$28,785 for reporting + $46,080 for disclosures).

Total annual capital or other non-labor costs: $314,000, rounded to the nearest thousand.

Total capital and start-up costs: The Rule imposes no appreciable current capital or start-up costs. The vast majority of warrantors have already developed systems to retain the records and provide the disclosures required by the Rule. Rule compliance does not require the use of any capital goods, other than ordinary office equipment, to which providers already have access.

The Rule imposes only one additional cost on IDSMs operating under the Rule that would not apply to other IDSMs: The annual audit requirement. According to representatives of the IDSMs, the vast majority of costs associated with this requirement consist of the fees paid to the auditors and their staffs to perform the annual audit. Representatives of the IDSMs previously estimated a combined cost of $300,000 for both IDSMs currently operating under the Rule. Staff retains that estimate.

Other non-labor costs: $13,707 in copying costs, based on estimated copying costs of 7 cents per page and several conservative assumptions. Staff estimates that the average dispute-related file contains 35 pages and a typical annual audit file contains approximately 200 pages. As discussed above, staff assumes that twenty percent of consumers using an IDSM currently operating under the Rule (approximately 2,303 consumers) request copies of the records relating to their disputes.

Staff also estimates that a very small minority of consumers request a copy of the annual audit. Staff bases this assumption on (1) the number of consumer requests received by the IDSMs in the past; and (2) the fact that the IDSMs' annual audits are available online. For example, annual audits are available on the FTC's Web site, where consumers may view and or print pages as needed, at no cost to the IDSM. In addition, the Better Business Bureau makes available on its Web site the annual audit of the BBB AUTO LINE. Therefore, staff conservatively estimates that only five percent of consumers using an IDSM covered by the Rule (approximately 576 consumers) will request a copy of the IDSM's audit report.

Thus, the total annual copying cost for dispute-related files is approximately $5,643 (35 pages per file × $.07 per page × 2,303 consumer requests) and the total annual copying cost for annual audit reports is approximately $8,064 (200 pages per audit report × $.07 per page × 576 consumer requests). Accordingly, the total cost attributed to copying under the Rule is approximately $13,707. Thus, the total non-labor cost under the Rule is approximately $314,000 ($300,000 for auditor fees + $13,707 for copying costs).

Request for Comments

You can file a comment online or on paper. Write ''Warranty Rules: Paperwork Comment, FTC File No. P044403'' on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission Web site, at *http://www.ftc.gov/os/publiccomments.shtm.* As a matter of discretion, the Commission tries to remove individuals' home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, like anyone's Social Security number, date of birth, driver's license number or other state identification number or foreign country equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any ''[t]rade secret or any commercial or financial information which is * * * privileged or confidential,'' as discussed in 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). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you must follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c).[14] Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, the Commission encourages you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at *https://ftcpublic.commentworks.com/ftc/idsrpra* by following the instructions on the web-based form. If this Notice appears at *http://www.regulations.gov,* you also may file a comment through that Web site.

If you file your comment on paper, write ''Warranty Rules: Paperwork Comment, FTC File No. P044403'' on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H–113 (Annex J), 600 Pennsylvania Avenue NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at *http://www.ftc.gov* to read this Notice. 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 February 10, 2014. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

David C. Shonka,

Principal Deputy General Counsel.

[FR Doc. 2013–29404 Filed 12–9–13; 8:45 am]

BILLING CODE 6750–01–P

[14] 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), 16 CFR 4.9(c).

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; National Survey of Older Americans Act Participants

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. 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 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 information collection requirements relating to the information collection requirements contained in consumer assessment surveys that are used by ACL to measure program performance for programs funded under Title III of the Older Americans Act.

DATES: Submit written or electronic comments on the collection of information by February 10, 2014.

ADDRESSES: Submit electronic comments on the collection of information to: *elena.fazio@acl.hhs.gov.*

Submit written comments on the collection of information to Elena Fazio, Administration for Community Living, Office of Performance and Evaluation, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Elena Fazio, 202–357–3583.

SUPPLEMENTARY 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. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency request 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 (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility; (2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (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 on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The National Survey of Older Americans Act (OAA) Participants information collection, which builds on earlier national pilot studies and surveys, as well as performance measurement tools developed by ACL grantees in the Performance Outcomes Measures Project (POMP), will include consumer assessment surveys for the Congregate and Home-delivered meal nutrition programs; Case Management, Homemaker, and Transportation Services; and the National Family Caregiver Support Program. This information will be used by ACL to track performance outcome measures; support budget requests; comply with GPRA Modernization Act of 2010 (GPRAMA) reporting requirements; provide national benchmark information; and inform program development and management initiatives. Descriptions of previous National Surveys of OAA Participants can be found under the section on OAA Performance Information on ACL's Web site at: *http://www.aoa.gov/AoARoot/Program_Results/OAA_Performance.aspx.* Copies of the survey instruments and data from previous National Surveys of OAA Participants can be found and queried using the AGing Integrated Database (AGID) at *http://www.agid.acl.gov/.* The proposed Ninth National Survey entitled National Survey of OAA Participants draft 2013 may be found on the ACL Web site at *http://www.aoa.gov/AoARoot/Program_Results/OAA_Performance.aspx.* ACL estimates the burden of this collection of information as follows: *Respondents:* Individuals; *Number of Respondents:* 6,250; *Number of Responses per Respondent:* one; *Average Burden per Response:* 6000 at 40 minutes, 250 at 4 hours; *Total Burden:* 5,000.

Dated: December 5, 2013.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2013–29436 Filed 12–9–13; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0853]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Medical Devices Current Good Manufacturing Practice Quality System Regulation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Fax written comments on the collection of information by January 9, 2014.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0073. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Medical Devices Current Good Manufacturing Practice Quality System Regulation—21 CFR Part 820 (OMB Control Number 0910–0073)—Extension

Under section 520(f) of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 360j(f)), the Secretary of the Department of Health and Human Services has the authority to prescribe regulations requiring that the methods used in, and the facilities and controls used for, the manufacture, preproduction design validation (including a process to assess the performance of a device but not including an evaluation of the safety