

Washington, DC 20427. Please note that at this time, mail is sometimes delayed. Therefore, we encourage emailed comments.

FOR FURTHER INFORMATION CONTACT: Sarah Cudahy, 202–606–8090, register@fmcs.gov.

SUPPLEMENTARY INFORMATION: Copies of the proposed questions are available below. Paper copies are available by emailing register@fmcs.gov. Please ask for the Stakeholder Survey.

I. Information Collection Request

Agency: Federal Mediation and Conciliation Service.

Form Number: Not yet assigned.

Type of Request: New collection; generic clearance.

Affected Entities: Private sector; state, local, and tribal governments; individuals or households; and federal government.

Frequency: These methods of engagement are utilized on an as-needed basis. Each engagement is completed once.

Abstract: Pursuant to the Administrative Dispute Resolution Acts of 1990 and 1996, 5 U.S.C. 561 *et seq.* and 571 *et seq.*, and 29 U.S.C. 173(f), the Federal Mediation and Conciliation Service provides conflict prevention, management, and resolution services, including, but not limited to, public policy facilitation and mediation services, to Federal agencies. As part of these services, sometimes FMCS employees need to survey or ask questions to determine the best process and participants to prevent, manage, or resolve the issue, particularly for public policy mediations, public policy or environmental facilitations, or negotiated rulemaking. To do so, FMCS has created a set of questions to ask various stakeholders about issues, concerns, engagement, and appropriate stakeholders relevant to the issues. The survey format will differ depending on the project but may be conducted in one or more of the following ways, both in-person and virtually: Individual or group interviews, individual or group discussions, or written surveys. The survey requests information such as stakeholder understanding of the particular issue, stakeholder interests in the particular issue, appropriate stakeholders, methods of engagement with the issue, and other similar information that will allow FMCS to best create a successful process. A link to the survey is found here: https://tags.fmcs.gov/4DAction/FC/DoAsynchTop?Fedreg*UPPJ*919/10300. To log in, go to: <https://tags.fmcs.gov/>, use username “Fedreg” and password

“UPPJ.” The collection of such information is critical for ensuring the appropriate process, stakeholders, and stakeholder input in the process. No other collections are being conducted that would provide this information to FMCS.

Burden: The current total annual burden estimate is that FMCS will receive information from approximately 15,000 respondents per year. Interviews and discussions would be approximately thirty minutes in duration. Written surveys would take approximately ten minutes to complete. FMCS expects the total burden to not exceed 2,535 hours per year.

II. Request for Comments

FMCS solicits comments to:

i. Evaluate whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information will have practical utility.

ii. Enhance the accuracy of the agency’s estimates of the burden of the proposed collection of information.

iii. Enhance the quality, utility, and clarity of the information to be collected.

iv. Minimize the burden of the collections of information on those who are to respond, including the use of appropriate automated, electronic collection technologies or other forms of information technology.

III. 60-Day Comment Period

This information was previously published in the **Federal Register** on March 16, 2022, allowing for a 60-day public comment period under Document 2022–05543 at 87 FR 14857. FMCS received no comments.

IV. The Official Record

The official records are electronic records.

List of Subjects

Information Collection Requests.

Dated: May 13, 2022.

Anna Davis,
Acting General Counsel.

[FR Doc. 2022–10752 Filed 5–18–22; 8:45 am]

BILLING CODE 6732–01–P

FEDERAL TRADE COMMISSION

[File No. 222 3023]

Lions Not Sheep; Analysis of Proposed Consent Order To Aid Public Comment

AGENCY: Federal Trade Commission.

ACTION: Proposed consent agreement; request for comment.

SUMMARY: The consent agreement in this matter settles alleged violations of federal law prohibiting unfair or deceptive acts or practices. The attached Analysis of Proposed Consent Order to Aid Public Comment describes both the allegations in the draft complaint and the terms of the consent order—embodied in the consent agreement—that would settle these allegations.

DATES: Comments must be received on or before June 21, 2022.

ADDRESSES: Interested parties may file comments online or on paper by following the instructions in the Request for Comment part of the **SUPPLEMENTARY INFORMATION** section below. Please write “Lions Not Sheep; File No. 222 3023” 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 D), 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 D), Washington, DC 20024.

FOR FURTHER INFORMATION CONTACT: Julia Solomon Ensor (202–326–2377), Bureau of Consumer Protection, Federal Trade Commission, 600 Pennsylvania Avenue NW, Washington, DC 20580.

