

help us prepare for the ICCR–11 meeting that will be held July 12–14, 2017, in Brasilia, Brazil.

DATES: The public meeting will be held on May 25, 2017, from 2 p.m. to 4 p.m. See the **SUPPLEMENTARY INFORMATION** section for registration date and information.

ADDRESSES: The public meeting will be held at the Food and Drug Administration, Center for Food Safety and Applied Nutrition, 5001 Campus Dr., Wiley Auditorium (first floor), College Park, MD 20740.

FOR FURTHER INFORMATION CONTACT: Jonathan Hicks, Office of Cosmetics and Colors, Food and Drug Administration, 5001 Campus Dr. (HFS–125), College Park, MD 20740, email: jonathan.hicks@fda.hhs.gov, 240–402–1375.

SUPPLEMENTARY INFORMATION:

I. Background

The purpose of the multilateral framework on the ICCR is to pave the way for the removal of regulatory obstacles to international trade while maintaining global consumer protection. The purpose of the meeting is to invite public input on various topics pertaining to the regulation of cosmetics. We may use this input to help us prepare for the ICCR–11 meeting that will be held July 12–14, 2017, in Brasilia, Brazil.

ICCR is a voluntary international group of cosmetics regulatory authorities from Brazil, Canada, the European Union, Japan, and the United States of America. These regulatory authority members will enter into constructive dialogue with their relevant cosmetics industry trade associations and public advocacy groups. Currently, the ICCR members are: The Brazilian Health Surveillance Agency; Health Canada; the European Commission Directorate-General for Internal Market, Industry, Entrepreneurship, and Small and Medium-sized Enterprises; the Ministry of Health, Labor, and Welfare of Japan; and FDA. All decisions made by consensus will be compatible with the laws, policies, rules, regulations, and directives of the respective administrations and governments. Members will implement and/or promote actions or documents within their own jurisdictions and seek convergence of regulatory policies and practices. Successful implementation will need input from stakeholders.

II. Topics for Discussion at the Public Meeting

We will make the agenda for the public meeting available on the Internet

at <http://www.fda.gov/Cosmetics/InternationalActivities/ICCR/default.htm>. Depending on the number of requests for oral presentations, we intend to have an agenda available by May 18, 2017.

III. Participating in the Public Meeting

Registration: To register for the public meeting, send registration information (including your name, title, affiliation, address, email, and telephone), to Jonathan Hicks by May 11, 2017. If you would like to listen to the meeting by phone, please submit a request for a dial-in number by May 11, 2017. If you need special accommodations due to a disability, please contact Jonathan Hicks by May 18, 2016.

Requests for Oral Presentations: If you wish to make an oral presentation, you should notify Jonathan Hicks by May 11, 2017, and submit a brief statement of the general nature of the evidence or arguments that you wish to present, your name, title, affiliation, address, email, and telephone, and indicate the approximate amount of time you need to make your presentation. You may present proposals for future ICCR agenda items, data, information, or views, in person or in writing, on issues pending at the public meeting. There will be no presentations by phone. Time allotted for oral presentations may be limited to 10 minutes or less for each presenter.

Transcripts: Please be advised that as soon as a transcript of the public meeting is available, it will be accessible at <https://www.regulations.gov>. It may also be viewed at the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20850.

Dated: April 3, 2017.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2017–06938 Filed 4–6–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Findings of Research Misconduct; Correction

AGENCY: Office of the Secretary, HHS.

ACTION: Correction of notice.

SUMMARY: This document corrects an error that appeared in the notice published in the June 8, 2016, **Federal Register** entitled “Findings of Research Misconduct.”

DATES:

Effective Date: April 7, 2017.

Applicability Date: The correction notice is applicable for the Findings of Research Misconduct notice published on June 8, 2016.

FOR FURTHER INFORMATION CONTACT: Ms. Karen Gorirossi at 240–453–8800.

SUPPLEMENTARY INFORMATION:

I. Background

In FR Doc. 2016–13541 of June 8, 2016 (81 FR 36932–36933), there was a typographical error involving one of the papers cited in the notice. The error is identified and corrected in the Correction of Errors section below.

II. Correction of Errors

In FR Doc. 2016–13541 of June 8, 2016 (81 FR 36932–36933), make the following correction:

1. On page 36932, third column, in FR Doc. 2016–13541, second to last paragraph, line 5, change “August” to “December” so that the text reads “falsified twenty-four (24) fluorescent image panels by drawing staining in Photoshop and falsely labeling them in Figures 5B, 5C, 5D, 5E, 7A, 7B, 7D, 8A, 8B, 9A, and 9B in the December 2015 *Development* paper and included some of the same images in four (4) figures in the *ASCB* 2015 poster and in two (2) figures in the *MARZ* 2015 poster”

Dated: March 30, 2017.

Kathryn Partin,

Director, Office of Research Integrity.

[FR Doc. 2017–07006 Filed 4–6–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the meeting of the Council of Councils.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The open session will be videocast and can be accessed from the NIH Videocasting and Podcasting Web site (<http://videocast.nih.gov>).

A portion of the meeting will be closed to the public in accordance with