

- c. The number of contractor direct labor hours expended on the services performed during the previous Government fiscal year; and
- d. Data reported by subcontractors.

The prime contractor shall require each first-tier subcontractor performing under the contract to provide annually—

- a. The subcontract number (including subcontractor name and if available, Unique Entity Identifier number; and
- b. The number of first-tier subcontractor direct-labor hours expended on the services performed during the previous Government fiscal year.

In order to invoice the government for time-and-material/labor-hour (T&M/LH) and cost-reimbursement contracts, contractors already track labor hours expended, so the rule will cover T&M/LH and cost-reimbursement contracts over the simplified acquisition threshold.

Fixed price contracts are covered if the estimated total value is at \$500,000 or more in FY 2016 and thereafter.

For indefinite-delivery contracts, including but not limited to, indefinite-delivery indefinite-quantity (IDIQ) contracts, Federal Supply Schedule (FSS) contracts, Governmentwide Acquisition contracts (GWACs), and multi-agency contracts, reporting requirements will be determined based on the expected dollar amount and type of the orders issued under the contracts.

The burden has increased from the one in **Federal Register** Notice 78 FR 16268 dated March 14, 2013 due to more respondents being included in the overall total based on FY 2016 FPDS data. The threshold for Fixed-price contract reports are now covered if the estimated total value is at \$500,000 or more.

B. Annual Reporting Burden

Respondents: 111,172.

Responses/respondent: 1.

Total annual responses: 111,172.

Preparation hours per response: 2.

Total response burden hours: 222,344.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the FAR, and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of

information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

Obtaining copies of proposals:

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0179, Service Contracts Reporting Requirements, in all correspondence.

Dated: May 22, 2017.

Lorin S. Curit,

*Director, Federal Acquisition Policy Division,
Office of Governmentwide Acquisition Policy,
Office of Acquisition Policy, Office of
Governmentwide Policy.*

[FR Doc. 2017–10849 Filed 5–25–17; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10454, CMS–10558 and CMS–10650]

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 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 July 25, 2017.

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' 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: 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–10454 Disclosure of State Rating Requirements

CMS–10558 Information Collection for Machine Readable Data for Provider Network and Prescription Formulary Content for FFM QHPs

CMS–10650 State Permissions for Enrollment in Qualified Health Plans in the Federally Facilitated Exchange & Non-Exchange Entities

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:* 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:* On Occasion; *Affected Public:* State, Local, or Tribal Governments, Private Sector; *Number of Respondents:* 47; *Number of Responses:* 47; *Total Annual Hours:* 2,239. (For policy questions regarding this collection, contact Russell Tipps at 301-492-4371.)

2. *Type of Information Collection Request:* Revision of a currently approved information collection request; *Title of Information Collection:* Information Collection for Machine Readable Data for Provider Network and Prescription Formulary Content for FFM QHPs; *Use:* Under 45 CFR 156.122(d)(1)(2) and 156.230(c) and in the final rule, *Patient Protection and Affordable Care Act; HHS Notice of Benefit and Payment Parameters for 2018* (CMS-9934-F), standards for qualified health plan (QHP) issuers are established for the submission of provider and formulary data in a machine-readable format to the Department of Health and Human Services (HHS) and for posting on issuer Web sites. These standards provide greater transparency for consumers, including by allowing software developers to access formulary and provider data to create innovative and informative tools. *Form Number:* CMS-10558 (OMB Control Number 0938-1284); *Frequency:* Annually; *Affected Public:* Private Sector, State, Business, and Not-for Profits; *Number of Respondents:* 397; *Number of Responses:* 397; *Total Hours:* 208. (For questions regarding this collection contact Joshua Annas at (301) 492-4407.)

3. *Type of Information Collection Request:* New collection of information request; *Title of Information Collection:* State Permissions for Enrollment in Qualified Health Plans in the Federally Facilitated Exchange & Non-Exchange Entities; *Use:* The Patient Protection and Affordable Care Act, Public Law 111-148, enacted on March 23, 2010, and the Health Care and Education Reconciliation Act, Public Law 111-152, enacted on March 30, 2010 (collectively, “Affordable Care Act”), expand access to health insurance for individuals and employees of small businesses through the establishment of new Affordable Insurance Exchanges (Exchanges), also called Marketplaces, including the Small Business Health Options Program (SHOP). The Exchanges, which became operational on January 1, 2014, enhance competition in the health insurance market, expand access to affordable health insurance for millions of Americans, and provide consumers with a place to easily compare and shop for health insurance coverage.

This Information Collection Request (ICR) serves as the formal request for a new data collection clearance associated with the HHS Notice of Benefit and Payment Parameters for 2018 Final Rule (2018 Payment Notice). This ICR includes data collections related to the

ability of states to permit agents and brokers to assist qualified individuals, qualified employers, or qualified employees enrolling in Qualified Health Plans in the Federally Facilitated Exchange (§ 155.220) and ICRs related to non-exchange entities (§ 155.260). *Form Number:* CMS-10650 (OMB Control Number 0939-NEW); *Frequency:* Annually; *Affected Public:* Private Sector, State, Business, and Not-for Profits; *Number of Respondents:* 107,207; *Number of Responses:* 107,207; *Total Annual Hours:* 512,141. (For questions regarding this collection contact Joshua Annas at (301-492-4407).

Dated: May 23, 2017.

William N. Parham, III,

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

[FR Doc. 2017-10944 Filed 5-25-17; 8:45 am]

BILLING CODE 4120-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0523]

Agency Information Collection Activities: Proposed Collection; Comment Request; Applications for Food and Drug Administration Approval To Market a New Drug

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the requirements governing applications for FDA approval to market a new drug.

DATES: Submit either electronic or written comments on the collection of information by June 26, 2017. Late, untimely filed comments will not be considered. Electronic comments must be submitted on or before July 25, 2017. The <https://www.regulations.gov> electronic filing system will accept comments until midnight Eastern Time