

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2018-D-0719]

**E6(R2) Good Clinical Practice: Integrated Addendum to E6(R1); International Council for Harmonisation; 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 entitled “E6(R2) Good Clinical Practice: Integrated Addendum to E6(R1).” The guidance was prepared under the auspices of the International Council for Harmonisation (ICH), formerly the International Conference on Harmonisation. The guidance amends the guidance entitled “E6 Good Clinical Practice: Consolidated Guidance (E6(R1))” to encourage implementation of improved and more efficient approaches to clinical trial design, conduct, oversight, recording, and reporting, and also updates standards regarding electronic records and essential documents. The guidance is intended to improve clinical trial quality and efficiency, while maintaining human subject protection and reliability of trial results.

**DATES:** The announcement of the guidance is published in the **Federal Register** on March 1, 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-2018-D-0719 for “E6(R2) Good Clinical Practice: Integrated Addendum to E6(R1).” 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 (CDER), Food and Drug Administration, 10001 New Hampshire Ave., Hillandale Building, 4th Floor, Silver Spring, MD 20993-0002, or 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. The guidance may also be obtained by mail by calling CBER at 1-800-835-4709 or 240-402-8010. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the guidance document.

**FOR FURTHER INFORMATION CONTACT:**

*Regarding the guidance:* Dianne Paraoan, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 3326, Silver Spring, MD 20993-0002, 301-796-2500; 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.

*Regarding the ICH:* Amanda Roache, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 1176, Silver Spring, MD 20993-0002, 301-796-4548.

**SUPPLEMENTARY INFORMATION:**

**I. Background**

In recent years, regulatory authorities and industry associations from around the world have participated in many important initiatives to promote international harmonization of regulatory requirements under the ICH. FDA has participated in several ICH

meetings designed to enhance harmonization, and FDA is committed to seeking scientifically based harmonized technical procedures for pharmaceutical development. One of the goals of harmonization is to identify and then reduce differences in technical requirements for drug development among regulatory agencies.

ICH was established to provide an opportunity for harmonization initiatives to be developed with input from both regulatory and industry representatives. FDA also seeks input from consumer representatives and others. ICH is concerned with harmonization of technical requirements for the registration of pharmaceutical products for human use among regulators around the world. The six founding members of the ICH are the European Commission; the European Federation of Pharmaceutical Industries Associations; FDA; the Japanese Ministry of Health, Labour, and Welfare; the Japanese Pharmaceutical Manufacturers Association; and the Pharmaceutical Research and Manufacturers of America. The Standing Members of the ICH Association include Health Canada and Swissmedic. Any party eligible as a Member in accordance with the ICH Articles of Association can apply for membership in writing to the ICH Secretariat. The ICH Secretariat, which coordinates the preparation of documentation, operates as an international nonprofit organization and is funded by the Members of the ICH Association.

The ICH Assembly is the overarching body of the Association and includes representatives from each of the ICH members and observers. The Assembly is responsible for the endorsement of draft guidelines and adoption of final guidelines. FDA publishes ICH guidelines as FDA guidance.

In the **Federal Register** of September 29, 2015 (80 FR 58492), FDA published a notice announcing the availability of a draft guidance entitled “E6(R2) Good Clinical Practice.” The notice gave interested persons an opportunity to submit comments on the “ADDENDUM” text added to ICH E6(R1) by November 30, 2015.

After consideration of the comments received and revisions to the guidance, a final draft of the guidance was submitted to the ICH Assembly and endorsed by the regulatory agencies in November 2016.

The guidance discusses approaches to clinical trial design, conduct, oversight, recording, and reporting as well as updated standards regarding electronic records and essential documents. This

guidance includes additions to ICH E6(R1) that are identified as “ADDENDUM” and are marked with vertical lines on both sides of the text. The additions to ICH E6(R1) are intended to encourage implementation of the described approaches and processes to improve clinical trial quality and efficiency while maintaining human subject protection and reliability of trial results. Evolutions in technology and risk management processes offer new opportunities to increase clinical trial efficiency, in part by focusing on trial activities essential to ensuring human subject protection and the reliability of trial results. For example, the guidance recommends sponsors implement a system to manage quality throughout clinical trials and recommends sponsors develop a systematic, prioritized, risk-based approach to monitoring clinical trials. The guidance provides additional detail regarding recommendations for use of electronic records and essential documents. The final guidance includes clarifications and additional detail on topics including, for example, validation of computerized systems and centralized monitoring.

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 “E6(R2) Good Clinical Practice: Integrated Addendum to E6(R1).” 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 guidance contains information collection provisions that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (21 U.S.C. 3501–3520). The collections of information in this guidance were approved under 0910–0843. This guidance also refers to previously approved collections of information found in FDA regulations. The collections of information found in 21 CFR part 11 have been approved under OMB control number 0910–0303; the collections of information found in 21 CFR part 56 have been approved under OMB control numbers 0910–0755; the collections of information found in 21 CFR part 312 have been approved under OMB control numbers 0910–0014 and 0910–0733; the collections of information found in 21 CFR part 314 have been approved under OMB control number 0910–0001; and the collections of information found 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 document at <https://www.regulations.gov>, <https://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm>, or <https://www.fda.gov/BiologicsBloodVaccines/GuidanceComplianceRegulatoryInformation/Guidances/default.htm>.

Dated: February 23, 2018.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2018–04154 Filed 2–28–18; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA–2017–N–5925]

#### Susceptibility Test Interpretive Criteria Recognized and Listed on the Susceptibility Test Interpretive Web Page; Reopening of a Public Docket; Request for Comments

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; reopening of a public docket; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA or the Agency) is reopening a docket for public comment on the susceptibility test interpretive criteria for antibacterial and antifungal drugs provided by FDA on its Susceptibility Test Interpretive Criteria web page (Interpretive Criteria web page) established on December 13, 2017. On the Interpretive Criteria web page, FDA recognizes, in whole or in part, susceptibility test interpretive criteria standards established by Standard Development Organizations (SDOs) and lists other susceptibility test interpretive criteria identified by FDA outside of the SDO process.

**DATES:** This notice is published in the **Federal Register** on March 1, 2018.

**ADDRESSES:** You may submit either electronic or written comments and information as follows:

#### *Electronic Submissions*

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- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://>