treatment/referrals, etc. Information on facility location, key populations served, and workforce capacity is also needed to identify areas in need of expanded support to deliver these services. There is no other data source that comprehensively collects this information.

The participation of respondents is voluntary. There is no cost to the respondents other than their time. Through their participation, respondents will help to improve programs to prevent HIV infection as well as services for those who already have HIV. The total estimated annualized burden is 618 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Facility administrative staff	MMP Facility Survey	1,200 225	1	30/60 5/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-21-1227]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled "Assessment of Ill Worker Policies Study" to the Office of Management and Budget (OMB) for review and approval. CDC previously published a "Proposed Data Collection Submitted for Public Comment and Recommendations" notice on September 14, 2020, to obtain comments from the public and affected agencies. CDC received one comment related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

- (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

- (c) Enhance the quality, utility, and clarity of the information to be collected:
- (d) Minimize the burden of the collection of information on those who are to respond, including, through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and
- (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570. Comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/ do/PRAMain Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202) 395-5806. Provide written comments within 30 days of notice publication.

Proposed Project

Assessment of Ill Worker Policies Study (OMB Control No. 0920–1227, Exp. 5/31/2021)—Revision—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The CDC is requesting a three-year Paperwork Reduction Act (PRA) clearance for a Revision information collection request (ICR) for a research program focused on identifying the environmental causes of foodborne illness and improving environmental

public health practice. This research program is conducted by the Environmental Health Specialists Network (EHS-Net), a collaborative project of the CDC, U.S. Food and Drug Administration (FDA), U.S. Department of Agriculture (USDA), and eight state and local public health programs (Franklin County, OH; Tennessee; Minnesota; Rhode Island; New York; New York City, NY; Southern Nevada Health District, NV; and Harris County, TX).

This ICR aims to assess whether an educational intervention will result in either the development or enhancement of restaurant ill worker policies. This will be accomplished by interviewing restaurant managers and observing restaurant practices in 320 randomly selected and assigned restaurants in the EHS-Net catchment areas. There will be two or three site visits depending upon which group the restaurants are assigned to, that is, the intervention or the control group. An initial visit will be used to observe baseline conditions and to provide the intervention only to the restaurants selected to receive it. A second visit will be used to determine if the policies have changed and to introduce the intervention to the control restaurants (if it is deemed successful), and a final follow up visit to the control restaurants that received the intervention on the second visit (if they receive the intervention). Initial success for the intervention will be measured by whether three or more intervention restaurants in each EHS-Net catchment area either develop a written ill worker management plan (if they did not have one at the pre-intervention evaluation) or enhanced their policies (e.g., added provisions addressing reasons why ill workers reported working while ill).

Although approved in 2018, NCEH and its program partners needed to prioritize other data collections over this study, and then delayed the current study due to the COVID-19 pandemic.

NCEH partners provided feedback to refine this research protocol, to revise the ICR, and to begin this study in 2021. NCEH is requesting approval for revisions which fall into three categories: (1) Changes to strengthen the study, based on recent experience and stakeholder feedback; (2) changes to respond to the COVID–19 pandemic, and (3) a change in one participating site.

NCEH is requesting a revised PRA clearance for 820 responses per year and

for a time burden of 261 hours per year. These changes result in a decrease of 1,307 responses and 91 hours per year relative to the 2018 PRA clearance. There is no cost to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Restaurant Managers (Intervention and Control Restaurants).	Manager Recruiting Script	237	1	3/60
Restaurant Managers (Intervention Restaurants)	Manager Informed Consent and Interview.	53	2	20/60
	Intervention Log	53	1	30/60
Restaurant Managers (Control Restaurants)	Manager Informed Consent and Interview.	53	3	20/60
	Intervention Log	53	1	30/60
Health Department Workers (Intervention and Control Restaurants).	Restaurant Observation Form	106	2	30/60

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-R-148]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 (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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by March 10, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

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: https://www.cms.gov/Regulations-and-Guidance/Legislation/
PaperworkReductionActof1995/PRA-Listing.html.

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

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment:

1. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Limitations on Provider Related Donations and Health Care Related Taxes, Medicaid and Supporting Regulations in 42 CFR 433.68 through 433.74; Use: States may elect to submit a waiver to CMS for the broad based and/or uniformity requirements for any health care related tax program which does not conform to the broad based and uniformity requirements. It is also the responsibility of each State to demonstrate that their tax program(s) do not violate the hold harmless provision. For a waiver to be approved and a determination that the hold harmless provision is not violated, States must submit written documentation which satisfies the regulatory requirements. Without this information, the amount of FFP (Federal financial participation) payable to a State cannot be correctly determined. Form Number: CMS-R-148