uses permitted by the Privacy Act, in the Commission's privacy policy, at *http://www.ftc.gov/ftc/privacy.htm.*

Analysis of Proposed Consent Order To Aid Public Comment

The Federal Trade Commission (''FTC'' or ''Commission'') has accepted, subject to final approval, an agreement containing a consent order from Made in the USA Brand, LLC. (''Respondent'').

The proposed consent order has been placed on the public record for thirty (30) days for receipt of comments by interested persons. Comments received during this period will become part of the public record. After thirty (30) days, the Commission will again review the agreement and the comments received, and will decide whether it should withdraw from the agreement or make final the agreement's proposed order.

This matter involves Respondent's marketing, sale, and distribution of licenses to use its ''Made in USA'' certification mark to companies wishing to make U.S.-origin claims for their products. According to the FTC's complaint, Respondent represented that products and entities using Respondent's certification mark were independently and objectively evaluated for compliance with Respondent's accreditation standard. These claims were false or misleading. Additionally, the complaint alleges that Respondent did not possess and rely upon a reasonable basis to substantiate its claims that entities promoted on its Web site sold products that are all or virtually all made in the United States. In fact, in numerous instances, entities promoted on Respondent's Web site have sold products containing significant imported content. Finally, the complaint alleges that Respondent distributed promotional materials to third-party marketers for use in the marketing and sale of those third parties' products, providing the means and instrumentalities to those marketers to commit deceptive acts or practices. Accordingly, the complaint concludes that Respondent engaged in deceptive acts or practices in violation of Section 5(a) of the FTC Act.

The proposed consent order contains provisions designed to prevent Respondent from engaging in similar acts and practices in the future. Specifically, Part I prohibits Respondent from representing, expressly or by implication, that covered entities meet Respondent's accreditation standard, unless: (1) An entity with no material connection to that covered entity conducted an independent and objective evaluation to confirm that the accreditation standard was met; or (2) Respondent's mark and marketing materials prominently disclose that the accreditation standard may be met through self-certification.

Part II prohibits Respondent from making any country of origin claim about a product authorized to use Respondent's certification mark unless: (1) The claim is true, not misleading, and Respondent has a reasonable basis substantiating the representation; or (2) for representations made through use of Respondent's certification mark, Respondents clearly and prominently disclose that covered entities may meet the accreditation standard through self-certification.

Part III prohibits Respondent from providing third-party retailers with the means and instrumentalities to make the claims prohibited in Part I.

Parts IV through VIII are reporting and compliance provisions. Part IV requires Respondent to keep and make available to the Commission on request: Copies of advertisements, labeling, packaging, and promotional materials containing the representations identified in Parts I and II; materials relied upon in disseminating those representations; evidence that contradicts, qualifies, or calls into question the representations or the basis relied upon for the representations; and all acknowledgments of receipt of the Order. Part V requires Respondent to disseminate the Order to principals, officers, directors, and managers, and to all current and future employees, agents, and representatives having responsibilities relating to the subject matter of the order. Part VI requires notification to the FTC of changes in Respondent's corporate status. Part VII requires Respondent to submit an initial compliance report to the FTC within sixty (60) days of service and subsequent reports upon request.

Finally, Part VIII is a ''sunset'' provision, terminating the order after twenty (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 proposed order or to modify its terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2014–17705 Filed 7–25–14; 8:45 am]

BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-14–13AAI]

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

ROPS Attributes Identified by Distribution Channel Intermediaries—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all people through research and prevention.

High rates of traumatic injury are associated with the use of older tractors that are not equipped with rollover protective structures (ROPS), which have been proven to reduce tractor-rollovers, a leading cause of injury to agricultural workers. To reduce the incidence of traumatic injury among farm workers, NIOSH proposes to administer stated-preference questionnaires designed to assess preference among a group of tractor-parts dealers in Pennsylvania, New York, New Hampshire and Vermont, who have membership in the Northeast Equipment Dealers' Association (NEDA). NEDA is a trade group for tractor parts dealers and is active in 12 States in the Northeast and Mid-Atlantic States. This information will be used to assess the impediments and barriers to adoption, as well as the incentives, for the distribution and sale of ROPS.

