

performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB control number.

**DATES:** Written PRA comments should be submitted on or before June 4, 2019. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

**ADDRESSES:** Direct all PRA comments to Cathy Williams, FCC, via email [PRA@fcc.gov](mailto:PRA@fcc.gov) and to [Cathy.Williams@fcc.gov](mailto:Cathy.Williams@fcc.gov).

**FOR FURTHER INFORMATION CONTACT:** For additional information about the

information collection, contact Cathy Williams at (202) 418-2918.

**SUPPLEMENTARY INFORMATION:**

*OMB Control Number:* 3060-1000.

*Title:* Section 87.147, Authorization of Equipment.

*Form Number:* N/A.

*Type of Review:* Extension of a currently approved collection.

*Respondents:* Business or other for-profit entities.

*Number of Respondents:* 25 respondents; 25 responses.

*Estimated Time per Response:* 1 hour.

*Frequency of Response:* One time and occasion reporting requirements and third party disclosure requirement.

*Obligation to Respond:* Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. 154, 303 and 307(e) of the Communications Act of 1934, as amended.

*Total Annual Burden:* 25 hours.

*Total Annual Cost:* N/A.

*Privacy Impact Assessment:* No impact(s).

*Nature and Extent of Confidentiality:* There is no need for confidentiality with this collection of information.

*Needs and Uses:* Section 87.147 is needed to require applicants for aviation equipment certification to submit a Federal Aviation Administration (FAA) determination of the equipment's compatibility with the National Airspace System (NAS). This will

ensure that radio equipment operating in certain frequencies is compatible with the NAS, which shares system components with the military. The notification must describe the equipment, along with a report of measurements, give the manufacturer's identification, antenna characteristics, rated output power, emission type and characteristics, the frequency or frequencies of operation, and essential receiver characteristics if protection is required.

Federal Communications Commission.

**Marlene Dortch,**

*Secretary, Office of the Secretary.*

[FR Doc. 2019-06710 Filed 4-4-19; 8:45 am]

**BILLING CODE 6712-01-P**

**FEDERAL DEPOSIT INSURANCE CORPORATION**

**Notice of Termination of Receiverships**

The Federal Deposit Insurance Corporation (FDIC or Receiver), as Receiver for each of the following insured depository institutions, was charged with the duty of winding up the affairs of the former institutions and liquidating all related assets. The Receiver has fulfilled its obligations and made all dividend distributions required by law.

**NOTICE OF TERMINATION OF RECEIVERSHIPS**

Fund	Receivership name	City	State	Termination date
10142 .....	Madisonville State Bank .....	Madisonville .....	TX	4/1/2019
10198 .....	Century Security Bank .....	Duluth .....	GA	4/1/2019
10214 .....	Innovative Bank .....	Oakland .....	CA	4/1/2019
10424 .....	Charter National Bank & Trust .....	Hoffman Estates .....	IL	4/1/2019
10522 .....	Allied Bank .....	Mulberry .....	AR	4/1/2019

The Receiver has further irrevocably authorized and appointed FDIC-Corporate as its attorney-in-fact to execute and file any and all documents that may be required to be executed by the Receiver which FDIC-Corporate, in its sole discretion, deems necessary, including but not limited to releases, discharges, satisfactions, endorsements, assignments, and deeds. Effective on the termination dates listed above, the Receiverships have been terminated, the Receiver has been discharged, and the Receiverships have ceased to exist as legal entities.

Dated at Washington, DC, on April 2, 2019.

Federal Deposit Insurance Corporation.

**Robert E. Feldman,**

*Executive Secretary.*

[FR Doc. 2019-06712 Filed 4-4-19; 8:45 am]

**BILLING CODE 6714-01-P**

**FEDERAL RESERVE SYSTEM**

**Solicitation of Applications for Membership on the Community Advisory Council**

**AGENCY:** Board of Governors of the Federal Reserve System.

**ACTION:** Notice.

**SUMMARY:** The Board of Governors of the Federal Reserve System (Board) established the Community Advisory

Council (CAC) as an advisory committee to the Board on issues affecting consumers and communities. This Notice advises individuals who wish to serve as CAC members of the opportunity to be considered for the CAC.

