

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Sunshine Act; Notice of Meeting July 24, 2018, Telephonic, 12:00 p.m., 10th Floor Board Meeting Room, 77 K Street NE, Washington, DC 20002

Agenda

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD MEMBER MEETING STATUS: All parts will be open to the public.

MATTERS TO BE CONSIDERED:

Open Session

1. Approval of the Minutes for the June 25, 2018 Board Member Meeting
2. Monthly Reports
 - (a) Participant Activity
 - (b) Legislative Report
3. Quarterly Reports
 - (c) Investment Performance
 - (d) Budget Review
 - (e) Audit Status
4. Enterprise Risk Management Update
5. IT Update

CONTACT PERSON FOR MORE INFORMATION: Kimberly Weaver, Director, Office of External Affairs, (202) 942-1640.

Dated: July 16, 2018.
Dharmesh Vashee,
Deputy General Counsel, Federal Retirement Thrift Investment Board.
 [FR Doc. 2018-15433 Filed 7-16-18; 4:15 pm]
BILLING CODE 6760-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB NO.: 0970-0383]

Submission for OMB Review; Comment Request; Evaluation of the Transitional Living Program (TLP)—Extension

Description:
 The Family and Youth Services Bureau (FYSB) and the Office of Planning, Research, Evaluation (OPRE) in the Administration for Children and Families (ACF) are requesting to continue collecting data as part of a currently approved information collection (OMB No. 0970-0383). The

purpose is to continue baseline data collection at study enrollment and follow-up data collection for the Evaluation of the Transitional Living Program (TLP). The TLP evaluation was designed to examine the effects of FYSB's Transitional Living Program on runaway and homeless youth, focusing on such outcomes as housing and homelessness, education or training, employment, social connections, socio-emotional well-being, and risk behaviors.

Data collection will include three primary surveys: (1) A survey administered at the time of TLP enrollment (baseline), (2) a survey administered 6 months after enrollment, which will collect information on short-term outcomes; and (3) a survey administered at 12 months, which will collect information on longer-term outcomes. Participants will be enrolled through the TLP study sites.

Respondents: Runaway and homeless youth ages 16 to 22 who agree to participate in the study upon enrollment into one of the TLP study sites.

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
Young Adult Baseline Survey	600	200	1	0.62	124
Young Adult 6-Month Follow Up Survey	600	200	1	0.61	122
Young Adult 12-Month Follow Up Survey	600	200	1	0.61	122
Estimated Total Burden Hours					368

Additional Information: Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. Email address: OPREinfocollection@acf.hhs.gov.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget
 Paperwork Reduction Project
 Email: OIRA_SUBMISSION@OMB.EOP.GOV

Attn: Desk Officer for the Administration for Children and Families

Mary B. Jones,
ACF/OPRE Certifying Officer.
 [FR Doc. 2018-15307 Filed 7-17-18; 8:45 am]
BILLING CODE 4184-33-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0742]

Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection for drug establishment registration and product listing.

DATES: Submit either electronic or written comments on the collection of information by September 17, 2018.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before September 17,

2018. The <https://www.regulations.gov> electronic filing system will accept comments until midnight Eastern Time at the end of September 17, 2018. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

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-2011-N-0742 for "Agency Information Collection Activities; Proposed Collection; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution." Received comments, those filed in a timely manner (see **ADDRESSES**), 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.

