

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: Terrie Crescenzi, Office of Pediatric Therapeutics, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, terrie.crescenzi@fda.hhs.gov or Elizabeth Sanford, Office of Pediatric Therapeutics, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, elizabeth.sanford@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

On July 9, 2012, the President signed into law the Food and Drug Administration Safety and Innovation Act (FDASIA) (Pub L. 112–144). Section 508 of FDASIA directs the Secretary of HHS to submit a report to Congress on the implementation of the Best Pharmaceuticals for Children Act (BPCA) and Pediatric Research Equity Act (PREA). The first report was required to be submitted to Congress by July 9, 2016, and subsequent reports are required every 5 years thereafter. FDASIA also requires FDA to obtain, at least 180 days prior to submission of the report, stakeholder input from patient groups, consumer groups, regulated

industry, academia, and any other interested parties to obtain any recommendations or information relevant to the report including suggestions for modifications that would improve pediatric drug research and pediatric labeling of drugs and biological products. In addition, on August 18, 2017, the Food and Drug Administration Reauthorization Act (FDARA) (Pub L. 115–52) was signed into law, which outlined additional requirements to be included in the report.

Some of the issues to be discussed at the meeting will include, but not be limited to:

- Hearing from patients/parents and patient/parent groups, industry, academia and other stakeholders about the public health impact that pediatric legislation may have had on them or their communities;
- Understanding the effects of the requirement of pediatric studies under PREA or the incentives under BPCA on drug/biologic development plans; and
- Understanding if there are any barriers or resource issues preventing undertaking or completing studies under PREA and BPCA.

II. Meeting Attendance and Participation

If you wish to attend this meeting, visit <https://www.eventbrite.com/e/stakeholder-input-on-pediatric-legislation-registration-74306461627>. Please register by November 15, 2019. Those who are unable to attend the meeting in person can register to view a live webcast of the meeting. You will be asked to indicate in your registration if you plan to attend in person or via the webcast. Your registration will also contain your complete contact information, including name, title, affiliation, address, email address, and phone number. Seating will be limited so early registration is recommended. Registration is free and will be on a first-come, first-served basis. Onsite registration on the day of the meeting will be based on space availability. To view the webcast, visit: <https://collaboration.fda.gov/pediatric-legislation/>. If you need special accommodations due to a disability, please contact Elizabeth Sanford (see **FOR FURTHER INFORMATION CONTACT**) at least 7 days in advance. Persons attending the meeting are advised that FDA is not responsible for providing access to electrical outlets.

Persons interested in presenting comments at the meeting will be asked to indicate this in their registration. If you intend to use a PowerPoint presentation, please email the

presentation to opt@fda.hhs.gov by November 15, 2019. FDA will try to accommodate all participant requests to speak, however the duration of comments may be limited by time constraints.

Comments: Regardless of attendance at the public meeting, you can submit electronic or written comments to the public docket (see **ADDRESSES**) by December 19, 2019. Received comments may be seen in the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: October 21, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–23264 Filed 10–24–19; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–D–0283]

Postmarketing Studies and Clinical Trials—Implementation of Section 505(o)(3) of the Federal Food, Drug, and Cosmetic Act; 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 “Postmarketing Studies and Clinical Trials—Implementation of Section 505(o)(3) of the Federal Food, Drug, and Cosmetic Act.” This draft guidance revises the guidance for industry of the same name issued April 1, 2011. The draft guidance is being revised to describe the multiple factors that FDA considers, before requiring a postmarketing study or clinical trial for the purposes described in the Federal Food, Drug, and Cosmetic Act (FD&C Act), when determining the sufficiency of the reports under the FD&C Act and the active postmarket risk identification and analysis (ARIA) system available under the FD&C Act to meet these purposes. The draft guidance is also being revised to reflect certain provisions enacted under the Substance Use-Disorder Prevention that Promotes Opioid Recovery and Treatment (SUPPORT) for Patients and Communities Act as they relate to postmarketing studies and clinical trials.

DATES: Submit either electronic or written comments on the draft guidance by December 24, 2019 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-2009-D-0283 for "Postmarketing Studies and Clinical Trials—Implementation of Section 505(o)(3) of the Federal Food, Drug, and Cosmetic Act; 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 or 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 assist that office in processing your requests. See the **SUPPLEMENTARY**

INFORMATION section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT:

Ayanna Augustus, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 6426, Silver Spring, MD 20993-0002, 301-796-3980; or Stephen Ripley, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 7301, Silver Spring, MD 20993-0002, 240-402-7911.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry entitled, "Postmarketing Studies and Clinical Trials—Implementation of Section 505(o)(3) of the Federal Food, Drug, and Cosmetic Act."

Section 505(o)(3) of the FD&C Act ¹ (21 U.S.C. 355(o)) authorizes FDA to require certain postmarketing studies or clinical trials for prescription drugs to obtain more information about a serious risk that may be associated with a drug. In some cases, FDA may be concerned about a serious risk that is potentially or known to be associated with a drug but may not know enough about the risk to determine if or how to address it, such as by describing the risk in labeling. Section 505(o)(3)(B) of the FD&C Act states that postmarketing studies and clinical trials may be required for any or all of the following purposes: (1) To assess a known serious risk related to the use of the drug; (2) to assess signals of serious risk related to the use of the drug; or (3) to identify an unexpected serious risk when available data indicates the potential for a serious risk.

