about the Board, its actions, and the economy. The responses to the FR 3076 help the Board determine how to most effectively communicate this information to the public in order to fulfill its statutory responsibilities. The FR 3076 is voluntary. The information collected by the FR 3076 is not considered to be confidential.

Board of Governors of the Federal Reserve System, April 2, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–06948 Filed 4–4–18; 8:45 am]

BILLING CODE 6210–01–P

GOVERNMENT ACCOUNTABILITY OFFICE

Request for Nominations for Board of Governors of the Patient-Centered Outcomes Research Institute (PCORI)

AGENCY: Government Accountability Office (GAO).

ACTION: Request for letters of nomination and resumes.

SUMMARY: The Patient Protection and Affordable Care Act gave the Comptroller General of the United States responsibility for appointing 19 members to the Board of Governors of the Patient-Centered Outcomes Research Institute. In addition, the Directors of the Agency for Healthcare Research and Quality and the National Institutes of Health, or their designees, are members of the Board. As the result of terms ending in September 2018, GAO is accepting nominations in the following categories required in statute: A physician, a nurse, a representative of patients and health care consumers, a representative of private payers, a representative of a state or a federal health program or agency, and a representative of pharmaceutical, device, or diagnostic manufacturers or developers. Nominations should be sent to the email or mailing address listed below. Acknowledgement of submissions will be provided within a week of submission.

DATES: Letters of nomination and resumes should be submitted no later than May 4, 2018, to ensure adequate opportunity for review and consideration of nominees prior to appointment.

ADDRESSES: Submit letters of nomination and resumes by either of the following methods: Email: *PCORI@gao.gov.* Include PCORI Nominations in the subject line of the message, or Mail: U.S. GAO, Attn: PCORI Board Nominations, 441 G Street NW, Washington, DC 20548.

FOR FURTHER INFORMATION CONTACT: Rashmi Agarwal at (202) 512–4077 or *agarwalr@gao.gov* if you do not receive an acknowledgement or need additional information. For general information, contact GAO's Office of Public Affairs, (202) 512–4800.

Authority: [Sec. 6301 and Sec. 10602, Pub. L. 111–148].

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2018–06999 Filed 4–4–18; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2002–D–0093]

Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation; 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 guidance for industry entitled ''Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation.'' This guidance document finalizes the revised draft of the same name that published on October 30, 2015. This guidance provides recommendations to applicants on the chemistry, manufacturing, and controls (CMC); pharmacokinetics and bioavailability; and labeling documentation for liposome drug products submitted in new drug applications (NDAs) and abbreviated new drug applications (ANDAs), reviewed by the Center for Drug Evaluation and Research (CDER).

DATES: The announcement of the guidance is published in the **Federal Register** on April 5, 2018.

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–2002–D–0093 (formerly 2002D–0337) for ''Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation.'' 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