captioning rules (47 CFR 79.1), which require that, with some exceptions, all new video programming, and 75 percent of “pre-rule” programming, be closed captioned. The existing collections include petitions by video programming providers, producers, and owners for exemptions from the closed captioning rules, responses by commenters, and replies; complaints by viewers alleging violations of the closed captioning rules, responses by video programming distributors (VPDs) and video programmers, recordkeeping in support of complaint responses, and compliance ladder obligations in the event of a pattern or trend of violations; records of monitoring and maintenance activities; caption quality best practices procedures; making video programming distributor contact information available to viewers in phone directories, on the Commission’s Web site and the Web sites of video programming distributors (if they have them), and in billing statements (to the extent video programming distributors issue them); and video programmers filing contact information and compliance certifications with the Commission.

On February 19, 2016, the Commission adopted the Closed Captioning Quality Second Report and Order, published at 81 FR 57473, August 23, 2016, amending its rules to allocate the responsibilities of VPDs and video programmers with respect to the provision and quality of closed captioning. The Commission took the following actions, among others:

(a) Required video programmers to file certifications with the Commission that (1) the video programmer (i) is in compliance with the rules requiring the inclusion of closed captions, and (ii) either is in compliance with the captioning quality standards or has adopted and is following related Best Practices; or (2) is exempt from the captioning obligation and specifies the exemption claimed.

(b) Revised the procedures for receiving, serving, and addressing television closed captioning complaints in accordance with a burden-shifting compliance model.

(c) Established a compliance ladder for the Commission’s television closed captioning quality requirements.

(d) Required VPDs to use the Commission’s web form when providing contact information to the VPD registry.

(e) Required video programmers to register their contact information with the Commission for the receipt and handling of written closed captioning complaints.

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board’s Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than November 3, 2017.

A. Federal Reserve Bank of Atlanta

Kathryn Haney, Director of Applications (1000 Peachtree Street NE., Atlanta, Georgia 30309). Comments can also be sent electronically to Applications.Comments@atl.frb.org:

1. SSX2, LLC, Tallahassee, Florida: to be added to the Smith family control group, which controls Capital City Bank Group, Inc., and its subsidiary, Capital City Bank, both of Tallahassee, Florida.
DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10054 and CMS–10106]

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 December 18, 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:


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:

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 Payments for APCs Under the Outpatient Prospective Payment System

CMS–10106 Medicare Authorization to Disclose Personal Health Information

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: Reinstatement without change of a previously approved collection; Title of Information Collection: New Technology Payments for APCs Under the Outpatient Prospective Payment System; Use: CMS needs to keep pace with emerging new technologies and make them accessible to Medicare beneficiaries in a timely manner. It is necessary that we continue to collect appropriate information from interested parties such as hospitals, medical device manufacturers, pharmaceutical companies and others that bring to our attention specific services that they wish us to evaluate for New Technology APC payment. We are making no changes to the information that we collect. The information that we seek to continue to collect is necessary to determine whether certain new services are eligible for payment in New Technology APCs, to determine appropriate coding and to set an appropriate payment rate for the new technology service. The intent of these provisions is to ensure timely beneficiary access to new and appropriate technologies Form Number: CMS–10054 (OMB control number: 0938–0860); Frequency: Annually; Affected Public: Private Sector; Business or Other for-profits; Number of Respondents: 10; Total Annual Responses: 10; Total Annual Hours: 160. (For policy questions regarding this collection contact Josh McFeeters at 410–786–9732).

2. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Medicare Authorization to Disclose Personal Health Information; Use: Unless permitted or required by law, the Health Insurance Portability and Accountability Act (HIPAA) Privacy Rule (§ 164.508) prohibits Medicare (a HIPAA covered entity) from disclosing an individual’s protected health information without a valid authorization. In order to be valid, an authorization must include specified core elements and statements. Medicare will make available to Medicare beneficiaries a standard, valid authorization to enable beneficiaries to request the disclosure of their protected health information. This standard authorization will simplify the process of requesting information disclosure for beneficiaries and minimize the response time for Medicare. Form CMS–10106 (OMB control number: 0938–0930); Frequency: Occasionally; Affected Public: