

## ESTIMATED ANNUALIZED BURDEN HOURS

| Type of respondent                          | Form name                        | Number of respondents | Number of responses per respondent | Average burden per response (in hours) |
|---------------------------------------------|----------------------------------|-----------------------|------------------------------------|----------------------------------------|
| Parent/guardian of children aged 6–12 years | Screener Script Guide .....      | 667                   | 1                                  | 10/60                                  |
| Child participants aged 6–12 years .....    | Seat Belt Fit Measurements ..... | 142                   | 1                                  | 2.5                                    |

**Leroy A. Richardson,**

*Chief, Information Collection Review Office,  
Office of Scientific Integrity, Office of the  
Associate Director for Science, Office of the  
Director, Centers for Disease Control and  
Prevention.*

[FR Doc. 2016–13849 Filed 6–10–16; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

**[Document Identifiers: CMS–R–142 and  
CMS–588]**

#### 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 any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) 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 9, 2016.

**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 the following:

1. Access CMS' Web site address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov).

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

**FOR FURTHER INFORMATION CONTACT:** Reports Clearance Office at (410) 786–1326.

#### 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–R–142 Examination and Treatment for Emergency Medical Conditions and Women in Labor;**

#### **CMS–588 Electronic Funds Transfer Authorization Agreement**

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.

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; *Use:* In accordance with regulation sections 488.18, 489.20 and 489.24, during Medicare surveys of hospitals and State agencies CMS will review hospital records for lists of on-call physicians, and will review and obtain the information which must be recorded on hospital medical records for individuals with emergency medical conditions and women in labor, and the emergency department reporting information Medicare participating hospitals and Medicare State survey agencies must pass on to CMS. Additionally, CMS will use the QIO Report assessing whether an individual had an emergency condition and whether the individual was stabilized to determine whether to impose a CMP or physician exclusion sanctions. Without such information, CMS will be unable to make the hospital emergency services compliance determinations that Congress expects CMS to make under sections 1154, 1866 and 1867 of the Act. *Form Number:* CMS–R–142 (OMB control number: 0938–0667); *Frequency:* Occasionally; *Affected Public:* Private Sector; *Number of Respondents:* 6,149; *Total Annual Responses:* 6,149; *Total Annual Hours:* 1. (For policy questions regarding this collection contact Renate Dombrowski at 410–786–4645.)

2. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: Electronic Funds Transfer Authorization Agreement; *Use*: The information is needed to allow providers to receive funds electronically in their bank accounts. *Form Number*: CMS-588 (OMB control number: 0938-0626); *Frequency*: On occasion; *Affected Public*: Business or other for-profit, Not-for-profit institutions; *Number of Respondents*: 45,807; *Total Annual Responses*: 45,807; *Total Annual Hours*: 22,543. (For policy questions regarding this collection contact Kimberly McPhillips at 410-786-4645.)

Dated: June 7, 2016.

**William N. Parham, III,**

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

[FR Doc. 2016-13800 Filed 6-10-16; 8:45 am]

**BILLING CODE 4120-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10105, CMS-10191, CMS-10525, CMS-10623, and CMS-R-246]

### 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 any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) 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, 2016.

**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 the following:

1. Access CMS' Web site address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov).

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

**FOR FURTHER INFORMATION CONTACT**: Reports Clearance Office at (410) 786-1326.

### 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-10105 National Implementation of In-Center Hemodialysis CAHPS Survey  
CMS-10191 Medicare Parts C and D Program Audit Protocols and Data Requests  
CMS-10525 Program of all-Inclusive Care for the Elderly (PACE) Quality Data Entry in CMS Health Plan Monitoring System  
CMS-10623 Testing Experience and Functional Tools Demonstration:

Personal Health Record (PHR) User Survey  
CMS-R-246 Medicare Advantage, Medicare Part D, and Medicare Fee-For-Service Consumer Assessment of Healthcare Providers and Systems (CAHPS) Survey

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.

1. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: National Implementation of the In-Center Hemodialysis CAHPS Survey; *Use*: Data collected in the national implementation of the In-center Hemodialysis Consumer Assessment of Healthcare Providers and Systems (CAHPS) Survey will be used to: (1) Provide a source of information from which selected measures can be publicly reported to beneficiaries as a decision aid for dialysis facility selection, (2) aid facilities with their internal quality improvement efforts and external benchmarking with other facilities, (3) provide CMS with information for monitoring and public reporting purposes, and (4) support the end-stage renal disease value-based purchasing program. *Form Number*: CMS-10105 (OMB control number: 0938-0926). *Frequency*: Occasionally; *Affected Public*: Individuals or households; *Number of Respondents*: 109,328; *Total Annual Responses*: 109,328; *Total Annual Hours*: 59,037. (For policy questions regarding this collection contact Elizabeth Goldstein at 410-786-6665.)

2. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: Medicare Parts C and D Program Audit Protocols and Data Requests; *Use*: Under the Medicare Prescription Drug, Improvement, and