Register Notice 2009–03, 74 FR 7285 (February 17, 2009).

CALENDAR OF REPORTING DATES FOR WISCONSIN SPECIAL ELECTION

Report	Close of books [1]	Reg./cert. & overnight mailing deadline	Filing deadline
Committees Involved in *Only* the Special Primary (02/18/2020) Must File:			
Year-End	Waived		
Pre-Primary	01/29/2020	02/03/2020	02/06/2020
April Quarterly	03/31/2020	04/15/2020	04/15/2020
Committees Involved in Both the Special Primary (02/18/2020) and Special General (05/12/2020) Must File:			
Year-End	Waived		
Pre-Primary	01/29/2020	02/03/2020	02/06/2020
April Quarterly	03/31/2020	04/15/2020	04/15/2020
Pre-General	04/22/2020	04/27/2020	04/30/2020
Post-General	06/01/2020	06/11/2020	06/11/2020
July Quarterly	06/30/2020	07/15/2020	07/15/2020
Committees Involved in *Only* the Special General (05/12/2020) Must File:			
Pre-General	04/22/2020	04/27/2020	04/30/2020
Post-General	06/01/2020	06/11/2020	06/11/2020
July Quarterly	06/30/2020	07/15/2020	07/15/2020

Dated: November 13, 2019.

On behalf of the Commission.

Ellen L. Weintraub,

Chair, Federal Election Commission.

[FR Doc. 2019–24999 Filed 11–18–19; 8:45 am]

BILLING CODE 6715–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–R–142]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send 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 January 21, 2020.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. 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: Document Identifier/OMB Control Number ____, 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 make your request using one of following:

1. Access CMS' website address at website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

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

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement

[1] The reporting period always begins the day after the closing date of the last report filed. If the committee is new and has not previously filed a report, the first report must cover all activity that occurred before the committee registered as a political committee up through the close of books for the first report due.

and associated materials (see **ADDRESSES**).

CMS–R–142 Examination and Treatment for Emergency Medical Conditions and Women in Labor (EMTALA)

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collection

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Examination and Treatment for Emergency Medical Conditions and Women in Labor (EMTALA); *Use:* Pursuant to section 1866(a)(1)(I) of the Act, Congress has mandated that the Secretary enforce section 1867 of the Act. Under section 1867, effective August 1, 1986, hospitals may continue to participate in the Medicare program only if they are not out of compliance with its provisions. Continued Paper Work Reduction Act (PRA) approval of the regulation sections cited below will promote uniform and thorough application of the section 1866 and 1867 requirements. They will also provide information when requested by Congress and other interested parties regarding the implementation of the statute. During 2004 through 2018, approximately 8,146 complaints were received, approximately 7,770 of those complaints were investigated, and approximately 3,567 EMTALA deficiencies were found. During Federal fiscal years 2001 through 2005 the Inspector General's Office imposed civil monetary penalties on hospitals in 105 cases, for a total of $2,645,750 in penalties. An audit completed by the Office of Inspector General (OIG) (entitled, Office of Inspector General: Implementation and Enforcement of the Examination and Treatment for Emergency Medical Conditions and Women in Labor by the Health Care Financing Administration, April 1995, A–06–93–00087) determined that CMS's implementation of the Act was generally effective, but Regional Offices (RO) were not consistent with conducting timely investigations, sending acknowledgments to complaints, ensuring that investigations were thorough, or ensuring that violations were referred to the OIG in accordance with CMS policy for possible civil monetary penalty action. OIG further concluded that without proper compliance, there is an increased risk that individuals with emergency medical conditions will not receive the treatment needed to stabilize their condition, which may place them in greater risk of death. *Form Number:* CMS–R–142 (OMB control number: 0938–0667); *Frequency:* Occasionally; *Affected Public:* Private Sector; Business or other for-profits, Not-for-profit institutions; *Number of Respondents:* 5,291; *Total Annual Responses:* 5,291; *Total Annual Hours:* 5,291. (For policy questions regarding this collection contact Renate Dombrowski at (410) 786–4645.)

Dated: November 14, 2019.

William N. Parham, III,

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–25065 Filed 11–18–19; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Evaluation of the National Human Trafficking Hotline Program (New Collection)

AGENCY: Office of Planning, Research, and Evaluation; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) is proposing a data collection activity as part of the Evaluation of the National Human Trafficking Hotline (NHTH) Program. This data collection activity will examine the experiences of individuals who seek assistance from the NHTH after their interactions with the NHTH. The study will collect information via voluntary phone and Web-based surveys at two time points: (1) Immediately after an individual has contacted the NHTH by phone, text, or live online chat; and (2) two weeks later.

DATES: *Comments due within 30 days of publication.* OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Copies of the proposed collection may be obtained by emailing *OPREinfocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The purpose of the proposed data collection activity is to document and examine the following: Why individuals contact the NHTH; hotline users' perceptions of hotline staff's knowledge and skills; the extent to which users felt their interaction was helpful, they were supported by the NHTH, they were satisfied with the NHTH, and their needs were met by the interaction; and outcomes from NHTH interactions (*e.g.,* users' knowledge and use of available resources and referrals). The proposed data collection activity includes a two-phase approach to obtain information from individuals after their contact (via phone, text, or live online chat) with the NHTH. The proposed information collection activities are (1) an integrated voice response telephone survey or Web-based survey immediately after NHTH contact; and (2) a telephone or Web-based survey approximately two weeks after completion of the first survey. The survey immediately after contact with the NHTH will be offered to all individuals who contact the NHTH during the data collection period and includes questions focused on users' experiences and satisfaction with their NHTH interaction. The follow-up survey will be administered two weeks later with a sample of respondents who completed the immediate survey and