Prior to requiring a postmarketing study or clinical trial, FDA must find that the reports under section 505(k)(1) of the FD&C Act and the ARIA system ² made available under section 505(k)(3) of the FD&C Act will not be sufficient to meet the purposes described in section 505(o)(3)(B) of the FD&C Act. ³ Similarly, before requiring a postmarketing clinical trial, FDA must find that a postmarketing study will not be sufficient to meet the purposes

¹ Section 901 of Title IX of the Food and Drug Administration Amendments Act of 2007 (Pub. L. 110-85) amended the FD&C Act by adding section 505(o).

² Section 505(k)(3) of the FD&C Act mandates that FDA establish an active surveillance system for monitoring drugs, using electronic data from healthcare information holders. The Sentinel System draws on existing healthcare data from multiple sources to actively monitor the safety of medical products. The ARIA system is a subcomponent of the Sentinel System.

³ Section 505(o)(3)(D)(i) of the FD&C Act.

described in section 505(o)(3)(B) of the FD&C Act.⁴

In April 2011, FDA issued a guidance describing how it would implement section 505(o) of the FD&C Act. At that time, the ARIA system was still in early development. The ARIA system is now officially launched, and FDA must consider the system's sufficiency to meet the purposes of section 505(o)(3)(B) of the FD&C Act to determine if a postmarketing study or clinical trial is necessary. This draft guidance revises the guidance for industry of the same name issued on April 1, 2011 (76 FR 18226). Significant changes from the 2011 version include explaining how FDA considers the reporting under section 505(k)(1) of the FD&C Act and the ARIA system when determining their sufficiency for the purposes under section 505(o)(3)(B) of the FD&C Act. The guidance is also being revised to provide examples of postmarketing requirements under section 505(o)(3) of the FD&C Act to assess a potential reduction in the expected effectiveness of a drug under certain circumstances. FDA's authority to require these types of studies or trials was clarified by a modification to the definition of *adverse drug experience* at section 505–1(b)(1)(E) of the FD&C Act (21 U.S.C. 505–1(b)(1)(E)) enacted under section 3041 of the SUPPORT Act (Pub. L. 115–271).

This draft guidance is being issued consistent with FDA's good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the current thinking on implementation of section 505(o)(3)(B) of the FD&C Act. 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.

II. Paperwork Reduction Act of 1995

This draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information 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 following collections of information for postmarketing reports and clinical data in 21 CFR 314.50, 314.80, 314.81, 314.98, 314.430, and 314.610(b), subpart I have been approved under OMB control number 0910–0001: (1) Preparing and submitting reports pertaining to safety, postmarketing commitments and

preparing and submitting spontaneous and periodic reports, including active postmarket risk identification (using electronic health care data) and any milestones or submissions for which projected dates were specified as part of the postmarketing commitment; (2) submitting a proposed timetable of the postmarketing commitments; (3) preparing registries and submitting them when appropriate; (4) designing meta-analyses to evaluate statistical analyses of data; (5) preparing assay procedures; and (6) prepare a plan or approach for approval an NDA when human efficacy studies are not ethical or feasible.

The following collections of information for postmarketing studies and clinical trials (including various patient populations) in 21 CFR 312.23 have been approved under OMB control number 0910–0014: (1) Conducting in vitro laboratory tests and studies to compare pregnancy incidence and pregnancy outcomes and/or child outcomes for patients exposed to a drug; (2) submitting an introductory statement and general investigational plan, including a drug's pharmacological class; and (3) submitting protocols for drug safety and pharmacology and toxicology information.

The collections of information in 21 CFR 310.305, 314.80, and 314.98 for submitting adverse event information to the FDA Adverse Event Reporting System have been approved under OMB control numbers 0910–0230 and 0910–0291; the collections of information in 21 CFR 312.47 and 312.82 for submitting a meeting request to appeal the conduct of a postmarketing study or clinical trial have been approved under OMB control number 0910–0430 (and guidance for industry and review staff entitled "Formal Dispute Resolutions: Appeals Above the Division Level" (available at <https://www.fda.gov/ucm/groups/fdagov-public/@fdagov-drugs-gen/documents/document/ucm343101.pdf>).

The following collection of information in § 314.510 has been approved under OMB control number 0910–0765: Requests for serious or life-threatening diseases or conditions that may be granted accelerated approval if FDA determines the product has an effect on a surrogate endpoint that is reasonably likely to predict clinical benefit or on a clinical endpoint that can be measured earlier than irreversible morbidity or mortality or other clinical benefit.

III. Electronic Access

Persons with access to the internet may obtain the draft guidance at either

<https://www.fda.gov/drugs/guidance-compliance-regulatory-information/guidances-drugs>, <https://www.fda.gov/vaccines-blood-biologics/guidance-compliance-regulatory-information-biologics/biologics-guidances>, or <https://www.regulations.gov>.

Dated: October 21, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–23312 Filed 10–24–19; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Evidence-based Screening in Diverse Adult Populations.

Date: November 15, 2019.

Time: 12:00 p.m. to 1:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Karen Nieves Lugo, MPH, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health Bethesda, MD 20892, 301–594–9088, karen.nieveslugo@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Cancer Diagnostics and Treatments (CDT).

Date: November 18–19, 2019.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW, Washington, DC 20015.

Contact Person: Zhang-Zhi Hu, MD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6186,

⁴ Section 505(o)(3)(D)(ii) of the FD&C Act.