FOR FURTHER INFORMATION CONTACT:

Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-5733, PRASStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: Under the 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. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests

or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution—21 CFR Part 207

OMB Control Number 0910-0045—Extension

This information collection supports FDA's drug establishment registration and listing regulations and associated guidance intended to assist respondents in this regard. Requirements for drug establishment registration and drug listing are set forth in section 510 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 360), and section 351 of the Public Health Service Act (42 U.S.C. 262). Section 224 of the Food and Drug Administration Amendments Act of 2007 (Pub. L. 110-85) amended section 510(p) of the FD&C Act to require electronic drug establishment registration and drug listing. Regulations implementing these provisions are established under 21 CFR part 207. Except as provided in § 207.65, all information submitted must be transmitted to FDA in electronic format by using our electronic drug registration and listing system, in a form that we can process, review, and archive. Establishment registration information helps FDA identify who is manufacturing, repacking, relabeling,

and salvaging drugs and where those operations are performed. Drug listing information gives FDA a current inventory of drugs manufactured, repacked, relabeled, or salvaged for commercial distribution. Both types of information facilitate implementation and enforcement of the FD&C Act and are used for many important public health purposes.

Registration Under Part 207

Unless otherwise exempt under section 510(g) of the FD&C Act or § 207.13, all manufacturers, repackers, relabelers, and salvagers must register each domestic establishment that manufactures, repacks, relabels, or salvages a drug, or an animal feed bearing or containing a new animal drug, and each foreign establishment that manufactures, repacks, relabels, or salvages a drug, or an animal feed bearing or containing a new animal drug, that is imported or offered for import into the United States. When operations are conducted at more than one establishment and common ownership and control among all the establishments exists, the parent, subsidiary, or affiliate company may submit registration information for all establishments.

Private label distributors who do not also manufacture, repack, relabel, or salvage drugs are not required to register under part 207. FDA will accept registration or listing information submitted by a private label distributor only if it is acting as an authorized agent for and submitting information that pertains to an establishment that manufactures, repacks, relabels, or salvages drugs.

Under § 207.21, domestic manufacturers, domestic repackers, domestic relabelers, and domestic drug product salvagers must complete initial registration of each establishment no later than 5 calendar days after beginning to manufacture, repack, relabel, or salvage a drug. In addition, foreign manufacturers, foreign repackers, foreign relabelers, and foreign drug product salvagers must register each establishment before the drug is imported or offered for import into the United States.

The information that must be provided to FDA for registration is described in § 207.25 and includes the following: (1) Name of the owner or operator of each establishment; if a partnership, the name of each partner; if a corporation, the name of each corporate officer and director, and the place of incorporation; (2) each

establishment's name, physical address, and telephone number(s); (3) all name(s) of the establishment, including names under which the establishment conducts business or names by which the establishment is known; (4) registration number of each establishment, if previously assigned by FDA; (5) a Unique Facility Identifier in accordance with the system specified under section 510 of the FD&C Act; (6) all types of operations performed at each establishment; (7) name, mailing address, telephone number, and email address of the official contact for the establishment, as provided in § 207.69(a); and (8) additionally, with respect to foreign establishments subject to registration, the name, mailing address, telephone number, and email address must be provided for: (a) The U.S. agent, as provided in § 207.69(b); (b) each importer in the United States of drugs manufactured, repacked, relabeled, or salvaged at the establishment that is known to the establishment; and (c) each person who imports or offers for import such drug to the United States.

Registrants must update their registration information as prescribed under § 207.29.

National Drug Code (NDC)

The NDC for a drug is a numeric code. Each finished drug product or unfinished drug subject to the listing requirements of part 207 must have a unique NDC to identify its labeler, product, and package size and type. The format of an NDC is described under § 207.33.

Under § 207.35, registrants must notify us of a change in any of the drug characteristics (except certain identifying information) for an NDC in § 207.33, and assign a new product code and package code for that drug.

Listing Under Part 207

Under § 207.41, registrants must list each drug that it manufactures, repacks, relabels, or salvages for commercial distribution. Each domestic registrant must list each such drug regardless of whether the drug enters interstate commerce. When operations are conducted at more than one establishment, and common ownership and control exists among all the establishments, the parent, subsidiary, or affiliate company may submit listing information for any drug manufactured, repacked, relabeled, or salvaged at any such establishment. A drug manufactured, repacked, or relabeled for

private label distribution must be listed in accordance with the requirements.

