this document for purposes of publication in the **Federal Register**.

Chyana Woodyard,

Federal Register Liaison, Centers for Medicare & Medicaid Services.

[FR Doc. 2026–24000 Filed 11–24–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[Office of Management and Budget #: 0970–0446]

Proposed Information Collection Activity; Temporary Assistance for Needy Families Expenditure Report, Form ACF–196R

AGENCY: Office of Family Assistance, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comments.

SUMMARY: The Administration for Children and Families' (ACF) Office of Family Assistance (OFA) is requesting a 3-year extension of the Temporary Assistance for Needy Families (TANF) Expenditure Report, Form ACF–196R (Office of Management and Budget #: 0970–0446, expiration January 31, 2027). ACF is proposing revisions to streamline instructions, improve organization and flow, and remove redundant reporting requirements.

DATES: Comments due January 25, 2027.

ADDRESSES: In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above. You can obtain copies of the proposed collection of information and submit comments by emailing *infocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: Form ACF–196R is used by states administering the TANF program to report quarterly expenditure data and to request quarterly grant funds. Failure to collect the data would seriously compromise OFA's and ACF's ability to monitor TANF expenditures and compliance with statutory requirements. These data are also needed to estimate outlays and to prepare reports and budget submissions for Congress. Since its last approval, the instructions have been streamlined in plain language and reorganized with formatting and clear section headers. Specifically, ACF has made the following updates to the form and instructions: Updated dates and references to more current status; simplified examples and explanations to support comprehension of reporting expectations; added table of contents and introduced plain language edits to improve clarity and the presentation of the document; and grayed out form fields where input is no longer required.

Respondents: State agencies administering the TANF program (50 states and the District of Columbia).

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
Form ACF–196R	51	4	9	1,836

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: 42 U.S.C. 609.

Samantha L. Illangasekare,

ACF/OPRE Certifying Officer.

[FR Doc. 2026–23967 Filed 11–24–26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–D–1188]

Over-the-Counter Monograph Order Requests: Format and Content; 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 final guidance for industry titled ''Over-the-Counter Monograph Order Requests: Format and Content.'' This guidance provides recommendations on the format and content of the information that a requestor should provide in an over-the-counter (OTC) monograph order request (OMOR) and identifies relevant guidance documents to assist requestors in preparing their OMORs. This guidance finalizes the draft guidance titled ''Over-the-Counter Monograph Order Requests (OMORs): Format and Content'' issued on April 13, 2023.

DATES: The announcement of the guidance is published in the **Federal Register** on November 25, 2026.

ADDRESSES: You may submit either electronic or written comments on any 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–2023–D–1188 for ''Over-the-Counter Monograph Order Requests: Format and Content.'' 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, 240–402–7500.

• 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, 240–402–7500.

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 Office of Drug Information, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 1109, 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: Cynthia Kim, Center for Drug Evaluation and Research (HFD–600), Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 22, Rm. 6464, Silver Spring, MD 20993–0002, 301–796–0879.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of a final guidance for industry titled ''Over-the-Counter Monograph Order Requests: Format and Content.'' As required by section 505G(l)(2) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 355h(l)(2)), this guidance discusses the format and content of data submissions, specifically OMORs, to FDA. This guidance is intended to assist requestors in preparing OMORs for submission to FDA under section 505G of the FD&C Act. This guidance provides FDA's recommendations on the format and content of the information that a requestor should provide in an OMOR and identifies relevant guidance documents to assist requestors in preparing their OMORs.

