DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2019-D-1917]

Drug Abuse and Dependence Section of Labeling for Human Prescription Drug and Biological Products--Content and Format; Draft Guidance for Industry; Availability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SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for industry entitled "Drug Abuse and Dependence Section of Labeling for Human Prescription Drug and Biological Products--Content and Format." This guidance is intended to assist applicants in writing the DRUG ABUSE AND DEPENDENCE section of the labeling, as described in the regulations for the content and format of labeling for human prescription drug and biological products. The recommendations in this draft guidance will help ensure that the labeling is clear, concise, useful and informative, and, to the extent possible, consistent in content and format within and across drug and therapeutic classes.

DATES: Submit either electronic or written comments on the draft guidance by [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER] to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.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-2019-D-1917 for "Drug Abuse and Dependence Section of Labeling for Human Prescription Drug and
Biological Products--Content and Format. 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 http://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 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 draft guidance document.

FOR FURTHER INFORMATION CONTACT:  Iris Masucci, Center for Drug Evaluation
and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, 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

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a draft guidance for industry entitled "Drug Abuse
and Dependence Section of Labeling for Human Prescription Drug and Biological Products--
Content and Format." This draft guidance provides recommendations on the general principles
to consider when drafting the DRUG ABUSE AND DEPENDENCE section of the labeling, and
how to write, organize, and format the information within the DRUG ABUSE AND DEPENDENCE section of the labeling. The draft guidance provides recommendations on what information to include in the DRUG ABUSE AND DEPENDENCE section, including common terminology and definitions related to abuse and dependence, and how information related to topics presented in the DRUG ABUSE AND DEPENDENCE section should be distributed elsewhere in labeling.

This draft guidance is one in a series of guidances FDA is developing or has developed to assist applicants with the content and format of labeling for human prescription drug and biological products. In the Federal Register of January 24, 2006 (71 FR 3922), FDA published a final rule on labeling for human prescription drug and biological products. The final rule and additional guidances on labeling can be accessed at https://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/LawsActsandRules/ucm084159.htm. The labeling requirements and these guidances are intended to make information in prescription drug labeling easier for health care practitioners to access, read, and use.

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 of FDA on "Drug Abuse and Dependence Section of Labeling for Human Prescription Drug and Biological Products--Content and Format." 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 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 collections of information in 21 CFR 201.56 and 201.57 have been approved under OMB control number 0910-0572; the collections of information in 21 CFR 312.41 have been approved under OMB control number 0910-0014; the collections of information in 21 CFR 314.126(c), 314.70, and 314.97 have been approved under OMB control number 0910-0001; and the collections of information in 21 CFR 601.12 have been approved under OMB control number 0910-0338.

III. Electronic Access


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Principal Associate Commissioner for Policy.

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