Registrants must provide listing information for each drug in accordance with the listing requirements described in §§ 207.49, 207.53, and 207.54 that correspond to the activity or activities they engage in for that drug. For both animal and human drugs, each registrant must list each drug it manufactures, repacks, or relabels for commercial distribution under the trade name or label of a private label distributor using an NDC that includes such private label distributor's labeler code.

Additionally, in the case of human drugs, each registrant must list each human drug it manufactures, repacks, or relabels using an NDC that includes the registrant's own labeler code, regardless of whether the drug is commercially distributed under the registrant's own label or trade name or under the label or trade name of a private label distributor.

Under § 207.45, for each drug being manufactured, repacked, relabeled, or salvaged for commercial distribution at an establishment at the time of initial registration, drug listing information must be submitted no later than 3 calendar days after the initial registration of the establishment.

Each registrant must provide the listing information described under § 207.49 for each drug it manufactures for commercial distribution. Each registrant must also provide the listing information for each drug it repacks or relabels under § 207.53. A registrant who also relabels or repacks a drug that it salvages must list the drug it relabels or repacks in accordance with § 207.53. Registrants who perform only salvaging with respect to a drug must provide the listing information for that drug as required under § 207.54. Additional information may be requested for a listed drug as described in § 207.55.

Under § 207.57, registrants must update drug listing information submitted previously (either when the change is made or, at a minimum, each June and December). Registrants must also notify FDA if any listed drug has been discontinued from marketing or if any discontinued drug has been reintroduced and provide listing information for any drug not yet listed (at the time of annual establishment registration if not sooner).

We estimate the burden of the information collection as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

Activity; 21 CFR section	Number of respondents	Number of responses per respondent	Total annual responses	Hours per response	Total hours
Initial establishment registration; §§ 207.17, 207.21, 207.25.	1,480	2	2,960	1	2,960
Annual review and update of registration information (including expedited updates); § 207.29.	10,000	1	10,000	.5 (30 minutes)	5,000
Initial listing (including NDC); §§ 207.33, 207.41, 207.45, 207.49, 207.53, 207.54, 207.55.	1,713	7.28	12,470	1.5	18,705
June and December review and update (or certification) of listing; §§ 207.35, 207.57.	5,300	20	106,000	.75 (45 minutes) ...	79,500
Waiver requests; § 207.65	1	1	1	.5 (30 minutes)	1
Public disclosure exemption requests; § 207.81(c)	100	1	100	1	100
Total					106,266

¹ There are no capital or operating and maintenance costs associated with the information collection.

TABLE 2—ESTIMATED ANNUAL RECORDKEEPING BURDEN ¹

Standard operating procedure (SOP) for creating and uploading the SPL file	Number of recordkeepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeping	Total hours
Preparation of SOP	1,000	1	1,000	40	40,000

¹ There are no capital or operating and maintenance costs associated with the information collection.

Based on FDA data, we estimate that 1,480 respondents will submit 2,960 new establishment registrations annually. Based on the number of registered establishments in our database, we estimate 10,000 registrants will provide 10,000 annual reviews and updates of registration information (including expedited updates) or reviews and certifications that no changes have occurred. The estimates include the registration of establishments for both domestic and foreign manufacturers, repackers, relabelers, and drug product salvagers, and registration information submitted by anyone acting as an authorized agent for an establishment that manufactures, repacks, relabels, or salvages drugs. The estimates include an additional 80 positron emission tomography (PET) drug producers who are not exempt from registration and approximately 30 manufacturers of plasma derivatives.

We estimate that it will take 1 hour for registrants to submit initial registration information electronically for each new establishment. We also estimate that it will take approximately 30 minutes for each annual review and update of registration information (including any expedited updates) or each review and certification that no changes have occurred. The burden hour estimates above are based on our familiarity with the amount of time it takes registrants to input registration information electronically since June 2009. The estimates are an average of the time it would take to register a

domestic or foreign establishment and an average of the time it would take to review registration information and update several registration items in the database or review registration information and only certify that no changes have occurred.