FDA agreed to specific performance goals and procedures described in the document ''Over-the-Counter Monograph User Fee Program Performance Goals and Procedures—Fiscal Years 2018–2022,'' commonly referred to as the OMUFA I Commitment Letter (the document can be accessed at *https://www.fda.gov/media/106407/download*). Among other things, in the OMUFA I Commitment Letter, FDA committed to issuing this guidance. The latest OMUFA Commitment Letter (OMUFA II Commitment Letter) can be found at *https://www.fda.gov/media/182750/download?attachment.*

This guidance finalizes the draft guidance titled ''Over-the-Counter Monograph Order Requests: Format and Content'' issued on April 13, 2023 (88 FR 22451). FDA considered comments received on the draft guidance as the guidance was finalized. Changes from the draft guidance to the final guidance include: (1) clarifying that because the type and quantity of data and information necessary to support a generally recognized as safe and effective determination may vary, not all of the information described in the guidance (*e.g.,* information identified for certain modules, section headings, or subsection headings described in the guidance) may be relevant to each particular OMOR; (2) clarifying the applicability of a financial certification (Form FDA 3454) or a financial disclosure (Form FDA 3455); and (3) clarifying quality information that may be relevant for an OMOR, including for a proposed active ingredient and for a drug product formulated using that active ingredient, as applicable in an OMOR.

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 ''Over-the-Counter Monograph Order Requests: Format and Content.'' 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.

FDA considered the applicability of Executive Order 14192, per OMB guidance in M–25–20, and finds this action to be neither an E.O. 14192 regulatory action nor an E.O. 14192 deregulatory action.

II. Paperwork Reduction Act of 1995

Under section 505G(o) of the FD&C Act, the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521) does not apply to collections of information made under section 505G of the FD&C Act. The information collections described in this guidance implement the following provisions of section 505G:

(1) Section 505G(b)(5) of the FD&C Act, which allows submission of administrative orders, OMORs, initiated at the request of a requestor.

(2) Section 505G(b)(6) of the FD&C Act, which allows requestors to provide certain information regarding safe nonprescription product marketing and use as a condition for filing a generally recognized as safe and effective request.

(3) Section 505G(d) of the FD&C Act confidentiality of information submitted to the Secretary, which requires FDA to make information submitted in support of an OMOR available to the public no later than the date of the proposed order unless it meets certain limitations on public availability.

(4) Section 505G(j) of the FD&C Act, which requires that all submissions under section 505G must be in electronic format.

(5) Section 505G(l)(1) of the FD&C Act, which requires FDA to issue guidance that specifies the procedures and principles for formal meetings between the Secretary and sponsors or requestors for drugs subject to section 505G.

(6) Section 505G(l)(2) of the FD&C Act, which requires FDA to issue guidance that specifies the format and content of data submissions to the Secretary under section 505G.

(7) Section 505G(l)(3) of the FD&C Act, which requires FDA to issue guidance that specifies the format of electronic submissions to the Secretary under section 505G.

Therefore, clearance by the Office of Management and Budget (OMB) under the PRA is not required for these collections of information. In addition, this guidance refers to previously approved FDA collections of information. The previously approved collections of information are subject to review by OMB under the PRA. The collections of information for OTC monograph products, OMORs, and the OTC Monograph User Fee Program have been approved under OMB control number 0910–0340. The information collections for submission of new drug applications and abbreviated new drug applications in 21 CFR part 314 are approved under OMB control number 0910–0001. The collections of information used by FDA to assess the environmental impact of Agency actions and to ensure that the public is informed of environmental analyses under 21 CFR part 25 have been approved under OMB control number 0910–0322.

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/regulatory-information/search-fda-guidance-documents*, or *https://www.regulations.gov*.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2018–D–4533]

Compounding Animal Drugs From Bulk Drug Substances: Compounding Under CGMP in Federally-Registered Facilities; Draft Guidance for Industry; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is extending the comment period for the notice of availability that appeared in the **Federal Register** of August 28, 2026. In the notice, FDA requested comments on the draft guidance for industry (GFI) #256B titled ''Compounding Animal Drugs from Bulk Drug Substances: Compounding under CGMP in Federally-Registered Facilities.'' The Agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: FDA is extending the comment period for the notice of availability published August 28, 2026 (91 FR 55602). Submit either electronic or written comments on the draft guidance by February 25, 2027, to ensure that the Agency considers your comments 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'').

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- 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–4533 for ''Compounding Animal Drugs from Bulk Drug Substances: Compounding under CGMP in Federally-Registered Facilities.'' 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, 240–402–7500.

- 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