continued existence of the receivership will serve no useful purpose. Consequently, notice is given that the receivership shall be terminated, to be effective no sooner than thirty days after the date of this Notice. If any person wishes to comment concerning the termination of the receivership, such comment must be made in writing and sent within thirty days of the date of this Notice to: Federal Deposit Insurance Corporation, Division of Resolutions and Receiverships, Attention: Receivership Oversight Department 34.6, 1601 Bryan Street, Dallas, TX 75201.

No comments concerning the termination of this receivership will be considered which are not sent within this time frame.

Dated: March 2, 2017.

Federal Deposit Insurance Corporation.

Valerie J. Best,

Assistant Executive Secretary.

[FR Doc. 2017–04491 Filed 3–7–17; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL MARITIME COMMISSION

[Docket No. 17–02]

Notice of Filing of Complaint and Assignment; Hangzhou Qianwang Dress Co., Ltd. V. RDD Freight International Inc.

Notice is given that a complaint has been filed with the Federal Maritime Commission (Commission) by Hangzhou Qianwang Dress Co. Ltd., hereinafter ''Complainant,'' against RDD Freight International Inc., hereinafter ''Respondent.'' Complainant states it is a People's Republic of China Corporation that ''manufactures apparel, including hats and gloves, and sells it to retailers in the United States.'' Complainant alleges that Respondent is a Commission licensed non-vessel operating common carrier, an international freight forwarder (''FF'') and a New York corporation.

Complainant alleges that Respondent failed to wait for receipt of the original Bills of Lading or to get the Complainant's consent before releasing certain shipments of merchandise to the consignee. The consignee had not paid the Complainant for the merchandise at the time of release nor have they paid as of the date of this filing. By releasing the goods to the consignee, Complainant alleges that the Respondent ''fail[ed] to establish, observe and enforce just and reasonable regulations and practices relating to or connected with receiving, handling, storing, or delivering property'' which violates 46 U.S.C. 41102(c).

Complainant seeks reparations in the amount of $134,207.70, and other relief. The full text of the complaint can be found in the Commission's Electronic Reading Room at *www.fmc.gov/17-02/.*

This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding officer in this proceeding shall be issued by March 2, 2018, and the final decision of the Commission shall be issued by September 17, 2018.

Rachel E. Dickon,

Assistant Secretary.

[FR Doc. 2017–04511 Filed 3–7–17; 8:45 am]

BILLING CODE 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 March 23, 2017.

A. Federal Reserve Bank of St. Louis (David L. Hubbard, Senior Manager) P.O. Box 442, St. Louis, Missouri 63166–2034. Comments can also be sent electronically to *Comments.applications@stls.frb.org:*

1. *John Christopher Harlin, Trustee of the John Christopher Harlin Revocable Trust, John L. Harlin Trustee of the Hugh T. Harlin Trust, John L. Harlin, Trustee UTA John L. Harlin Trust, Linda K. Harlin Trustee UTA Linda K. Harlin Trust, Abigail Christen Harlin, Bradley J. Hardcastle Trustee of the Bradley J. Hardcastle Trust, all of Gainesville, Missouri, Lisa M. Gables of Clifton, Virginia, and Joe D. Hardcastle and B. Sherrill Hardcastle Trustees of the Joe D. Hardcastle Revocable Trust, B. Sherrill Hardcastle and Joe D. Hardcastle Trustees of the B. Sherrill Hardcastle Revocable Trust, Sherrill Hardcastle Custodian under MO–UTMA FBO Faith Morgan Harlin, all of Lebanon, Missouri;* collectively as a group acting in concert, to retain shares of Century Bancshares, Inc., Gainesville, Missouri, and thereby indirectly retain shares of, Century Bank of the Ozarks, Gainesville, Missouri, and Ozarks Heritage Financial Group, Inc., Gainesville, Missouri and thereby retain shares of Legacy Bank & Trust Company, Rogersville, Missouri.

Board of Governors of the Federal Reserve System, March 3, 2017.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2017–04564 Filed 3–7–17; 8:45 am]

BILLING CODE 6210–01–P

GOVERNMENT ACCOUNTABILITY OFFICE

Health Information Technology Advisory Committee Nominations; Request for Letters of Nomination and Resumes

AGENCY: Government Accountability Office (GAO).

ACTION: Request for letters of nomination and resumes.

SUMMARY: The 21st Century Cures Act established the Health Information Technology Advisory Committee to provide recommendations to the National Coordinator for Health Information Technology on policies, standards, implementation specifications, and certification criteria relating to the implementation of a health information technology infrastructure that advances the electronic access, exchange, and use of health information. The Act gave the Comptroller General responsibility for appointing 14 of the committee's members. The Act requires that members at least reflect providers, ancillary health care workers, consumers, purchasers, health plans, health information technology developers, researchers, patients, relevant Federal agencies, and individuals with technical expertise on health care quality, system functions, privacy, security, and on the electronic exchange and use of health information, including the use standards for such activity. GAO is accepting nominations of individuals for this committee. Letters of nomination and resumes should be submitted by April 14, 2017 for appointments that will be made in July 2017. Acknowledgement of submissions will be provided within a week of submission. Please contact Will

Simerl at (404) 679–1888 if you do not receive an acknowledgement.

ADDRESSES:

Email: HITCommittee@gao.gov.

Mail: ATTN: HITC Appointments, U.S. GAO, 441 G Street NW., Washington, DC 20548.

FOR FURTHER INFORMATION CONTACT: GAO Office of Public Affairs, (202) 512–4800.

Authority: Pub. L. 114–255, § 3002(d) (2016).

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2017–04456 Filed 3–7–17; 8:45 am]

BILLING CODE 1610–02–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–17–0263; Docket No. CDC–2017–0021]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a revision request for the information collection titled ''Requirements for the Importation of Nonhuman Primates into the United States.'' This information collection contains the reporting and documentation requirements for registered importers of nonhuman primates.

DATES: Written comments must be received on or before May 8, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0021 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov.*

Please note: All public comment should be submitted through the Federal eRulemaking portal (Regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited 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) ways to enhance the quality, utility, and clarity of the information to be collected; (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; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

Proposed Project

Requirements for the Importation of Nonhuman Primates into the United States (OMB Control No. 0920–0263, Expiration Date, 09/30/2017)—Revision—National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Under the 42 CFR 71.53, CDC collects information pertaining to importers and imported nonhuman primates. This information collection enables CDC to evaluate compliance with pre-arrival of shipment notification requirements and to investigate the number and species of imported nonhuman primates. Also, it enables CDC to determine if adequate measures are being taken for the prevention of exposure to persons and animals during importation.

Since May 1990, CDC has monitored the arrival and/or uncrating of certain shipments of non-human primates imported into the United States. In February 2013, CDC promulgated two regulations pertaining to the importation of nonhuman primates. The first rule, Establishment of User Fees for Filovirus Testing of Nonhuman Primate Liver Samples, outlines a process by which importers can send liver tissues to CDC from primates that die during importation from reasons other than trauma (2/12/2013, Vol.78, No. 29, p. 9828). CDC performs these tests due to the absence of a private sector option. The second rule, Requirements for Importers of Nonhuman Primates, consolidates into 42 CFR 71.53 the requirements previously found in 42 CFR part 71.53 with those found in the Special Permit to Import Cynomolgus, African Green, or Rhesus Monkeys into the United States (2/15/2013, Vol. 78, No. 32/p. 11522). It also rescinded the six-month special-permit requirements for cynomolgus, African green, and rhesus monkeys and extended the time period for registration/permit renewal from 180 days to 2 years, reducing much of the respondent burden. CDC feels these regulatory changes and reporting