, Washington, DC 20573, within twelve days of the date this notice appears in the **Federal Register**. Copies of the agreements are available through the Commission's website (*www.fmc.gov*) or by contacting the Office of Agreements at (202) 523–5793 or *tradeanalysis@fmc.gov*.

Agreement No.: 012356–001.

Title: Matson/Mell Space Charter Agreement (Pacific Islands).

Parties: Matson Navigation Company, Inc. and Mariana Express Lines Pte. Ltd.

Filing Party: David J. Tubman, Assistant General Counsel; Matson; 555 12th Street; Oakland, CA 94607.

Synopsis: The amendment updates the Agreement to reflect reciprocal space charter authority, and makes other administrative changes to the Agreement.

Agreement No.: 201250.

Title: Marine Terminal Services Agreement between Port of Houston Authority and Zim Integrated Shipping Services Ltd.

Parties: Port of Houston Authority and Zim Integrated Shipping Services Ltd.

Filing Party: Chasless Yancy; Port of Houston Authority; 111 East Loop North; Houston, TC 77029.

Synopsis: The Agreement sets forth certain discounted rates and charges applicable to Zim's container vessels calling at PHA's Barbours Cut and Bayport Container Terminals in the Port of Houston. The MTSA will commence upon filing with the Federal Maritime Commission, and the term of the MTSA is for 10 years following such filing, with an option to jointly agree upon a five-year extension.

Dated: May 14, 2018.

Rachel E. Dickon,

Secretary.

[FR Doc. 2018–10558 Filed 5–16–18; 8:45 am]

BILLING CODE 6731–AA–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

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 shares of 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 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 June 4, 2018.

A. Federal Reserve Bank of Dallas (Robert L. Triplett III, Senior Vice President) 2200 North Pearl Street, Dallas, Texas 75201–2272:

1. *Sid Ridlehuber, Corpus Christi, Texas; and Sid Ridlehuber, Corpus Christi, Texas, Ryan Ridlehuber, San Antonio, Texas, and Robyn Totah, Austin, Texas,* (together known as the Ridlehuber Family Group, a group acting in concert); to retain and acquire voting shares of Charter Bancshares, Inc., Corpus Christi, Texas, which controls Charter Bank, Corpus Christi, Texas.

Board of Governors of the Federal Reserve System, May 14, 2018.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2018–10575 Filed 5–16–18; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: How TANF Agencies Support Families Experiencing Homelessness.

OMB No.: New Collection.

Description: The Office of Planning, Research, and Evaluation (OPRE), Administration for Children and Families (ACF) at the U.S. Department of Health and Human Services (HHS) is conducting the, ''How TANF Agencies Support Families Experiencing Homelessness,'' project through a contract with Abt Associates in partnership with MEF Associates. This project will assist HHS in understanding the extent to which TANF agencies across the country are using TANF funds to serve and support families experiencing or are at-risk of homelessness. It also will document the approaches and strategies used by TANF agencies to serve these families. We are seeking OMB approval for four elements of the study: (1) The TANF Administrator Web Survey (tailored for both state and county respondents), (2) a Site Visit Discussion Guide for TANF staff, 3) a Site Visit Discussion Guide for Staff at Continuums of Care (CoC)/ Partner Organizations, and 4) a Site Visit Focus Group Guide.

TANF Administrator Web Survey. We will administer an online survey to all state and territory TANF administrators as well as a selection of three county TANF administrators from each state. The survey will collect information about the agencies' overall approaches toward addressing family homelessness and the extent to which TANF funds, assessments, tools, additional services, and partners are used in these efforts.

Discussion protocols during site visits to TANF agencies. The study team will visit five purposefully selected TANF agencies. During these two-day visits, the project staff will use the Site Visit Discussion Guide for TANF Staff to conduct interviews with TANF office staff, use the Site Visit Focus Group Guide to convene focus groups of TANF participants experiencing or at-risk of homelessness, and use the Site Visit Discussion Guide for Staff at CoC/ Partner Organizations to interview representatives from relevant homelessness organization partners, including CoCs.

Respondents: State, territory, and selected county TANF administrators; TANF agency staff who provide case management or services to address family homelessness; representatives from the local CoC, and as applicable, staff from other partner organizations that serve homeless families; TANF recipients experiencing or at-risk of homelessness.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
TANF Administrator Web Survey (State and County)	206	69	1	.5	35
Site Visit Discussion Guide for TANF Staff	50	17	1	1.5	26
Site Visit Discussion Guide for Staff at CoC/Partner Organizations	20	7	1	1.5	11
Site Visit Focus Group Guide	20	7	1	1.5	11

Estimated Total Annual Burden Hours: 83.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. Email address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–10550 Filed 5–16–18; 8:45 am]

BILLING CODE 4184–09–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–N–1561]

Anesthesiology and Respiratory Therapy Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Anesthesiology and Respiratory Therapy Devices Panel of the Medical Devices Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public.

DATES: The meeting will be held on June 14, 2018 from 8 a.m. to 6 p.m.

ADDRESSES: Gaithersburg Holiday Inn, Grand Ballroom, 2 Montgomery Village Ave., Gaithersburg, MD 20879. The hotel's telephone number is 301–948–8900. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FOR FURTHER INFORMATION CONTACT: Evella Washington, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. G640, Silver Spring, MD 20993–0002, *Evella.Washington@fda.hhs.gov,* 301–796–6683, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at *https://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

SUPPLEMENTARY INFORMATION:

Agenda: The committee will discuss, make recommendations and vote on information related to PneumRx, Inc.'s premarket approval application for the PNEUMRX ELEVAIR Endobronchial Coil System, which is a first of a kind implantable lung reduction coil for the proposed indication for use in patients with homogeneous and/or heterogeneous severe emphysema to improve quality of life, lung function, and exercise capacity.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's website after the meeting. Background material is available at *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before June 7, 2018. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those individuals interested in making formal oral presentations should notify the contact person 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 requested to make their presentation on or before May 30,