Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5441, Silver Spring, MD 20993–0002, 301–796–8398; or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code PEAC. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 26, 2017 (82 FR 34681), FDA announced that a meeting of the Patient Engagement Advisory Committee would be held on October 11 and 12, 2017. On page 34681, in the third column, the **DATES** section is changed to reflect the time of these meetings on the announced dates.

On page 34682, in the first column, in the *Procedure* section, the third sentence is changed to reflect new times for oral presentations on October 11 and 12.

On page 34682, in the second column, a *Webcast Information* section is added before the last paragraph of the document. The amendments read as follows:

DATES: The meeting will be held on October 11, 2017, from 12:30 p.m. to 6 p.m. and October 12, 2017, from 8 a.m. to 5 p.m.

Procedure: Oral presentations from the public will be scheduled between approximately 2:30 p.m. to 3 p.m. on October 11, 2017, and approximately 1 p.m. to 2:30 p.m. on October 12, 2017.

Webcast Information: This meeting will also be made available to the public via webcast. The links for the webcasts are below: October 11, 2017: ''Patient Engagement Advisory Committee Meeting, Day 1,'' *https://event.webcasts.com/starthere.jsp?ei=1157277&tp_key=5580d0c7a5.* October 12, 2017: ''Patient Engagement Advisory Committee Meeting, Day 2,'' Morning Session—*https://event.webcasts.com/starthere.jsp?ei=1157280&tp_key=dfcde848fe;* and Afternoon Session—*https://event.webcasts.com/starthere.jsp?ei=1157282&tp_key=6d832a247e.*

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: September 29, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–21317 Filed 10–3–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Findings of Research Misconduct

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: Notice is hereby given that the Office of Research Integrity (ORI) has taken final action in the following case:

Azza El-Remessy, Ph.D., University of Georgia, College of Pharmacy: Based on the report of an investigation conducted by the University of Georgia, College of Pharmacy (UGCP) and additional analysis conducted by ORI in its oversight review, ORI found that Dr. Azza El-Remessy, former Associate Professor, Department of Clinical and Administrative Pharmacy, UGCP, engaged in research misconduct in research supported by National Eye Institute (NEI), National Institutes of Health (NIH), grants R01 EY011766, R01 EY022408, and R01 EY04618, National Heart, Lung, and Blood Institute (NHLBI), NIH, grant R01 HL056259, and National Cancer Institute (NCI), NIH, grant K01 CA89689.

ORI found that false Western blot data were included in:

- *J Cell Sci.* 118(Pt. 1):243–52, 2005 (hereafter referred to as ''*J Cell Sci.* 2005''). Retraction in: *J Cell Sci.* 129(16):3203, 2016.
- *FASEB J.* 21(10):2528–39, 2007 (hereafter referred to as ''*FASEB J.* 2007''). Retraction in: *FASEB J.* 31(1):421, 2017.
- *PLoS One* 8(8):e71868, 2013 (hereafter referred to as ''*PLoS One* 2013'').

As a result of its investigation, UGCP recommended that *PLoS One* 2013 be corrected. As a result of the investigation, *J Cell Sci.* 2005 and *FASEB J.* 2007 have been retracted.

ORI found that Respondent intentionally, knowingly, or recklessly used the same Western blot bands to represent different experimental results. Specifically, Respondent reused and relabeled bands in:

1. Figure 3B, *J Cell Sci.* 2005, to represent p38 bands from retinal cultured endothelial cells in high glucose in the absence of exogenous VEGF and also cells in peroxynitrite in the presence of exogenous VEGF.

2. Figure 4A, *J Cell Sci.* 2005, to represent nitrotyrosine immunoprecipitations from retinal endothelial cells cultured in normal glucose in the presence or absence of FeTTP; the Respondent also duplicated controls for p85 immunoprecipitation by using three bands representing 2 normal glucose and 1 high glucose treatments, flipping them horizontally (mirror images) to also represent 2 high glucose and 1 peroxynitrite treatments.

3. Figure 4B, *J Cell Sci.* 2005, to represent p85 immunoprecipitations from retinal endothelial cells stimulated with VEGF and also cells treated with either high glucose or peroxynitrite.

4. Figure 4A, *PLoS One* 2013, to represent immunoprecipitations for phosphorylated GSK-3 (p-GSK-3) in cells with normal glucose or high glucose for day 1 and to also represent cells treated with VEGF or VEGF+VEGFI (inhibitor); the Respondent also duplicated GSK-3 controls by using the same bands to represent high glucose treatment for day 1 and day 3 treatments, flipping them horizontally, to also represent for VEGF and VEGFRI treatments.

5. Figure 3, *FASEB J.* 2007, to represent phosphorylated VEGF2 (P-VEGF2) protein expression in microvascular endothelial cells in: Lanes 1 and 8, lanes 2 and 5, and lanes 6 and 7, where each lane represents different experimental conditions.

