annually, as required by 5 U.S.C. 552a(o)(2)(A)(i), (r), and (u)(3)(ID).

5. Publish advance notice of the matching program in the Federal Register as required by 5 U.S.C. 552a(e)(12).

This matching program meets these requirements.

Barbara Demopulos,
Privacy Advisor, Division of Security, Privacy Policy and Governance, Office of Information Technology, Centers for Medicare & Medicaid Services.

PARTICIPATING AGENCIES:
The Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS) is the recipient agency, and the Social Security Administration (SSA) is the source agency.

AUTHORITY FOR CONDUCTING THE MATCHING PROGRAM:
The statutory authority for the matching program is 42 U.S.C. secs. 18081 and 18083.

PURPOSE(S):
The purpose of the matching program is to provide CMS with SSA information which CMS and state-based administering entities will use to determine individuals’ eligibility for initial enrollment in a Qualified Health Plan through an Exchange established under the Patient Protection and Affordable Care Act, for Insurance Affordability Programs (IAPs), and certificates of exemption from the shared responsibility payment; and to make eligibility determinations and redetermination of insurance coverage when CMS requests it.

The purpose of the matching program is to support this matching program is identified below:

(1) Master Files of SSN Holders and SSN Applications, 60–0058, last fully published at 75 FR 82121 (Dec. 29, 2010) and amended at 78 FR 40542 (July 5, 2013), 79 FR 8780 (Feb. 13, 2014), 83 FR 31250 (July 3, 2018), and 83 FR 54969 (Nov. 1, 2018);

(2) Prisoner Update Processing System (PUPS), 60–0269, last fully published at 64 FR 11076 (Mar. 8, 1999) and amended at 72 FR 69723 (Dec. 10, 2007), 78 FR 40542 (July 5, 2013), and 83 FR 54969 (Nov. 1, 2018);

(3) Master Beneficiary Record, 60–0090, last fully published at 71 FR 1826 (Jan. 11, 2006), and amended at 72 FR 69723 (Dec. 10, 2007), 78 FR 40542 (July 5, 2013), 83 FR 31250 (July 3, 2018) and 83 FR 54969 (Nov. 1, 2018);


The SSA SORNs and routine uses that support this matching program are identified below:

B. SYSTEMS OF RECORDS MAINTAINED BY SSA

The SSA SORNs and routine uses that support this matching program are identified below:

(1) Master Files of SSN Holders and SSN Applications, 60–0058, last fully published at 75 FR 82121 (Dec. 29, 2010) and amended at 78 FR 40542 (July 5, 2013), 79 FR 8780 (Feb. 13, 2014), 83 FR 31250 (July 3, 2018), and 83 FR 54969 (Nov. 1, 2018);

(2) Prisoner Update Processing System (PUPS), 60–0269, last fully published at 64 FR 11076 (Mar. 8, 1999) and amended at 72 FR 69723 (Dec. 10, 2007), 78 FR 40542 (July 5, 2013), and 83 FR 54969 (Nov. 1, 2018);

(3) Master Beneficiary Record, 60–0090, last fully published at 71 FR 1826 (Jan. 11, 2006), and amended at 72 FR 69723 (Dec. 10, 2007), 78 FR 40542 (July 5, 2013), 83 FR 31250 (July 3, 2018) and 83 FR 54969 (Nov. 1, 2018);


DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services


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 January 19, 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,
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:

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

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–10764 Evaluation of Risk Adjustment Data Validation (RADV) Appeals and Health Insurance Exchange Outreach Training Sessions
CMS–10454 Disclosure of State Rating Requirements
CMS–R–71 Quality Improvement Organization (QIO) Assumption of Responsibilities and Supporting Regulations
CMS–370/CMS–377 ASC Forms for Medicare Program Certification
CMS–1572 Home Health Agency Survey and Deficiencies Report
CMS–10332 Disclosure Requirement for the In-Office Ancillary Services Exception

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: New collection (Request for a new OMB control number); Title of Information Collection: Evaluation of Risk Adjustment Data Validation (RADV) Appeals and Health Insurance Exchange Outreach Training Sessions; Use: CMS recognizes that the success of accurately identifying risk-adjustment payments and payment errors is dependent upon the data submitted by Medicare Advantage Organizations (MAOs), and is strongly committed to providing appropriate education and technical outreach to MAOs and third-party administrators (TPAs). In addition, CMS strongly committed to providing appropriate education and technical outreach to States, issuers, self-insured group health plans and TPAs participating in the Marketplace and/or market stabilization programs mandated by the Affordable Care Act (ACA).
CMS will strengthen outreach and engagement with MAOs and stakeholders in the Marketplace through satisfaction surveys following contract-level (CON) RADV audit and Health Insurance Exchange training events. The survey results will help to determine stakeholders’ level of satisfaction with trainings, identify any issues with training and technical assistance delivery, clarify stakeholders’ needs and preferences, and define best practices for training and technical assistance.

Form Number: CMS–10764 (OMB control number: 0938–NEW); Frequency: Occasionally; Affected Public: Private Sector; Number of Respondents: 4,270; Total Annual Responses: 4,270; Total Annual Hours: 1,068.

(For questions regarding this collection contact Melissa Barkai at 410–786–4305.)

2. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Disclosure of State Rating Requirements; Use: The final rule “Patient Protection and Affordable Care Act; Health Insurance Market Rules; Rate Review” implements sections 2701, 2702, and 2703 of the Public Health Service Act (PHS Act), as added and amended by the Affordable Care Act, and sections 1302(e) and 1312(c) of the Affordable Care Act. The rule directs that states submit to CMS certain information about state rating and risk pooling requirements for their individual, small group, and large group markets, as applicable. Specifically, states will inform CMS of age rating ratios that are narrower than 3:1 for adults; tobacco use rating ratios that are narrower than 1.5:1; a state-established uniform age curve; geographic rating areas; whether premiums in the small and large group market are required to be based on average enrollee amounts (also known as composite premiums); and, in states that do not permit any rating variation based on age or tobacco use, uniform family tier structures and corresponding multipliers. In addition, states that elect to merge their individual and small group market risk pools into a combined pool will notify CMS of such election. This information will allow CMS to determine whether state-specific rules apply or Federal default rules apply. It will also support the accuracy of the federal risk adjustment methodology.

Form Number: CMS–10454 (OMB control number 0938–1258); Frequency: Occasionally; Affected Public: State, Local, or Tribal Governments; Number of Respondents: 3; Total Annual Responses: 3; Total Annual Hours: 17.

(For policy questions regarding this collection contact Russell Tipps at 301–869–3502.)

3. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Quality Improvement Organization (QIO) Assumption of Responsibilities and Supporting Regulations; Use: The Peer Review Improvement Act of 1982 amended Title XI of the Social Security Act to create the Utilization and Quality Control Peer Review Organization (PRO) program which replaces the Professional Standards Review Organization (PSRO) program and streamlines peer review activities. The term PRO has been renamed Quality Improvement Organization (QIO). This information collection describes the review functions to be performed by the QIO. It outlines relationships among QIOs, providers, practitioners, beneficiaries, intermediaries, and carriers.

Form Number: CMS–R–71 (OMB control number: 0938–0445); Frequency: Yearly; Affected Public: Business or other for-profit and Not-for-profit institutions; Number of Respondents: 6,939; Total Annual Responses: 972,478; Total Annual Hours: 1,034,655.

(For policy questions regarding this collection contact Kimberly Harris at 401–837–1118.)

4. Type of Information Collection Request: Extension of a currently approved collection; Titles of Information Collection: ASC Forms for Medicare Program Certification: Use: the form CMS–370 titled “Health Insurance Benefits Agreement” is used for the purpose of establishing an ASC’s eligibility for payment under Title XVIII of the Social Security Act (the “Act”). This agreement, upon acceptance by the Secretary of Health & Human Services, shall be binding on the ASC and the Secretary. The agreement may be
the ACA amended section 1877(b)(2) of the 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.

Physicians who provide certain imaging services (MRI, CT, and PET) under the in-office ancillary services exception 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: Private Sector—Business or other for-profit and Not-for-profit institutions; Number of Respondents: 1,567; Total Annual Responses: 1,567; Total Annual Hours: 1,012. (For policy questions regarding this collection contact Caroline Gallaher at 410–786–8705.)

5. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: Home Health Agency Survey and Deficiencies Report; Use: In order to participate in the Medicare Program as a Home Health Agency (HHA) provider, the HHA must meet federal standards. This form is used to record information and patients’ health and provider compliance with requirements and to report the information to the federal government. Form Number: CMS–1572 (OMB control number: 0938–0355); Frequency: Yearly; Affected Public: State, Local or Tribal Government; Number of Respondents: 3,833; Total Annual Responses: 3,833; Total Annual Hours: 1,917. (For policy questions regarding this collection contact Tara Lemons at 410–786–3030.)

6. 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 Affordable Care Act (ACA) established a new disclosure requirement that a physician must perform for certain imaging services to meet the in-office ancillary services exception to the prohibition of the physician self-referral law. This section

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity: Addition of New Instruments to Existing Information Collections by the Office of Refugee Resettlement (OMB #s: 0970–0553, 0970–0554, and 0970–0547)

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, Department of Health and Human Services.

ACTION: Request for public comment.

SUMMARY: The Office of Refugee Resettlement (ORR), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is inviting public comments on several proposed instrument efforts. The instruments will be added to the following existing information collections: Services Provided to Unaccompanied Alien Children (OMB #0970–0553), Placement and Transfer of Unaccompanied Alien Children into ORR Care Provider Facilities (OMB #0970–0554), and Administration and Oversight of the Unaccompanied Alien Children Program (OMB #0970–0547).

DATES: Comments due within 60 days of publication. In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described in this notice.

ADDRESSES: Copies of the proposed collection of information can be obtained and comments may be forwarded by emailing infocollection@acf.hhs.gov. Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation (OPRE), 330 C Street SW, Washington, DC 20201, Attn: ACF Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: The components of these requests and the existing information collections to which each component will be added are as follows:

Services Provided to Unaccompanied Alien Children Into ORR Care Provider Facilities (OMB #0970–0553)

1. Admission: This instrument is used by ORR grantee case managers and clinicians to document the UAC’s initial needs, functioning, and history. The Admission Details tab includes a case status timeline; biographic information on the UAC; admission and educational information; medical clearance information; influx transfer information, if applicable; system-generated information; a clickable, auto-generated list of Admission Assessments and the ability to create a new assessment; a clickable, auto-generated list of Transfer Requests and the ability to create a new transfer requests, if applicable; and a clickable, auto-generated list of Long Term Foster Care (LFTC) Travel Requests and the ability to create a new transfer requests, if applicable. The Related tab includes areas to upload case management, education, and medical documents; an area to add Entry Team members (individuals granted read/write access to the Admission instrument); and an auto-generated list of changes made to the