you to submit your comment online through the *https://www.regulations.gov* website. To ensure the Commission considers your online comment, please follow the instructions on the web-based form.

If you file your comment on paper, write ''Regs BEMZ, PRA Comments, P084812'' on your comment and on the envelope, and mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580; or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024. If possible, please submit your paper comment to the Commission by courier or overnight service.

Because your comment will be placed on *https://www.regulations.gov,* you are solely responsible for making sure that your comment does not include any sensitive or confidential information. In particular, your comment should not include any sensitive personal information, such as your or anyone else's Social Security number, date of birth, driver's license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include sensitive health information, such as medical records or other individually identifiable health information. In addition, your comment should not include any ''trade secret or any commercial or financial information which . . . is privileged or confidential,'' as provided by section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2), including in particular, competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled ''Confidential,'' and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. Your comment will be kept confidential only if the FTC General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted on *https://www.regulations.gov,* we cannot redact or remove your comment from that website, unless you submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before July 16, 2021. For information on the Commission's privacy policy, including routine uses permitted by the Privacy Act, see *https://www.ftc.gov/siteinformation/privacy-policy.*

Josephine Liu,

Assistant General Counsel for Legal Counsel.

[FR Doc. 2021–10285 Filed 5–14–21; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2021–0053]

The Systematic Review Report for Diagnosis and Treatment of Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (ME/CFS): Request for Comment

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC) in the Department of Health and Human Services (HHS) announces the opening of a docket to obtain comment on the systematic review draft report for Diagnosis and Treatment of Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (ME/CFS). The draft report describes inclusion and exclusion criteria to identify relevant literature, outlines the approach for evaluating study quality, and summarizes the systematic review results. The report, once finalized, is intended to support the anticipated development of future clinical practice guidelines, which would guide physicians in managing and providing care for patients with ME/CFS. Currently there are no federal guidelines for management of ME/CFS. CDC has commissioned the Pacific Northwest Evidence-Based Practice Center at Oregon Health & Science University to conduct a systematic review of the publicly available scientific literature and now seeks public comment to inform the final report. In particular, CDC seeks data and information, including reports and manuscripts that are pending publications or are not available through indexed bibliographic databases. Access to pertinent scientific information from research and evidence-based clinical practice may be used to inform the final report. The anticipated CDC guideline would assist clinicians by outlining management practices for patients with ME/CFS.

DATES: Written comments must be received on or before August 16, 2021.May 17, 2021.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2021–0053 by either of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Anindita Issa, MD, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24–12, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *http://regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov. Do not submit comments by email. CDC does not accept comments by email.*

FOR FURTHER INFORMATION CONTACT: For technical information on the systematic review report for ME/CFS, contact Anindita Issa, MD, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24–12, Atlanta, Georgia 30329. Telephone: 404–718–3959; email: *cfs@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Public Participation

Interested persons or organizations are invited to participate by submitting written views, recommendations, and data related to the draft report, including perspectives on and experiences with diagnosis and management of ME/CFS illness. In addition, CDC invites comments specifically on topics for pharmacological or non-pharmacological treatments.

Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore,

do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted in preparation of the final systematic review report and may revise it as appropriate. Do not submit comments by email. CDC does not accept comments by email.

Background

CDC has commissioned the Pacific Northwest Evidence-Based Practice Center at Oregon Health and Science University to conduct a systematic review of the publicly available scientific literature for ME/CFS (systematic review). Once finalized, the systematic review report is intended to support the anticipated development of future agency clinical practice guidelines. The anticipated CDC guideline would assist physicians and other clinicians by outlining management practices for caring for patients with ME/CFS.

Public comment may inform the final systematic review report and may be used to inform development of a clinical guideline and related materials, which would help clinicians diagnose and treat patients with ME/CFS.

Dated: May 11, 2021.

Sandra Cashman,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2021–10271 Filed 5–14–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10398 #71]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance [1] related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. On October 23, 2011, OMB approved our initial request to use the generic clearance process under control number 0938–1148 (CMS–10398). It was last approved on April 26, 2021, via the standard PRA process which included the publication of 60- and 30-day **Federal Register** notices. The scope of the April 2021 umbrella accounts for Medicaid and CHIP State plan amendments, waivers, demonstrations, and reporting. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit comments regarding our burden estimates or any other aspect of this collection of information, including: The necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by June 1, 2021.

ADDRESSES: When commenting, please reference the applicable form number (see below) and the OMB control number (0938–1148). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address:

CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: CMS–10398 (#71)/OMB control number: 0938–1148, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may access CMS' website at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION: Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

Generic Information Collection

1. *Title of Information Collection:* Reporting Requirements for State Planning Grants for Qualifying Community-Based Mobile Crisis Intervention Services During the COVID–19 Emergency; *Type of Information Collection Request:* New collection; *Use:* On March 11, 2021, President Biden signed the American Rescue Plan Act of 2021 (ARP) (Pub. L. 117–2). Under ARP section 9813, planning grantees must submit quarterly progress reports and a final progress report. The reports should include narrative updates on planning grant activities, as well as information on each recipient's approved work plan, as specified in each recipient's approved application in accordance with the ''Section 9813 State Planning Grants for Mobile Crisis Intervention Services Cooperative Agreement.'' To ensure maximum state flexibility and to reduce the reporting burden on states as much as possible, states will submit quarterly and final progress reports in their own preferred format. CMS will not require states to use a standardized template or form. When ready, the Notice of Funding Opportunity will be posted on the *Grants.gov* website; *Form Number:* CMS–10398 (#71) (OMB control number: 0938–1148); *Frequency:* Quarterly; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 20; *Total Annual Responses:* 80; *Total Annual Hours:*

[1] *https://www.whitehouse.gov/sites/whitehouse.gov/files/omb/assets/inforeg/PRA_Gen_ICRs_5-28-2010.pdf.*