**DATES:** Applications received between Monday, April 8, 2019 and Friday, May 31, 2019 will be considered for selection to the CAC for terms beginning January 1, 2020.

**ADDRESSES:** Individuals who are interested in being considered for the CAC may submit an application via the Board's website or via email. The application can be accessed at <https://www.federalreserve.gov/secure/CAC/Application/>. Emailed submissions can

be sent to [CCA-CAC@frb.gov](mailto:CCA-CAC@frb.gov). The information required for consideration is described below.

If electronic submission is not feasible, submissions may be mailed to the Board of Governors of the Federal Reserve System, Attn: Community Advisory Council, Mail Stop I-305, 20th Street and Constitution Ave. NW, Washington, DC 20551.

**FOR FURTHER INFORMATION CONTACT:**

Jennifer Fernandez, Community Development Analyst, Division of Consumer and Community Affairs, Board of Governors of the Federal Reserve System, 20th Street and Constitution Ave. NW, Washington, DC 20551, or (202) 452-2412, or [CCA-CAC@frb.gov](mailto:CCA-CAC@frb.gov). Telecommunications Device for the Deaf (TDD) users may contact (202) 263-4869.

**SUPPLEMENTARY INFORMATION:** The Board created the Community Advisory Council (CAC) as an advisory committee to the Board on issues affecting consumers and communities. The CAC is composed of a diverse group of experts and representatives of consumer and community development organizations and interests, including from such fields as affordable housing, community and economic development, employment and labor, financial services and technology, small business, and asset and wealth building. CAC members meet semiannually with the members of the Board in Washington, DC to provide a range of perspectives on the economic circumstances and financial services needs of consumers and communities, with a particular focus on the concerns of low- and moderate-income consumers and communities. The CAC complements two of the Board's other advisory councils—the Community Depository Institutions Advisory Council (CDIAC) and the Federal Advisory Council (FAC)—whose members represent depository institutions.

The CAC serves as a mechanism to gather feedback and perspectives on a wide range of policy matters and emerging issues of interest to the Board of Governors and aligns with the Federal Reserve's mission and current responsibilities. These responsibilities include, but are not limited to, banking supervision and regulatory compliance (including the enforcement of consumer protection laws), systemic risk oversight and monetary policy decision-making, and, in conjunction with the Office of the Comptroller of the Currency (OCC) and Federal Deposit Insurance Corporation (FDIC), responsibility for implementation of the Community Reinvestment Act (CRA).

This Notice advises individuals of the opportunity to be considered for appointment to the CAC. To assist with the selection of CAC members, the Board will consider the information submitted by the candidate along with other publicly available information that it independently obtains.

**Council Size and Terms**

The CAC consists of at least 15 members. The Board will select members in the fall of 2019 to replace current members whose terms will expire on December 31, 2019. The newly appointed members will serve three-year terms that will begin on January 1, 2020. If a member vacates the CAC before the end of the three-year term, a replacement member will be appointed to fill the unexpired term.

**Application**

Candidates may submit applications by one of three options:

- **Online:** Complete the application form on the Board's website at <https://www.federalreserve.gov/secure/CAC/Application/>.
- **Email:** Submit all required information to [CCA-CAC@frb.gov](mailto:CCA-CAC@frb.gov).
- **Postal Mail:** If electronic submission is not feasible, submissions may be mailed to the Board of Governors of the Federal Reserve System, Attn: Community Advisory Council, Mail Stop I-305, 20th Street and Constitution Ave. NW, Washington, DC 20551.

Interested parties can view the current Privacy Act Statement at: <https://www.federalreserve.gov/aboutthefed/cac-privacy.htm>.

Below are the application fields. Asterisks (\*) indicate required fields.

