Respondents from engaging in the same or similar acts or practices in the future.

Part I of the proposed order prohibits Respondents from selling a monitoring product unless: (1) The monitoring product does not circumvent security protections implemented by the mobile device operating system or manufacturer; (2) prior to the sale of the monitoring product, express written attestation is obtained from the purchaser that the monitoring product will be used for legitimate and lawful purposes; and (3) documentation is obtained proving that the purchaser is an authorized user on the monitored mobile device’s service carrier account. The proposed order also requires that Respondents display an application icon, including the name of the monitoring product, when the monitoring product is on the mobile device. Moreover, a clear and conspicuous notice must be presented when the application icon is clicked.

Part II of the order restrains Respondents from distributing monitoring products unless Respondents have: (1) A home page notice stating that the monitoring product may only be used for legitimate and lawful purposes by authorized users; and (2) a purchase page notice stating that the monitoring product may only be used for legitimate and lawful purposes by authorized users, and that installing or using the monitoring product for any other purpose may violate local, state, and/or federal law.

Part III of the proposed order prohibits Respondents from violating the Children’s Online Privacy Protection Rule. Part IV of the proposed order prohibits Respondents from misrepresenting the extent to which Respondents maintain and protect the privacy, security, confidentiality, or integrity of consumers’ personal information. Part V requires that Respondents delete all personal information collected from a monitoring product prior to entry of the proposed order within 120 days.

Part VI of the proposed order prohibits Respondents, and any business that a Respondent controls, from selling, sharing, collecting, maintaining, or storing personal information unless Respondents establish and implement, and thereafter maintain, a comprehensive information security program that protects the security confidentiality, and integrity of such personal information. Part VII requires Respondents to obtain initial and biennial data security assessments for twenty years. Part VIII of the proposed order requires Respondents to disclose all material facts to the assessor and prohibits Respondents from misrepresenting any fact material to the assessments required by Part VII. Part IX requires Respondents to submit an annual certification from a senior corporate manager (or senior officer responsible for its information security program), that Respondents have implemented the requirements of the proposed order, are not aware of any material noncompliance that has not been corrected or disclosed to the Commission, and includes a brief description of any covered incident involving unauthorized access to or acquisition of personal information. Part X requires Respondents to submit a report to the Commission of their discovery of any covered incident.

Parts XI through XIV of the proposed order are reporting and compliance provisions, which include recordkeeping requirements and provisions requiring Respondents to provide information or documents necessary for the Commission to monitor compliance. Part XV states that the proposed order will remain in effect for 20 years, with certain exceptions. The purpose of this analysis is to aid public comment on the proposed order. It is not intended to constitute an official interpretation of the complaint or proposed order, or to modify in any way the proposed order’s terms.

By direction of the Commission.

April J. Tabor.
Acting Secretary.

[FR Doc. 2019–23809 Filed 10–30–19; 8:45 am]
BILLING CODE 6750–01–P

FEDERAL TRADE COMMISSION
Agency Information Collection Activities; Proposed Collection; Comment Request; Extension

AGENCY: Federal Trade Commission.
ACTION: Notice.

SUMMARY: The Federal Trade Commission ("FTC" or "Commission") is seeking public comment on its proposal to extend for an additional three years, the current PRA clearance for its shared enforcement authority with the Consumer Financial Protection Bureau ("CFPB") for information collection requirements contained in the CFPB’s Regulation O. That clearance expires on February 29, 2020.

DATES: Comments must be filed by December 30, 2019.

ADDRESSES: Interested parties may file a comment online or on paper, by following the instructions in the Request for Comment part of the SUPPLEMENTARY INFORMATION section below. Write “MARS (Regulation O) PRA Comment, FTC File No. P134812” on your comment, and file your comment online at https://www.regulations.gov by following the instructions on the web-based form. If you prefer to file your comment on paper, mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024.


SUPPLEMENTARY INFORMATION: Title X of the Dodd-Frank Wall Street Reform and Consumer Protection Act ("Dodd-Frank Act"), Public Law 111–203, 124 Stat. 1376 (2010), transferred the Commission’s rulemaking authority under the mortgage provisions in section 626 of the 2009 Omnibus Appropriations Act, as amended,1 to the CFPB.2 On December 16, 2011, the CFPB republished the Mortgage Assistance Relief Services (“MARS”) Rule as Regulation O (12 CFR 1015).3 As a result, the Commission subsequently rescinded its MARS Rule (16 CFR part 322).4 Nonetheless, under the Dodd-Frank Act, the FTC retains its authority to bring law enforcement actions to enforce Regulation O.5 Regulation O contains information collection requirements that have been approved by OMB under the PRA, 44 U.S.C. 3501 et seq. (OMB Control Number 3084–0157). The FTC, as a co-enforcer, seeks OMB clearance for its share of the estimated PRA burden for the information collection requirements of Regulation O. The Rule includes disclosure requirements to assist purchasers of mortgage assistance relief services in making well-informed decisions and avoiding unfair or
deceptive acts and practices. The information that must be retained under Regulation O’s recordkeeping requirements is used by the CFPB and the FTC for enforcement purposes and to ensure compliance by MARS providers with Regulation O.

