

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 (CBER), 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 guidance document.

FOR FURTHER INFORMATION CONTACT: Dorothy West, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6332, Silver Spring, MD 20993–0002, 301–796–0164; 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 guidance for industry entitled “Providing Regulatory Submissions in Electronic Format—Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications (Revision 7).” This guidance provides information regarding submission types that warrant an exemption or long-term waiver from Agency eCTD requirements. In addition, this guidance outlines certain circumstances where FDA will consider granting short-term waivers from eCTD submission requirements. This guidance is intended to address current concerns raised with FDA regarding the burden of complying with eCTD submission requirements, which could have unintended public health consequences.

This guidance finalizes the draft guidance entitled “Providing Regulatory Submissions in Electronic Format—Certain Human Pharmaceutical Product Applications and Related Submissions Using the Electronic Common Technical

Document Specifications (Revision 7)” issued on July 16, 2019 (84 FR 33949). The Agency received comments on the draft guidance requesting that FDA clarify certain requirements relating to Type III DMF submissions. The Agency considered these comments and made technical and editorial changes for clarity, where appropriate. For example, “noncommercial products” was changed to “noncommercial INDs” to clarify the type of submissions referenced in that section of the guidance. In addition, the FDA Electronic Submissions Gateway (ESG) waiver request submission option has been removed to avoid confusion when selecting from the ESG drop-down options.

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 “Providing Regulatory Submissions in Electronic Format—Certain Human Pharmaceutical Product Applications and Related Submissions Using the eCTD Specifications (Revision 7).”

FDA guidances ordinarily contain standard language explaining that guidances should be viewed only as recommendations unless specific regulatory or statutory requirements are cited. FDA is not including this standard language in this guidance because this guidance contains binding provisions. In section 745A(a) of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 379k–1(a)), Congress granted explicit authorization to FDA to specify in guidance the format for the electronic submissions required under that section and required that FDA “shall” issue such guidance. Accordingly, this guidance explains such requirements under section 745A(a) of the FD&C Act, indicated by the use of the words *must* or *required*, and therefore is not subject to the usual restrictions in FDA’s good guidance practice regulations, such as the requirement that guidances not establish legally enforceable responsibilities. See *e.g.*, 21 CFR 10.115(d).

II. Paperwork Reduction Act of 1995

This 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. 3501–3521). The collections of information in 21 CFR part 312 have been approved under OMB control number 0910–0014; the collections of information in 21 CFR part 314 have been approved under

OMB control number 0910–0001; and the collections of information in 21 CFR part 601 have been approved under OMB control number 0910–0338.

III. Electronic Access

Persons with access to the internet may obtain the guidance at <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: February 18, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020–03522 Filed 2–21–20; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–D–1216]

Electronic Common Technical Document v4.0 Technical Conformance Guide; Food and Drug Administration Electronic Common Technical Document v4.0 Module 1 Implementation Package; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is requesting comment on the draft Electronic Common Technical Document (eCTD) v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package. The eCTD v4.0 Technical Conformance Guide will provide specifications, recommendations, and general considerations on how to submit eCTD v4.0-based electronic submissions to the Center for Drug Evaluation and Research (CDER) or the Center for Biologics Evaluation and Research (CBER) using the International Council for Harmonisation eCTD v4.0 Implementation Package and the FDA eCTD v4.0 Module 1 Implementation Package. The Agency is seeking comment on the eCTD v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package for the accuracy, suitability, and appropriateness of these specifications for the submission of eCTD v4.0 submissions. These versions of the documents are not for implementation.

DATES: Submit either electronic or written comments by April 9, 2020.

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 April 9, 2020. The <https://www.regulations.gov> electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of April 9, 2020. 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 "Instruction."

Instructions: All submissions received must include the Docket No. FDA-2018-D-1216 for "Electronic Common Technical Document (eCTD) v4.0

Technical Conformance Guide; FDA eCTD v4.0 Module 1 Implementation Package." 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.govinfo.gov/content/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. The eCTD v4.0 Technical Conformance Guide and FDA eCTD v4.0 Module 1 Implementation Package are available on FDA's eCTD v4.0 web page at: <https://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionRequirements/ElectronicSubmissions/ucm309911.htm>.

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 eCTD v4.0 Technical Conformance Guide or FDA eCTD v4.0 Module 1 Implementation Package 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 assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to these documents.

FOR FURTHER INFORMATION CONTACT:

Jonathan Resnick, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 3160, Silver Spring, MD 20993-0002, 301-796-7997; or Stephen Ripley, Center for Biologics Evaluation and Research, Food and Drug Administration, Bldg. 71, Rm. 7301, Silver Spring, MD 20993-0002, 240-402-7911.

SUPPLEMENTARY INFORMATION:

I. Background

The eCTD v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package are draft versions of the eCTD standard format. FDA will continue to only accept eCTD v3.2.2 submissions until eCTD version 4.0 is finalized. Once eCTD v4.0 is finalized, FDA will accept both eCTD v3.2.2 and eCTD v4.0 submissions for a lengthy phase-in period before eventually only accepting eCTD v4.0 submissions. FDA is requesting comments on the draft eCTD v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package for eCTD v4.0 submissions only. After receiving comments, the Agency will update the eCTD v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package to facilitate the Agency's future acceptance of eCTD v4.0 submissions.

The eCTD v4.0 Technical Conformance Guide will provide specifications, recommendations, and general considerations on how to submit eCTD v4.0-based electronic submissions to CDER or CBER when the Agency implements eCTD v4.0. The eCTD v4.0 Technical Conformance Guide is organized as follows:

Section 1: Introduction

- Provides information on regulatory policy and guidance background, purpose, document control, new features of eCTD v4.0, and guidelines for an eCTD v4.0 submission.

Section 2: Submission Contents

- Recommends and provides details on specific topics organized by their placement (by module) in the eCTD submission.

Section 3: Combination Products

- Recommends and provides details on device combination product information organized by their placement in the eCTD submission.

Section 4: Two-Way Communications

- Provides details on the two-way communication process.

Section 5: Rules for Submission Tracking Information

- Provides details on the submission tracking relationships for an FDA eCTD submission.

The FDA eCTD v4.0 Module 1 Implementation Package will provide the detailed specifications to create Module 1 of an eCTD v4.0-based electronic submission for CDER or CBER. The Implementation Package will provide the technical specifications and the necessary components to create a valid FDA eCTD v4.0 submission. The Implementation Package contains the following components:

FDA eCTD Module 1 Implementation Guide

- The technical specification for the FDA eCTD v4.0 Module 1 using the Health Level Seven Regulated Product Submission Release 2, Normative standard.

FDA Regional Generic Code Controlled Vocabulary Files

- Includes region-specific vocabulary and the files intended for implementers to use as a computable version of the controlled vocabulary content.

FDA Regional Module XML Samples

- Includes samples of M1 eCTD v4.0 xml.

FDA Object Identifiers (OID) Listing

- Provides the OIDs to be used for the FDA Module 1 controlled vocabulary.

FDA Regional Controlled Vocabulary

- Includes region-specific vocabulary and these files are intended as the human readable version of the controlled vocabulary content.

FDA Regional Controlled Vocabulary for Transition Mapping Message DTD 2.01

- Provides a human readable version of the controlled vocabulary transition mapping for the transition from Module 1 DTD 2.01.

FDA Regional Controlled Vocabulary for Transition Mapping Message DTD 3.3

- Provides a human readable version of the controlled vocabulary transition mapping for the transition from Module 1 DTD 3.3.

II. Electronic Access

Persons with access to the internet may obtain the eCTD v4.0 Technical Conformance Guide and the FDA eCTD v4.0 Module 1 Implementation Package at either <https://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionsRequirements/ElectronicSubmissions/ucm309911.htm> or <https://www.regulations.gov>.

Dated: February 19, 2020.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2020-03574 Filed 2-21-20; 8:45 am]

BILLING CODE 4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES**Food and Drug Administration**

[Docket No. FDA-2020-N-0008]

Allergenic Products Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Allergenic Products Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. The meeting will be open to the public.

DATES: The meeting will be held on May 15, 2020, from 8:30 a.m. to 3 p.m.

ADDRESSES: The meeting will be held at FDA White Oak Campus, 10903 New Hampshire Ave., Bldg. 31, Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993-0002. Entrance for the public meeting participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. For security information, please refer to <https://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/>

[WhiteOakCampusInformation/ucm241740.htm](https://www.fda.gov/WhiteOakCampusInformation/ucm241740.htm). For those unable to attend in person, the meeting will also be webcast and will be available at the following link: <https://collaboration.fda.gov/apac051520/>. Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: <https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm>.

FOR FURTHER INFORMATION CONTACT:

Kathleen Hayes or Monique Hill, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6307C, Silver Spring, MD 20993-0002, 301-796-7864, Kathleen.Hayes@fda.hhs.gov, or 301-796-4620, monique.hill@fda.hhs.gov, respectively; or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at <https://www.fda.gov/AdvisoryCommittees/default.htm> and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

SUPPLEMENTARY INFORMATION:

Agenda: On May 15, 2020, the Center for Biologics Evaluation and Research's (CBER) Allergenic Products Advisory Committee (APAC) will meet in open session to discuss and make recommendations on the safety and efficacy of Peanut (*Arachis hypogaea*) Allergen Extract manufactured by DBV Technologies, S.A for treatment of patients 4 through 11 years old with a confirmed diagnosis of peanut allergy.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's website after the meeting. Background material is available at <https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee meeting link.