withdraw from the agreement or make final the agreement's proposed order.

AmeriFreight is an automobile shipment broker—that is, it arranges shipment of consumers' automobiles through third-party freight carriers. This matter involves AmeriFreight's online advertising for those services. The Commission's complaint alleges that the Respondents violated Section 5(a) of the Federal Trade Commission Act by misrepresenting that AmeriFreight was a highly rated or top-ranked automobile shipment broker based on its customers' unbiased reviews. The complaint also alleges that AmeriFreight failed to disclose that it paid consumers to post reviews.

The proposed order includes injunctive relief that prohibits these alleged violations and fences in similar and related violations.

Part I of the Order prohibits the Respondents from misrepresenting that their products or services are highly rated or top-ranked based on unbiased customer reviews or that their customer reviews are unbiased.

Part II of the Order requires the Respondents, when using an endorsement to advertise any product or service, to clearly and prominently disclose a material connection, if one exists, between the person providing the endorsement and Respondents.

Part III contains recordkeeping requirements for advertisements and other documents relevant to the order.

Parts IV through VII of the proposed order require Respondents to: Deliver a copy of the order to principals, officers, directors, managers, employees, agents, and representatives having responsibilities with respect to the subject matter of the order; notify the Commission of changes in corporate structure, discontinuance of current business or employment, or affiliation with any new business or employment that might affect compliance obligations under the order; and file compliance reports with the Commission.

Part VIII provides that the order will terminate after twenty (20) years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the proposed order, and it is not intended to constitute an official interpretation of the complaint or proposed order, or to modify the proposed order's terms in any way.

By direction of the Commission.

**Donald S. Clark,**

*Secretary.*

[FR Doc. 2015–05105 Filed 3–5–15; 8:45 am]

**BILLING CODE 6750–01–P**

## FEDERAL TRADE COMMISSION

### Agency Information Collection Activities; Submission for OMB Review; Comment Request; Extension

**AGENCY:** Federal Trade Commission (''FTC'' or ''Commission'').

**ACTION:** Notice.

**SUMMARY:** The information collection requirements described below will be submitted to the Office of Management and Budget (''OMB'') for review, as required by the Paperwork Reduction Act (''PRA''). The FTC intends to ask OMB to extend for an additional three years the current PRA clearance for the FTC's enforcement of the information collection requirements in its Fair Packaging and Labeling Act regulations (''FPLA Rules''). That clearance expires on May 31, 2015.

**DATES:** Comments must be filed by April 6, 2015.

**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 ''FPLA Rules, PRA Comment, P074200'' on your comment and file your comment online at *https://ftcpublic.commentworks.com/ftc/fplaregspra2* 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.

**FOR FURTHER INFORMATION CONTACT:** Megan Gray, Attorney, Division of Enforcement, Bureau of Consumer Protection, (202) 326–3405, 600 Pennsylvania Ave. NW., Room 9541, Washington, DC 20580.

**SUPPLEMENTARY INFORMATION:** On December 16, 2014, the FTC sought public comment on the information collection requirements associated with the FPLA Rules (December 16, 2014 Notice[1]), 16 CFR parts 500–503 (OMB Control Number 3084–0110).[2] No relevant comments were received. Pursuant to the OMB regulations, 5 CFR part 1320, that implement the PRA, 44 U.S.C. 3501 *et seq.,* the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for the Rule. All comments should be filed as prescribed herein, and must be received on or before April 6, 2015.

#### Burden Statement

As detailed in the December 16, 2014 Notice, the FTC estimates cumulative annual burden on affected entities to be 8,015,140 hours and $185,149,734 in labor costs. Commission staff believes that the FPLA Rules impose negligible capital or other non-labor costs, as the affected entities are likely to have the necessary supplies and/or equipment already (*e.g.,* offices and computers) to implement the packaging and labeling disclosure requirements under the FPLA Rules.

#### Request for Comment

You can file a comment online or on paper. For the FTC to consider your comment, we must receive it on or before April 6, 2015.

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before April 6, 2015. Write ''FPLA Rules, PRA Comment, P074200'' 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 doesn't 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 doesn't 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

[1] 79 FR 74722.

[2] Section 4 of the FPLA specifically requires packages or labels to be marked with: (1) A statement of identity; (2) a net quantity of contents disclosure; and (3) the name and place of business of the company responsible for the product. The FPLA Rules, 16 CFR parts 500–503, specify how manufacturers, packagers, and distributors of ''consumer commodities'' must do this.

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, don't include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to treat your comment as confidential, you must file it in paper form, with a request for confidentiality, and you have to follow the procedure explained in FTC Rule 4.9(c).[3] Your comment will be kept confidential only if the FTC General Counsel 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, we encourage 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/fplaregspra2,* by following the instructions on the web-based form. When this Notice appears at *http://www.regulations.gov/#!home,* you also may file a comment through that Web site.

If you file your comment on paper, write ''FPLA Rules, PRA Comment, P074200'' 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.

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 April 6, 2015. For information on the Commission's privacy policy, including routine uses permitted by the Privacy Act, see *http://www.ftc.gov/ftc/privacy.htm.* For supporting documentation and other information underlying the PRA discussion in this Notice, see *http://www.reginfo.gov/public/jsp/PRA/praDashboard.jsp.*

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW., Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead should be sent by facsimile to (202) 395–5806.

**David C. Shonka,**

*Principal Deputy General Counsel.*

[FR Doc. 2015–05194 Filed 3–5–15; 8:45 am]

**BILLING CODE 6750–01–P**

[3] 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

### Centers for Disease Control and Prevention

**[30-Day–15–14LA]**

### Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies' estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

**Proposed Project**

Annual Survey of Colorectal Cancer Control Activities Conducted by States and Tribal Organizations—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

In July 2009, the Centers for Disease Control and Prevention's (CDC's) Division of Cancer Prevention and Control, National Center for Chronic Disease Prevention and Health Promotion, funded the Colorectal Cancer Control Program (CRCCP) for a 5-year period. The purpose of the CRCCP is to promote colorectal cancer (CRC) screening to increase population-level screening rates to 80% and, subsequently, to reduce CRC incidence and mortality. The current awardees are 25 states and 4 tribal organizations.

The CRCCP includes two program components: (1) CRC screening of low-income, uninsured and underinsured people (screening provision) and (2) implementation of interventions to increase population-level screening rates (screening promotion).

As a comprehensive, organized screening program, the CRCCP supports activities including program management, partnership development, public education and targeted outreach, screening and diagnostic services, patient navigation, quality assurance and quality improvement, professional development, data management and utilization, and program monitoring and evaluation. For clinical service delivery, grantees fund health care providers in their state/territory/tribe to deliver colorectal cancer screening, diagnostic evaluation, and treatment referrals for those diagnosed with cancer.

An annual survey of CRCCP grantees was fielded from 2011–2013 through the