Based on the number of drugs listed annually since June 2009, we estimate that approximately 1,713 registrants will report 12,469 new listings annually (including the information submitted to obtain a labeler code and to reserve an NDC for future use).

Based on the number of drugs in our listing database and the current number of changes to listing information submitted, we estimate 5,300 registrants will each report 20 reviews and updates (including the information submitted to revise an NDC) for a total of 106,000 annually.

The estimates for the number of drug listings include both domestic and foreign listings, listings submitted by registrants for products sold under their own names as well as products intended for private label distribution, and information submitted related to an NDC and to obtain a labeler code. The estimate for the number of drugs subject to the listing requirements includes PET drugs and approximately 30 plasma derivatives. The estimates for the number of June and December reviews and updates of listing information include the number of changes to drug characteristics pertaining to the drug product code to obtain a new NDC and the reports of the withdrawal of an

approved drug from sale under 21 CFR 314.81(b)(3)(iii).

Based on our familiarity with the time required to input listing information electronically since June 2009, we estimate that it will take registrants 1 hour and 30 minutes to submit information electronically for each drug they list for the first time (for both foreign and domestic registrant listings). These estimates are an average of the time it will take manufacturers, repackers, relabelers, and drug product salvagers, with drug product salvagers taking considerably less time than manufacturers. The estimates include the time for submitting the content of labeling and other labeling in electronic format. (For drugs subject to an approved marketing application, the electronic submission of the content of labeling under § 314.50(l)(1)(i) is approved under OMB control number 0910–0001). We also estimate that it will take 45 minutes for each June and December review and update. These estimates represent the average amount of time to review and update listing information or to review and certify that no changes have occurred. The estimates include the time for submitting any labeling for each drug, changes to the drug's characteristics submitted for a new NDC, and reports of the withdrawal of an approved drug from sale under § 314.81(b)(3)(iii).

In 2009, to help respondents transition to the current electronic reporting requirements, FDA issued the guidance for industry entitled

“Providing Regulatory Submissions in Electronic Format—Drug Establishment Registration and Drug Listing.” The document provides guidance to industry on the statutory requirement to submit electronically drug establishment registration and drug listing information. The guidance describes the types of information to include for purposes of drug establishment registration and drug listing and how to prepare and submit the information in an electronic format (Structured Product Labeling (SPL) files) that FDA can process, review, and archive. The burden attributed to the guidance includes the preparation of an SOP for creating and uploading the SPL file. Although most firms will already have prepared an SOP for the electronic submission of drug establishment registration and drug listing information, each year additional firms will need to create an SOP. As reflected in table 2, FDA estimates that approximately 1,000 firms will expend 40 hours to prepare, review, and approve an SOP, for a total of 40,000 hours annually.

Dated: July 12, 2018.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2018–15298 Filed 7–17–18; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–D–2281]

Innovative Approaches for Nonprescription Drug Products; Draft 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 draft guidance for industry entitled “Innovative Approaches for Nonprescription Drug Products.” This draft guidance describes two innovative approaches that may be useful to consider for demonstrating safety and effectiveness for a nonprescription drug product in cases where the drug facts labeling (DFL) alone is not sufficient to ensure that the drug product can be used safely and effectively in a nonprescription setting: The development of labeling in addition to the DFL and the implementation of additional conditions so that consumers

appropriately self-select and use the product.

DATES: Submit either electronic or written comments on the draft guidance by September 17, 2018 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance 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–2018–D–2281 for “Innovative Approaches for Nonprescription Drug Products; Draft Guidance for Industry; Availability.” 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 the draft 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 draft guidance document.

FOR FURTHER INFORMATION CONTACT: Chris Wheeler, Center for Drug Evaluation and Research, 10903 New