

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]

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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

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. Send one self-addressed adhesive label to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

FOR FURTHER INFORMATION CONTACT:

Richard T. Lostritto, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 4132, Silver Spring, MD 20993–0002, 301–796–1697.

SUPPLEMENTARY INFORMATION:

I. Background

FDA 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 provides recommendations to applicants on the CMC, human

pharmacokinetics and bioavailability, and labeling documentation for liposome drug products submitted in NDAs and ANDAs reviewed by CDER. Although this guidance does not intend to provide recommendations specific to liposome drug products to be marketed under biologics license applications (BLAs), many scientific principles described in this guidance may also apply to these products.

In the **Federal Register** of August 21, 2002 (67 FR 54220), FDA announced the availability of a draft version of this guidance. FDA published a revised draft guidance on October 30, 2015 (80 FR 66906), because of the need to address changes in technology since the draft was first published in 2002, and to add ANDAs to the scope. Most of the changes to the 2015 revised draft guidance were made to clarify statements in the 2002 draft guidance. FDA received comments in response to the draft and revised draft guidance, and this guidance reflects FDA’s careful consideration of those comments.

The guidance does not provide recommendations on clinical efficacy and safety studies, nonclinical pharmacology and/or toxicology studies, liposome formulations of vaccine adjuvants or biologics, or drug-lipid complexes.

This guidance is being issued consistent with FDA’s good guidance practices regulation (21 CFR 10.115). The guidance represents the current thinking of FDA on “Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation.” It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations. This guidance is not subject to Executive Order 12866.

II. The Paperwork Reduction Act of 1995

This guidance refers to previously approved collections of information that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR part 314 have been approved under OMB control number 0910–0001.

III. Electronic Access

Persons with access to the internet may obtain the document at either <https://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm> or <https://www.regulations.gov>.

Dated: March 29, 2018.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2018–06926 Filed 4–4–18; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0438]

Agency Information Collection Request. 60-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before June 4, 2018.

ADDRESSES: Submit your comments to Sherrette.Funn@hhs.gov or by calling (202) 795–7714.

FOR FURTHER INFORMATION CONTACT:

When submitting comments or requesting information, please include the document identifier 0990–0438–60D and project title for reference, to Sherrette.funn@hhs.gov, or call the Reports Clearance Officer.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection: Teen Pregnancy Prevention Performance Measures Data Collection, Office of Adolescent Health.

Type of Collection: Revision.
OMB No.: 0990–0438.

Abstract: The Office of Adolescent Health (OAH), U.S. Department of Health and Human Services (HHS) is requesting approval by OMB on a revision of the Teen Pregnancy Prevention (TPP) Performance Measures from funded grantees. The performance measures data include grantee-level measures (dissemination, partners,