SUPPLEMENTARY INFORMATION: Pursuant to Section 6(f) of the Federal Trade Commission Act, 15 U.S.C. 46(f), and FTC Rule 2.34, 16 CFR 2.34, notice is hereby given that the above-captioned consent agreement containing a consent order to cease and desist, having been filed with and accepted, subject to final approval, by the Commission, has been placed on the public record for a period of thirty (30) days. The following Analysis of Aid Public Comment describes the terms of the consent agreement and the allegations in the complaint. An electronic copy of the full text of the consent agreement package can be obtained at <https://www.ftc.gov/news-events/commission-actions>.

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before June 21, 2022. Write “Lions Not Sheep; File No. 222 3023” 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 <https://www.regulations.gov> website.

Because of heightened security screening, postal mail addressed to the Commission will be subject to delay. We strongly encourage you to submit your comments online through the <https://www.regulations.gov> website.

If you prefer to file your comment on paper, write “Lions Not Sheep; File No. 222 3023” on your comment and on the envelope, and mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex D), Washington, DC 20580.

Because your comment will be placed on the publicly accessible website at <https://www.regulations.gov>, you are solely responsible for making sure your comment does not include any sensitive or confidential information. In particular, your comment should not include sensitive personal information, such as your or anyone else’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 your comment does not include sensitive health 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 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 on the <https://www.regulations.gov> website—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from that website, 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.

Visit the FTC website at <http://www.ftc.gov> to read this document and the news release describing the proposed settlement. The FTC Act and other laws 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 June 21, 2022. 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>.

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 Lions Not Sheep Apparel, LLC; Lions Not Sheep Products, LLC; Lions Not Sheep Ventures, LLC; Lions Not Sheep LLC; and Sean Whalen (“Respondents”).

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 decide whether it should withdraw from the agreement or make final the agreement’s proposed order.

This matter involves Respondents’ advertising of hats, accessories, and apparel as “Made in USA.” According to the FTC’s complaint, Respondents represented that hats and non-apparel accessories were all or virtually all made in the United States. However, the complaint alleges that, in numerous instances, those hats and non-apparel accessories are wholly imported or contain significant imported content. Based on the foregoing, the complaint alleges Respondents engaged in deceptive acts or practices in violation of Section 5(a) of the FTC Act.

The complaint further alleges Respondents violated the Textile Fiber

Products Identification Act by (1) advertising articles of wearing apparel as of U.S. origin despite the fact they are wholly imported or incorporate significant imported materials, and (2) removing tags containing information required pursuant to the Textile Fiber Products Identification Act and replacing those tags with false country-of-origin designations.

The proposed consent order contains provisions designed to prevent Respondents from engaging in similar acts and practices in the future. Consistent with the FTC’s Made in USA Labeling Rule, 16 CFR part 323, and Enforcement Policy Statement on U.S.-Origin Claims, Part I prohibits Respondents from making U.S.-origin claims for their products unless: (1) The final assembly or processing of the product occurs in the United States, all significant processing that goes into the product occurs in the United States, and all or virtually all ingredients or components of the product are made and sourced in the United States; (2) a clear and conspicuous qualification appears immediately adjacent to the representation that accurately conveys the extent to which the product contains foreign parts, ingredients or components, and/or processing; or (3) for a claim that a product is assembled in the United States, the product is last substantially transformed in the United States, the product’s principal assembly takes place in the United States, and United States assembly operations are substantial.

Part II prohibits Respondents from making any representation about a product or service, including any representation regarding country of origin, unless the representation is not misleading and Respondents have a reasonable basis substantiating it.

Part III requires Respondents to make certain disclosures about the country of origin of any product subject to the Textile Fiber Products Identification Act.

Parts IV through VI are monetary provisions. Part IV imposes a judgment of \$211,335. Part V includes additional monetary provisions relating to collections. Part VI requires Respondents to provide sufficient customer information to enable the Commission to administer consumer redress, if appropriate.

Part VII is a notice provision requiring Respondents to identify and notify certain consumers of the FTC’s action within 30 days after the issuance of the order, or within 30 days of the consumer’s identification, if identified later. Respondents are also required to

submit reports regarding their notification program.