Dr. El-Remessy entered into a Voluntary Settlement Agreement (Agreement) to resolve this matter without further expenditure of time or other resources. Dr. El-Remessy accepts ORI's findings of research misconduct as set forth above but neither admits nor denies ORI's findings of research misconduct. The settlement is not an admission of liability on the part of the Respondent. Dr. El-Remessy voluntarily agreed, beginning on September 12, 2017:

(1) To have her research supervised for a period of three (3) years beginning with the effective date of the Agreement; Respondent agreed that prior to the submission of an application for U.S. Public Health Service (PHS) support for a research project on which the Respondent's participation is proposed and prior to Respondent's participation in any capacity on PHS-supported research, Respondent shall ensure that a plan for supervision of Respondent's duties is submitted to ORI for approval; the supervision plan must be designed to ensure the scientific integrity of Respondent's research contribution; Respondent agreed that she shall not participate in any PHS-supported research until such a supervision plan is submitted to and approved by ORI; Respondent agreed to maintain responsibility for compliance with the agreed upon supervision plan;

(2) that for three (3) years beginning with the effective date of the Agreement,

any institution employing her shall submit, in conjunction with each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent is involved, a certification to ORI that the data provided by Respondent are based on actual experiments or are otherwise legitimately derived and that the data, procedures, and methodology are accurately reported in the application, report, manuscript, or abstract;

(3) to exclude herself voluntarily from serving in any advisory capacity to PHS including, but not limited to, service on any PHS advisory committee, board, and/or peer review committee, or as a consultant for a period of three (3) years, beginning with the effective date of the Agreement; and

(4) that as a condition of the Agreement, Respondent will request that *PLoS One* 8(8):e71868, 2013 be corrected or retracted.

FOR FURTHER INFORMATION CONTACT: Director, Office of Research Integrity, 1101 Wootton Parkway, Suite 750, Rockville, MD 20852, (240) 453–8200.

Kathryn M. Partin,

Director, Office of Research Integrity.

[FR Doc. 2017–21367 Filed 10–3–17; 8:45 am]

BILLING CODE 4150–31–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Human Genome Research Institute Special Emphasis Panel, SEP U24 Research Resource.

Date: November 30, 2017.

Time: 11:00 a.m. to 2:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Human Genome Research Institute, 5635 Fishers Lane, Suite 3146, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Rudy O. Pozzatti, Ph.D., Scientific Review Officer, Scientific Review Branch, National Human Genome Research Institute, 5635 Fishers Lane, Suite 4076, MSC 9306, Rockville, MD 20852, (301) 402–0838, *pozzattr@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.172, Human Genome Research, National Institutes of Health, HHS)

Dated: September 28, 2017.

Sylvia L. Neal,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–21263 Filed 10–3–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Multi-site Clinical Trials.

Date: October 19, 2017.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Shang-Yi Anne Tsai, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, Division of Extramural Research, National Institute on Drug Abuse, NIH, DHHS, 6001 Executive Boulevard, Room 4228, MSC 9550, Bethesda, MD 20892, 301–827–5842, *shangyi.tsai@nih.gov*.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; NIDA Mentored Clinical Scientists Development Program Award in Drug Abuse and Addiction (K12).

Date: October 25, 2017.

Time: 9:00 a.m. to 12:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852.

Contact Person: Susan O. McGuire, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, National Institutes of Health, DHHS, 6001 Executive Blvd., Room 4245, Rockville, MD 20852, (301) 827–5817, *mcguireso@mail.nih.gov*.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; NIH Pathway to Independence Award (K99/R00).

Date: October 25, 2017.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852.

Contact Person: Susan O. McGuire, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, National Institutes of Health, DHHS, 6001 Executive Blvd., Room 4245, Rockville, MD 20852, (301) 827–5817, *mcguireso@mail.nih.gov*.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; (T32) Ruth L. Kirschstein National Research Service Award (NRSA) Institutional Research Training Grants.

Date: October 26–27, 2017.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852.

Contact Person: Susan O. McGuire, Ph.D. Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, National Institutes of Health, DHHS, 6001 Executive Blvd., Room 4245, Rockville, MD 20852, (301) 827–5817, *mcguireso@mail.nih.gov*.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Exploring Novel RNA Modifications in HIV/AIDS and Substance Use Disorders (R01, R21).

Date: November 1, 2017.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852,

Contact Person: Susan O. McGuire, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, National Institutes of Health, DHHS, 6001 Executive Blvd., Room 4245, Rockville, MD 20852, (301) 827–5817, *mcguireso@mail.nih.gov*.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Cutting-Edge Basic Research Awards (CEBRA) (R21).

Date: November 29, 2017.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Bethesda North Marriott Hotel & Conference Center, 5701 Marinelli Road, Bethesda, MD 20852.