

**FEDERAL FINANCIAL INSTITUTIONS
EXAMINATION COUNCIL****[Docket No. AS23–03]****Appraisal Subcommittee Notice of
Meeting****AGENCY:** Appraisal Subcommittee of the Federal Financial Institutions Examination Council.**ACTION:** Notice of Special Meeting.

Description: In accordance with Section 1104(b) of Title XI of the Financial Institutions Reform, Recovery, and Enforcement Act of 1989, as amended, notice is hereby given that the Appraisal Subcommittee (ASC) met for a Special Meeting on this date.

Location: Virtual meeting via Webex.

Date: April 12, 2023.

Time: 12:15 p.m. ET.

Action and Discussion Item**Fiscal Year 2023 Budget Amendment**

The ASC convened a Special Meeting to vote on a budget amendment in the amount of \$267,065 to the ASC's Fiscal Year 2023 budget. The vote passed 6–0 with the FDIC abstaining.

James R. Park,

Executive Director.

[FR Doc. 2023–08507 Filed 4–20–23; 8:45 am]

BILLING CODE 6700–01–P

FEDERAL TRADE COMMISSION**[File No. 232 3007]****Cycra Inc.; 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 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 May 22, 2023.

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 “Cycra Inc.; File No. 232 3007” 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, please mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex P), Washington, DC 20580.

FOR FURTHER INFORMATION CONTACT: Julia Solomon Ensor (202–326–2377), Attorney, Division of Enforcement, 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 30 days. The following Analysis to 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 May 22, 2023. Write “Cycra Inc.; File No. 232 3007” 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 “Cycra Inc.; File No. 232 3007” 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 P), 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 it receives on or before May 22, 2023. 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 (the “Commission”) has accepted, subject to final approval, an agreement containing a consent order from Cycra Inc. and

Steven Chadwick James (“Respondents”). The proposed consent order has been placed on the public record for 30 days for receipt of comments from interested persons. Comments received during this period will become part of the public record. After 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’ labeling and advertising of motorcycle, motocross, and all-terrain vehicle products as “Made in USA.” According to the FTC’s complaint, Respondents labeled and advertised their products as made in the United States even though, in numerous instances, those products were wholly imported or contained significant imported content. Based on the foregoing, the complaint alleges Respondents violated Section 5 of the Federal Trade Commission Act, 15 U.S.C. 45(a), and section 323.2 of the Made in USA Labeling Rule, 16 CFR 323.2.

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 its 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 the country of origin of a product or service, unless the representation is not misleading and Respondents have a reasonable basis substantiating it.

Parts III through V are monetary provisions. Part IV imposes a judgment of \$872,577 and partially suspends that judgment based on the Respondents’

sworn financial statements. If the Commission concludes any Respondent made a material misrepresentation or omission in that Respondent’s sworn financial statement, the suspension as to that Respondent is lifted and the full judgment is immediately due. Part IV includes additional monetary provisions relating to collections. Part V requires Respondents to provide sufficient customer information to enable the Commission to administer consumer redress, if appropriate.

Part VI 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 VII through VIII are reporting and compliance provisions. Part VII 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 VIII 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 IX requires Respondents to maintain certain records, including records necessary to demonstrate compliance with the order. Part X 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 XI is a “sunset” provision, terminating the order after 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,

Secretary.

[FR Doc. 2023–08471 Filed 4–20–23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10302]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 June 20, 2023.

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.