Parts VIII through XI are reporting and compliance provisions. Part VIII requires Respondents to acknowledge receipt of the order, to provide a copy of the order to certain current and future principals, officers, directors, and employees, and to obtain an acknowledgement from each such person that they have received a copy of the order. Part IX requires Respondents to file a compliance report within one year after the order becomes final and to notify the Commission within 14 days of certain changes that would affect compliance with the order. Part X requires Respondents to maintain certain records, including records necessary to demonstrate compliance with the order. Part XI requires Respondents to submit additional compliance reports when requested by the Commission and to permit the Commission or its representatives to interview Respondents' personnel.

Finally, Part XII 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.

April J. Tabor,
Secretary.

[FR Doc. 2022-10748 Filed 5-18-22; 8:45 am]

BILLING CODE 6750-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Solicitation of Nominations for Appointment to CDC's Advisory Committee to the Director (ACD) Data and Surveillance Workgroup (DSW); Re-Opening of Solicitation Period

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: Notice is hereby given of a change in the solicitation of CDC's Advisory Committee to the Director (ACD) Data and Surveillance Workgroup (DSW). In the **Federal Register** notice published on May 4, 2022, nominations for appointment to CDC's ACD DSW workgroup were due May 16, 2022. Nominations are now due May 27, 2022.

DATES: Nominations for membership on the DSW workgroup must be received no later than May 27, 2022. Late nominations will not be considered for membership.

ADDRESSES: All nominations (cover letters and curriculum vitae) should be emailed to DSWACD@cdc.gov with the subject line: "Nomination for CDC ACD DSW Workgroup."

FOR FURTHER INFORMATION CONTACT: Rachel Holloway, MPH, Office of the Chief of Staff, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H21-10, Atlanta, Georgia 30329-4027; Telephone: (404) 639-7000; Email: DSWACD@cdc.gov.

SUPPLEMENTARY INFORMATION:

Background: The purpose of the ACD, CDC is to advise the Secretary, HHS, and the Director, CDC, on policy and broad strategies that will enable CDC to fulfill its mission of protecting health through health promotion, prevention, and preparedness. The ACD, CDC consists of up to 15 non-federal members, including the Chair, knowledgeable in areas pertinent to the CDC mission, such as health policy, public health, global health, preparedness, preventive medicine, the faith-based and community-based sector, and allied fields.

Purpose: The establishment and formation of the DSW is to provide input to the ACD, CDC on agency-wide activities related to the scope and implementation of CDC's data modernization strategy across the agency, ultimately playing a key role in the agency's work with public health, healthcare, and academic and private sector partners and with the promotion of equity. The DSW membership will consist of approximately 15 members. It will be co-chaired by two current ACD, CDC Special Government Employees. The DSW co-chairs will present their findings, observations, and work products at one or more ACD, CDC meetings for discussion, deliberation, and decisions (final recommendations to CDC).

Nomination Criteria: DSW members will serve terms ranging from six months to one year and be required to attend DSW meetings approximately one to two times per month (virtually or in person), and contribute time between meetings for research, consultation, discussion, and writing assignments.

Nominations are being sought for individuals who have the expertise and qualifications necessary to contribute to the accomplishments of the committee's/workgroup's objectives. Nominees will be selected based on expertise in the fields of public health

science and practice; public health preparedness and response; public health policy development, analysis, and implementation; public health surveillance and informatics; data analysis, data science, and forecasting; health information technology; and healthcare delivery from jurisdictional government agencies, non-government organizations, academia, and the private sector. To ensure a diverse workgroup composition, nominees with front line and field experience at the local, state, tribal, and territorial levels are encouraged to apply. This includes nominees with experience working for, and with, community-based organizations and other non-profit organizations. Federal employees will not be considered for membership. Selection of members is based on candidates' qualifications to contribute to the accomplishment of the DSW's objectives.

HHS policy stipulates that membership be balanced in terms of points of view represented and the workgroup's function. Appointments shall be made without discrimination based on age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status. Nominees must be U.S. citizens and cannot be full-time employees of the U.S. Government. Current participation on federal workgroups or prior experience serving on a federal advisory committee does not disqualify a candidate; however, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships. Interested candidates should submit the following items:

- A one-half to one-page cover letter that includes your understanding of, and commitment to, the time and work necessary; one to two sentences on your background and experience; and one to two sentences on the skills/perspective you would bring to the DSW.

- Current curriculum vitae which highlights the experience and work history being sought relevant to the criteria set forth above, including complete contact information (telephone numbers, mailing address, email address).

Nominations may be submitted by the candidate him or herself, or by the person/organization recommending the candidate.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other