providers exploring their experiences with the intervention and thoughts about providing PrEP clinical services; and a clinic assessment completed by clinic staff every six months to describe the current implementation of PrEP services at their clinical site. These data will inform ongoing practice improvement in PrEP clinical services and increase understanding of provider experiences with providing PrEP clinical services.

It is expected that half of screened persons will meet study eligibility. For all Aims we anticipate that screening and completion of the locator form will each take five minutes. Study staff will assist Aim 1 participants with onboarding the CleverCap device and mChoice app, a process that will take 20 minutes. Aim 1 participants will complete the baseline survey once (anticipated 30 minutes completion time) and the follow-up survey four times (anticipated completion time 30 minutes each) over their 12-month participation period. Total study enrollment for Aim 1 is 400, over the three-year data collection period the estimated annual enrollment is 134. Aims 2 and 3 interviews will take 60 minutes to complete. For Aim 2, total study enrollment is 30, over the three-year data collection period the estimated annual enrollment is 10. For Aim 3, total study enrollment is 20, over the three-year data collection period the estimated annual enrollment is seven. Additionally, a single Aim 3 participant at each of the four participating clinic sites will complete a clinic assessment form every six months throughout the study period.

The total number of burden hours is 1,323 across 36 months of data collection. The total estimated annualized burden hours are 441. There are no costs to the participants other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)	Total burden (in hr)
Aim 1 participants—YMSM General public, adults.	Aim 1 Participant Eligibility Screener	268	1	5/60	22
Aim 1 participants—YMSM General public, adults.	Aim 1 Participant Locator Form	134	1	5/60	12
Aim 1 participants—YMSM General public, adults.	Aim 1 mChoice Onboarding Guide ..	134	1	20/60	45
Aim 1 participants—YMSM General public, adults.	Aim 1 Participant Baseline Survey ..	134	1	30/60	67
Aim 1 participants—YMSM General public, adults.	Aim 1 Participant Follow-up Survey	134	4	30/60	268
Aim 2 participants—YMSM General public, adults.	Aim 2 Participant Eligibility Screener	20	1	5/60	2
Aim 2 participants—YMSM General public, adults.	Aim 2 Participant Locator Form	10	1	5/60	1
Aim 2 participants—YMSM General public, adults.	Aim 2 Participant Interview Guide ...	10	1	1.0	10
Aim 3 participants—providers General public, adults.	Aim 3 Participant Eligibility Screener	14	1	5/60	2
Aim 3 participants—providers General public, adults.	Aim 3 Participant Locator Form	7	1	5/60	1
Aim 3 participants—providers General public, adults.	Aim 3 Participant Interview Guide ...	7	1	1.0	7
Aim 3 participant—clinic staff respondent, 1 per clinic site General public, adults.	Aim 3 Clinic Assessment	4	2	30/60	4
TOTAL	.....	.....	.....	.....	441

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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BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10237 and CMS–10407]

Agency Information Collection Activities: Proposed Collection; 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 (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 August 12, 2022.

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.*

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

CMS–10237—Applications for Part C Medicare Advantage, 1876 Cost Plans, and Employer Group Waiver Plans to Provide Part C Benefits

CMS–10407—Summary of Benefits and Coverage and Uniform Glossary

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:* Revision of a currently approved collection; *Title of Information Collection:* Applications for Part C Medicare Advantage, 1876 Cost Plans, and Employer Group Waiver Plans to Provide Part C Benefits; *Use:* Collection of this information is mandated by the Code of Federal Regulations, MMA, and CMS regulations at 42 CFR 422, subpart K, in ''Application Procedures and Contracts for Medicare Advantage Organizations.'' In addition, the Medicare Improvement for Patients and Providers Act of 2008 (MIPPA) further amended titles XVII and XIX of the Social Security Act.

This information collection includes the process for organizations wishing to provide healthcare services under MA plans. These organizations must complete an application annually (if required), file a bid, and receive final approval from CMS. The MA application process has two options for applicants that include (1) request for new MA product or (2) request for expanding the service area of an existing product. CMS utilizes the application process as the means to review, assess and determine if applicants are compliant with the current requirements for participation in the MA program and to make a decision related to contract award. This collection process is the only mechanism for organizations to complete the required MA application process. *Form Number:* CMS–10237 (OMB control number: 0938–0935); *Frequency:* Yearly; *Affected Public:* Private Sector (Business or other for-profits, Not-for-Profit Institutions); *Number of Respondents:* 500; *Total Annual Responses:* 500; *Total Annual Hours:* 9,173. (For policy questions regarding this collection contact Keith Penn-Jones at 410–786–3104.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Summary of Benefits and Coverage and Uniform Glossary; *Use:* This information collection will ensure that over 30 million consumers shopping for or enrolled in private, individually purchased, or non-federal governmental group health plan coverage receive the consumer protections of the Affordable Care Act. Employers, employees, and individuals will use this information to compare coverage options prior to selecting coverage and to understand the terms of, and extent of medical benefits offered by, their coverage (or exceptions to such coverage or benefits) once they have coverage. *Form Number:* CMS–10407 (OMB control number 0938–1146); *Frequency:* Annually; *Affected Public:* Private Sector—Business or other for-profits and Not-for-profit institutions; *Number of Respondents:* 90,805; *Number of Responses:* 10,507,165; *Total Annual Hours:* 204,140. (For policy questions regarding this collection contact Daniel Kidane at *daniel.kidane@cms.hhs.gov.*)

Dated: June 7, 2022.

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10668 and CMS–10455]

Agency Information Collection Activities: Proposed Collection; 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 (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