

confidential; marketing materials; study types not included in the review; or information on indications not included in the review cannot be used by the EPC Program. This is a voluntary request for information, and all costs for complying with this request must be borne by the submitter.

The draft of this review will be posted on AHRQ's EPC Program website and available for public comment for a period of 4 weeks. If you would like to be notified when the draft is posted, please sign up for the email list at: <https://www.effectivehealthcare.ahrq.gov/email-updates>.

The systematic review will answer the following questions. This information is provided as background. AHRQ is not requesting that the public provide answers to these questions.

Key Questions (KQs)

KQ 1. What is the effectiveness of provider-to-provider telehealth for rural patients?

a. What is the impact of provider-to-provider telehealth on *rural patient and population* outcomes?

b. What is the impact of provider-to-provider telehealth on *healthcare providers*?

c. What is the impact of provider-to-provider telehealth on *private and public* (ex. CMS, TriCare, VA, etc.) payers?

d. What adverse events or unintended consequences are associated with provider-to-provider telehealth for rural patients?

e. What are the methodological weaknesses of the identified effectiveness studies of provider-to-provider telehealth for rural patients and what improvements in study design (e.g., focus on relevant comparisons and outcomes) might increase the impact of future research?

KQ 2. What is the effectiveness of implementation strategies for provider-to-provider telehealth in rural areas?

a. What is the uptake of different types of provider-to-provider telehealth in rural areas?

○ Who are the current patients, providers, and payers engaged in provider-to-provider telehealth in rural areas?

○ What factors affect whether provider-to-provider telehealth in rural areas can be sustained?

b. Which barriers and facilitators impact adoption and implementation of provider-to-provider telehealth in rural areas?

c. Which strategies are effective in sustaining provider-to-provider telehealth in rural areas?

d. What are the methodological weaknesses of the identified studies of implementation and sustainability of provider-to-provider telehealth in rural areas and what improvements in study design (e.g., focus on relevant comparisons and outcomes) might increase the impact of future research?

Populations, Interventions, Comparators, Outcomes, Settings

• Population(s)

○ Rural individual patients, patient families/care partners, and patient populations.

○ Healthcare providers (individuals and organizations) who provide health care services to rural patients or populations.

• Providers include any profession or occupation providing formal, paid services.

• Family or informal care partners are not considered providers.

○ Payers who pay for healthcare services for rural patients or populations.

• Interventions

○ Provider-to-provider telehealth defined as: Any telecommunications facilitated interaction among, or support for, healthcare professionals designed to improve access, quality of care, or health outcomes for rural patients and populations.

• Comparators

○ *KQ1*: Other telehealth facilitated care (not provider-to-provider), usual (in-person) provider-to-provider supports, no interaction or no care.

○ *KQ2*: Different strategies for dissemination, implementation, or spread; no strategies; time periods prior to implementation.

• Outcomes

○ *KQ1*: Clinical outcomes for the identified conditions (patient-reported outcomes, mortality, morbidity, such as function, illness recovery, infection); Economic outcomes such as return on investment, cost, volume of visits, and resource use, including length of stay and readmissions; Intermediate Outcomes: Patient satisfaction, behavior (such as care-seeking and compliance), and decisions such as completion of treatment, or satisfaction with less travel to access healthcare; Provider satisfaction, behavior, and decisions such as choice of treatment or antibiotic stewardship; Access measures and indicators including but not limited to time to diagnosis or time to treatment.

○ *KQ2*: Indicators and measures of uptake (e.g., rates of use, timing to implementation) and characteristics of users; categories and descriptors of barriers and facilitators; categories and descriptors of strategies.

• Settings

○ Outpatient (primary care and specialty care), inpatient, prehospital and emergency care, post-acute and long-term care.

○ Civilian, Veterans Administration, or military.

○ Health care and non-healthcare settings where health services are delivered including in the home.

○ U.S. relevant settings [Note that studies from countries with significantly different healthcare systems and fewer resources (e.g., low-income countries) are excluded.]

Dated: February 24, 2021.

Marquita Cullom,

Associate Director.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10175]

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 April 1, 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 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:* Certification Statement for Electronic File Interchange Organizations (EFIOs) that submit National Provider Identifier (NPI) data to the National Plan and Provider Enumeration System (NPPES); *Use:* the EFI process allows organizations to submit NPI application

information on large numbers of providers in a single file. Once it has obtained and formatted the necessary provider data, the EFIO can electronically submit the file to NPPES for processing. As each file can contain up to approximately 25,000 records, or provider applications, the EFI process greatly reduces the paperwork and overall administrative burden associated with enumerating providers. It is essential to collect this information from the EFIO to ensure that the EFIO understands its legal responsibilities as an EFIO and attests that it has the authority to act on behalf of the providers for whom it is submitting data. In short, the certification statement, which must be signed by an authorized official of the EFIO, serves as a safeguard against EFIOs attempting to obtain NPIs for illicit or inappropriate purposes. *Form Number:* CMS–10175 (OMB control number 0938–0984); *Frequency:* Once, Annually; *Affected Public:* Private Sector, State, Business, and Not-for Profits; *Number of Respondents:* 32; *Number of Responses:* 32; *Total Annual Hours:* 8. (For questions regarding this collection contact DaVona Boyd at 410–786–7483.)

Dated: February 25, 2021.

William N. Parham, III,

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

[FR Doc. 2021–04274 Filed 3–1–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10054 and CMS–10632]

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 May 3, 2021.

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 <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:

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–10054 New Technology Services for Ambulatory Payment Classifications under the Outpatient Prospective Payment System
CMS–10632 Evaluating Coverage to Care in Communities