**Burden Statement**

Because the FTC and CFPB share enforcement authority for this rule, the FTC is seeking clearance for one-half of the following burden estimates. These estimates are based on the agencies’ law enforcement experience and the recent analysis conducted as part of the CFPB’s clearance renewal for the information collections associated with Regulation O. The FTC and CFPB estimate that there are approximately 120 for-profit, non-attorney entities offering MARS services and subject to Regulation O’s requirements.7

Estimated annual hours burden: 360 (FTC share).

FTC staff estimates that compliance with Regulation O’s disclosure requirements for MARS providers requires 6 hours of labor annually.8 Multiplying this figure by 120 entities yields a total burden for covered providers of 720 hours annually.9 For PRA purposes, the FTC and CFPB share enforcement authority and split the information collection burden associated with the Rule equally. As a result, the FTC assumes 360 hours of this total annual hours burden.

Estimated associated labor cost: $11,747 (FTC share).

In calculating the associated labor costs, FTC staff estimates that a compliance officer or equivalent will prepare the required disclosures at an hourly rate of $32.63/hr.10 Thus, the estimated annual labor cost is $23,494 (120 providers × 6 hours × $33.26) of which the FTC assumes half, or $11,747.

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<table>
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<tr>
<th>Information collection</th>
<th>Number of respondents</th>
<th>Annual burden hours per respondent</th>
<th>Total burden hours</th>
<th>Associated hourly labor cost</th>
<th>Total respondent cost</th>
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<tr>
<td>FTC 50% Share</td>
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<td>120</td>
<td>6</td>
<td>720</td>
<td>$32.63</td>
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**Request for Comment**

Pursuant to Section 3506(c)(2)(A) of the PRA, the FTC invites comments on: (1) Whether the disclosure and recordkeeping requirements are necessary, including whether the information will be practically useful; (2) the accuracy of our burden estimates, including whether the methodology and assumptions used are valid; (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.

You can file a comment online or on paper. For the FTC to consider your comment, we must receive it on or before December 30, 2019. Write “MARS (Regulation O) PRA Comment, FTC File No. P134812” on your comment and on the envelope, and mail it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

Because your comment will be placed on the publicly accessible website at www.regulations.gov, you are solely responsible for making sure that your comment does not include any sensitive personal information, such as medical records or other individually identifiable health information. In addition, your comment should not include any “trade secret or any commercial or financial information which . . . is privileged or confidential”—as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled “Confidential,” and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c). Your comment will be kept confidential only if the General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted publicly at www.regulations.gov, we cannot redact
or remove your comment unless you submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before December 30, 2019. For information on the Commission’s privacy policy, including routine uses permitted by the Privacy Act, see https://www.ftc.gov/site-information/privacy-policy.

Heather Hippsley, Deputy General Counsel.

[FR Doc. 2019–23797 Filed 10–30–19; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–416]

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 30, 2019.

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 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–416 Annual Early and Periodic Screening, Diagnostic and Treatment (EPSDT) Participation Report

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: Annual Early and Periodic Screening, Diagnostic and Treatment (EPSDT) Participation Report; Use: The collected baseline data is used to assess the effectiveness of state early and periodic screening, diagnostic and treatment (EPSDT) programs in reaching eligible children (by age group and basis of Medicaid eligibility) who are provided initial and periodic child health screening services, referred for corrective treatment, and receiving dental, hearing, and vision services. This assessment is coupled with the state’s results in attaining the participation goals set for the state. The information gathered from this report, permits federal and state managers to evaluate the effectiveness of the EPSDT law on the basic aspects of the program. Form Number: CMS–416 (OMB control number 0938–0354); Frequency: Yearly and on occasion; Affected Public: State, Local, or Tribal Governments; Number of Respondents: 56; Total Annual Responses: 56; Total Annual Hours: 1,512. (For policy questions regarding this collection contact Karen Matsuoka at 410–786–9726.)

Dated: October 25, 2019.

William N. Parham, III, Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–23773 Filed 10–30–19; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Meeting of the National Advisory Council on Nurse Education and Practice

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In accordance with the Federal Advisory Committee Act, this