- Full Name \*
- Email Address \*
- Phone Number \*
- Postal Mail Street Address \*
- Postal Mail City \*
- Postal Zip Code \*
- Organization \*
- Title \*
- Organization Type (select one) \*
  - For Profit
    - Community Development Financial Institution (CDFI)
    - Non-CDFI Financial Institution
    - Financial Services
    - Professional Services
    - Other
  - Non-Profit
    - Advocacy
    - Association
    - Community Development Financial Institution (CDFI)
    - Educational Institution
    - Foundation
    - Service Provider

- Think Tank/Policy Organization
- Other
  - Government
- Primary Area of Expertise (select one) \*
  - Civil rights
  - Community development finance
  - Community reinvestment and stabilization
  - Consumer protection
  - Economic and small business development
  - Labor and workforce development
  - Financial technology
  - Household wealth building and financial stability
  - Housing and mortgage finance
  - Rural issues
  - Other (please specify)
- Secondary Area of Expertise (select one)
  - Civil rights
  - Community development finance
  - Community reinvestment and stabilization
  - Consumer protection
  - Economic and small business development
  - Labor and workforce development
  - Financial technology
  - Household wealth building and financial stability
  - Housing and mortgage finance
  - Rural issues
  - Other (please specify)
- Resume \*
  - The resume should include information about past and present positions you have held, dates of service for each, and a description of responsibilities.
- Cover Letter \*
  - The cover letter should explain why you are interested in serving on the CAC as well as what you believe are your primary qualifications.
- Additional Information
  - At your option, you may also provide additional information about your qualifications.

**Qualifications**

The Board is interested in candidates with knowledge of fields such as affordable housing, community and economic development, employment and labor, financial services and technology, small business, and asset and wealth building, with a particular focus on the concerns of low- and moderate-income consumers and communities. Candidates do not have to be experts on all topics related to consumer financial services or community development, but they should possess some basic knowledge of these areas and related issues. In

appointing members to the CAC, the Board will consider a number of factors, including diversity in terms of subject matter expertise, geographic representation, and the representation of women and minority groups.

CAC members must be willing and able to make the necessary time commitment to participate in organizational conference calls and prepare for and attend meetings two times per year (usually for two days). The meetings will be held at the Board's offices in Washington, DC. The Board will provide a nominal honorarium and will reimburse CAC members only for their actual travel expenses subject to Board policy.

By order of the Board of Governors of the Federal Reserve System, acting through the Director of the Division of Consumer and Community Affairs under delegated authority, March 26, 2019.

**Ann E. Misback,**  
Secretary of the Board.

[FR Doc. 2019-06406 Filed 4-4-19; 8:45 am]

BILLING CODE 6210-01-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2016-D-2730]

#### **Risk Evaluation and Mitigation Strategy: The Food and Drug Administration's Application of Statutory Factors in Determining When a Risk Evaluation and Mitigation Strategy Is Necessary; Guidance for Industry; Availability**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice of availability.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance for industry entitled "Risk Evaluation and Mitigation Strategy: FDA's Application of Statutory Factors in Determining When a Risk Evaluation and Mitigation Strategy Is Necessary." This guidance is intended to clarify how FDA applies the factors set forth in the Federal Food, Drug, and Cosmetic Act (FD&C Act) in determining whether a risk evaluation and mitigation strategy (REMS) is necessary to ensure that the benefits of a drug outweigh its risks. This guidance is one of several developed to fulfill performance goals that FDA agreed to satisfy in the reauthorization of the prescription drug user fee program (the Prescription Drug User Fee Act (PDUFA) V). This

guidance finalizes the draft guidance entitled "FDA's Application of Statutory Factors in Determining When a REMS Is Necessary," issued September 21, 2016.

**DATES:** The announcement of the guidance is published in the **Federal Register** on April 5, 2019.

**ADDRESSES:** You may submit either electronic or written comments on Agency guidances at any time as follows:

#### *Electronic Submissions*

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

#### *Written/Paper Submissions*

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

*Instructions:* All submissions received must include the Docket No. FDA-2016-D-2730 for "REMS: FDA's Application of Statutory Factors in Determining When a REMS Is Necessary." Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at

<https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- **Confidential Submissions**—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

*Docket:* For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of this guidance to the Division of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10001 New Hampshire Ave., Hillandale Building, 4th Floor, Silver Spring, MD 20993-0002; or to the Office of Communication, Outreach and Development, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 3128, Silver Spring, MD 20993-0002. Send one self-addressed adhesive label to