ROPS are generally provided to end users by tractor parts dealers, who constitute distribution channel intermediaries between the manufacturer and the consumer. However, little is known about the decision processes that tractor parts dealers follow in deciding whether or not to provide ROPS to end users. The current project will generate ranking scores for the importance given to various items of concern to tractor parts dealers; these most-important items were previously developed through review of relevant research studies.

CDC proposes to collect customized information, from 520 NEDA establishments, over a one-month period. This information will be of three kinds: 1. General screening information as to the appropriateness of administering a survey to the respondent organization; 2. Limited respondent perception of the demographic characteristics on the client base served by the NEDA establishment, and 3. Importance ranking of attributes of the process of providing ROPS, or the ROPS configuration itself.

This information will allow CDC to compile a systematic, quantifiable inventory of preference data for a group that is considered representative of tractor parts dealers nationwide. It will also allow CDC to develop recommendations for overcoming the barriers that have compromised the effectiveness of occupational health and safety programs.

The total estimated burden for the one-time retrospective data collection is 43 hours as indicated in the table below. The average burden per response is 5 minutes. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs)
Tractor Parts Dealers	ROPS Questionnaire for Tractor Parts Dealers	520	1	5/60

Leroy Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2014–17651 Filed 7–25–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–3296–NC]

RIN 0938–ZB14

Medicare Program; Evaluation Criteria and Standards for Beneficiary and Family Centered Care Quality Improvement Organization Contract

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice with comment period.

SUMMARY: This notice with comment period describes the general criteria we intend to use to evaluate the effectiveness and efficiency of Beneficiary and Family Centered Care (BFCC) Quality Improvement Organizations (QIOs) that will enter into contracts with CMS under the 11th Statement of Work (SOW) in May 2014 titled, ''Beneficiary and Family Centered Care (BFCC) Quality Improvement Organization (QIO) Contract'' (HHSM–500–2014–RFP–BFCC–QIO). This contract allows for a transition period from the incumbent QIOs to the successor QIOs. The activities for the BFCC–QIO SOW begin August 1, 2014. The evaluation of a BFCC–QIO's performance related to the SOW will be based on evaluation criteria specified for the tasks set forth in Attachment J–10 of the BFCC–QIOs' SOW contract.

DATES: *Effective Date:* August 1, 2014 to July 31, 2019.

Comment Date: To be assured consideration, comments must be received at one of the addresses provided below, no later than 5 p.m. on August 27, 2014.

ADDRESSES: In commenting, refer to file code CMS–3296–NC. Because of staff and resource limitations, we cannot accept comments by facsimile (FAX) transmission.

You may submit comments in one of four ways (please choose only one of the ways listed):

1. *Electronically.* You may submit electronic comments on this regulation to *http://www.regulations.gov.* Follow the ''Submit a comment'' instructions.

2. *By regular mail.* You may mail written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–3296–NC, P.O. Box 8010, Baltimore, MD 21244–1850.

Please allow sufficient time for mailed comments to be received before the close of the comment period.

3. *By express or overnight mail.* You may send written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS–3296–NC, Mail Stop C4–26–05, 7500 Security Boulevard, Baltimore, MD 21244–1850.

4. *By hand or courier.* Alternatively, you may deliver (by hand or courier) your written comments ONLY to the following addresses:

a. For delivery in Washington, DC—Centers for Medicare & Medicaid Services, Department of Health and Human Services, Room 445–G, Hubert H. Humphrey Building, 200 Independence Avenue SW., Washington, DC 20201

(Because access to the interior of the Hubert H. Humphrey Building is not readily