

EXHIBIT 2—ESTIMATED COST BURDEN OVER 3 YEARS—Continued

Type of information collection	Number of respondents	Total burden hours	Average hourly wage rate *	Total cost burden
In-person	600	500	33.51	16,755
Total	32,700	10,150	na	340,127

*Based upon the average wages for 29-000 (Healthcare Practitioner and Technical Occupations), "National Compensation Survey: Occupational Wages in the United States, May 2009," U.S. Department of Labor, Bureau of Labor Statistics.

Request for Comments

In accordance with the Paperwork Reduction Act, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ healthcare research and healthcare information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information upon the respondents, including the use of automated collection techniques or other forms of information technology.

Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.

Sharon B. Arnold,
Deputy Director.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-R-138 and CMS-10332]

Agency Information Collection Activities: Submission for OMB Review; 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 (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 October 16, 2017.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395-5806 or, Email: *OIRA_submission@omb.eop.gov*.

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' Web site 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 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:* Medicare Geographic Classification Review Board Procedures and Criteria; *Use:* During the first few years of IPPS, hospitals were paid strictly based on their physical geographic location concerning the wage index (Metropolitan Statistical Areas (MSAs)) and the standardized amount (rural, other urban, or large urban). However, a growing number of hospitals became concerned that their payment rates were not providing accurate compensation. The hospitals argued that they were not competing with the hospitals in their own geographic area, but instead that they were competing with hospitals in neighboring geographic areas. At that point, Congress enacted Section 1886(d)(10) of the Act which enabled hospitals to apply to be considered part of neighboring geographic areas for payment purposes based on certain criteria. The application and decision process is administered by the MGCRB which is not a part of CMS so that CMS could not be accused of any untoward

action. However, CMS needs to remain apprised of any potential payment changes. Hospitals are required to provide CMS with copy of any applications that they made to the MGCRB. CMS also developed the guidelines for the MGCRB that were the interim final issue of the **Federal Register**, and must ensure that the MGCRB properly applied the guidelines. This check and balance process also contributes to limiting the number of hospitals that ultimately need to appeal their MGCRB decisions to the CMS Administrator. *Form Number:* CMS-R-138 (OMB control number: 0938-0573); *Frequency:* Occasionally; *Affected Public:* Businesses or other for-profits and Not-for-profit institutions; *Number of Respondents:* 300; *Total Annual Responses:* 300; *Total Annual Hours:* 300. (For policy questions regarding this collection contact Noel Manlove at 410-786-5161.)

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Disclosure Requirement for the In-Office Ancillary Services Exception; *Use:* Section 6003 of the ACA established a disclosure requirement for the in-office ancillary services exception to the prohibition of physician self-referral for certain imaging services. This section of the ACA amended section 1877(b)(2) of the Social Security Act by adding a requirement that the referring physician informs the patient, at the time of the referral and in writing, that the patient may receive the imaging service from another supplier. The implementing regulations are at 42 CFR 411.355(b)(7).

Physicians who provide certain imaging services (MRI, CT, and PET) under the in-office ancillary services exception to the physician self-referral prohibition are required to provide the disclosure notice as well as the list of other imaging suppliers to the patient. The patient will then be able to use the disclosure notice and list of suppliers in making an informed decision about his or her course of care for the imaging service. CMS would use the collected information for enforcement purposes. Specifically, if we were investigating the referrals of a physician providing advanced imaging services under the in-office ancillary services exception, we would review the written disclosure in order to determine if it satisfied the requirement. *Form Number:* CMS-10332 (OMB control number: 0938-1133); *Frequency:* Occasionally; *Affected Public:* State, Local, and Tribal Governments; *Number of Respondents:* 7,100; *Total Annual Responses:*

759,700; *Total Annual Hours:* 19,638. (For policy questions regarding this collection contact Laura Dash at 410-786-8623.)

Dated: September 11, 2017.

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

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Public Comment Request; Revision of a Currently Approved Information Collection (ICR-REV); State Plan for Assistive Technology (OMB Approval Number 0985-0048)

AGENCY: Administration for Community Living, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to comment on ACL's intention to collect information necessary to determine grantee compliance with Section 4 of the Assistive Technology Act of 1998, as amended (AT Act). Under the Paperwork Reduction Act of 1995 (PRA), Federal agencies are required to publish a notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the proposed action. This notice solicits comments on a proposed revision to an existing information collection related to the State Grants for Assistive Technology Program State Plan for AT, formerly the 664 Report.

DATES: Submit written or electronic comments on the collection of information by November 14, 2017.

ADDRESSES: Submit electronic comments on the collection of information to: *Robert.Groenendaal@acl.hhs.gov*. Submit written comments on the collection of information by mail to Robert Groenendaal, U.S. Department of Health and Human Services, Administration for Community Living, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Robert Groenendaal at (202) 795-7356 or *Robert.Groenendaal@acl.hhs.gov*.

SUPPLEMENTARY INFORMATION: Section 4 of the AT Act establishes formula grants to states to support comprehensive statewide programs (Statewide AT Programs) that conduct activities that improve access to and acquisition of AT devices and services for individuals with disabilities across the lifespan and across a wide array of disabilities, and their family members, guardians, advocates, and authorized representatives. State Grants for AT program conducts the following state-level and state leadership activities: State financing, device demonstration, device loans, device reutilization, training and technical assistance, public awareness, and information and referral. 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. "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 provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or update of an existing collection of information, before submitting the collection to OMB for approval. The proposed data collection represents a revision of a currently approved collection (ICR-Rev). In order to comply with the above requirement, ACL is requesting approval of a revision of a previously approved collection, the State Grants for Assistive Technology Program State Plan for AT, formerly known as the 664 report (0985-0048).

With respect to the following collection of information, ACL invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility; (2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The State Plan for AT is submitted every three years and updated annually by all State Grants for AT programs