

indicate that the settlement is inappropriate, improper, or inadequate. The Agency's response to any comments received will be available for public inspection at the City of Los Angeles Central Library, Science and Technology Department, 630 West 5th Street, Los Angeles CA 90071 and at the EPA Region 9 Superfund Records Center, Mail Stop SFD-7C, 95 Hawthorne Street, Room 403, San Francisco, CA 94105.

DATES: Comments must be submitted on or before February 2, 2012.

ADDRESSES: The proposed settlement is available for public inspection at the EPA Region 9 Superfund Records Center, Mail Stop SFD-7C, 95 Hawthorne Street, Room 403, San Francisco, CA 94105. A copy of the proposed settlement may also be obtained from the EPA Region 9 Superfund Record Center, 95 Hawthorne Street, Mail Stop SFD-7C, Room 403, San Francisco, CA 94105, (415) 820-4700. Comments should reference the North Hollywood Operable Unit of the San Fernando Valley Area 1 Superfund Site, and EPA Docket No. 9-2011-0015 and should be addressed to Michael Massey, EPA Region 9, 75 Hawthorne Street, Mail Stop ORC-3, San Francisco, CA 94105.

FOR FURTHER INFORMATION CONTACT: Kelly Manheimer, EPA Region 9, 75 Hawthorne Street, Mail Stop SFD-7-1, San Francisco, CA 94105, (415) 972-3290.

Dated: December 22, 2011.

Kathleen Salyer,

Acting Superfund Division Director.

[FR Doc. 2011-33667 Filed 12-30-11; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[FRL-9612-9]

Biological Processors of Alabama; Decatur, Morgan County, AL; Notice of Settlement

AGENCY: Environmental Protection Agency.

ACTION: Notice of Settlement.

SUMMARY: Under Section 122(h)(1) of the Comprehensive Environmental Response, Compensation and Liability Act (CERCLA), the United States Environmental Protection Agency has entered into a settlement for reimbursement of past response costs concerning the Biological Processors of Alabama Superfund Site located in Decatur, Morgan County, Alabama.

DATES: The Agency will consider public comments on the settlement until February 2, 2012. The Agency will consider all comments received and may modify or withdraw its consent to the settlement if comments received indicate that the settlement is inappropriate, improper, or inadequate.

ADDRESSES: Copies of the settlement are available from Ms. Paula V. Painter. Submit your comments by Site name Biological Processors of Alabama Superfund Site by one of the following methods:

- www.epa.gov/region4/waste/sf/enforce.htm.

- *Email. Painter.Paula@epa.gov.*

FOR FURTHER INFORMATION CONTACT: Paula V. Painter at (404) 562-8887.

Dated: December 14, 2011.

Anita L. Davis,

Chief, Superfund Enforcement & Information Management Branch, Superfund Division.

[FR Doc. 2011-33680 Filed 12-30-11; 8:45 am]

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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:

Jennifer Jamieson, State University of New York, Upstate Medical University: Based on the report of an investigation conducted by the State University of New York, Upstate Medical University (SUNY US) and additional analysis conducted by ORI in its oversight review, ORI found that Ms. Jennifer Jamieson, former graduate student, Department of Cell and Developmental Biology, SUNY US, engaged in research misconduct in research supported by National Institute of General Medical Sciences (NIGMS), National Institutes of Health (NIH), grant R01 GM047607-18A1, and National Heart, Lung, and Blood Institute (NHLBI), NIH, grants R01 HL70244-05.

ORI found that Respondent engaged in research misconduct by falsifying data that were included in grant application R01 GM047607-18A1, in a manuscript submitted for publication to the *Journal of Cell Biology*, and in several interdepartmental data presentations. Specifically, ORI found that:

- Respondent falsified Figure 1A in a manuscript submitted for publication to the *Journal of Cell Biology*, by altering immunoprecipitation Western blot data to make this experiment appear that no Vav2 SH2 was associated with PKL 3YF, when in fact it did. In addition, the Respondent falsified five figures depicting Western blots of similar experiments in four laboratory meeting presentations. The purpose of the falsifications was to show that the experimental results were as described when they were not, or to show that the results were of greater significance than they actually were.

- Respondent falsified Figure 3I in a manuscript submitted for publication to the *Journal of Cell Biology* by falsely labeling a Western blot to indicate levels of expression for various Vav2 mutants, when the experimental data were taken from a completely unrelated experiment.

- Respondent falsified Figure 6A in an interdepartmental laboratory presentation by falsifying Western blot data to falsely depict Paxillin and Hic-5 expression and phosphorylation levels after siRNA treatment.

- Respondent falsified Figure 5 from NIGMS, NIH, grant application GM047607-18A1, by falsifying Western blot data to support the hypothesis that co-transfection of PKL plus RhoA GEF Vav2 induces RhoA activation and signaling upon plating on fibronectin.

Ms. Jamieson has entered into a Voluntary Settlement Agreement (Agreement). Ms. Jamieson neither admits nor denies ORI's finding of scientific misconduct nor any particular finding of fact asserted in support of that finding. The settlement is not an admission of liability on the part of the Respondent.

Ms. Jamieson has voluntarily agreed for a period of three (3) years, beginning on December 20, 2011:

(1) To have her research supervised if employed by an institution that receives or applies for U.S. Public Health Service (PHS) funding; Respondent agrees that prior to the submission of an application for 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 her duties is submitted to ORI for approval; the supervision plan must be designed to ensure the scientific integrity of Respondent's research contribution; Respondent agrees that she shall not participate in any PHS-supported research until such a supervision plan is submitted to and approved by ORI;

Respondent agrees to maintain responsibility for compliance with the agreed upon supervision plan;

(2) that 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 were accurately reported in the application, report, manuscript, or abstract; and

(3) to exclude herself 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 FURTHER INFORMATION CONTACT:

Director, Division of Investigative Oversight, Office of Research Integrity, 1101 Wootton Parkway, Suite 750, Rockville, MD 20852, (240) 453-8800.

John Dahlberg,

Director, Division of Investigative Oversight, Office of Research Integrity.

[FR Doc. 2011-33650 Filed 12-30-11; 8:45 am]

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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:

Mahesh Visvanathan, Ph.D., Kansas University: Based on an inquiry conducted and written admission obtained by Kansas University (KU) and additional analysis conducted by ORI in its oversight review, ORI found that Dr. Mahesh Visvanathan, Research Assistant Professor in the K-INBRE¹ Bioinformatics Core Facility, KU, engaged in research misconduct in research supported by U.S. Public Health Service (PHS) funds, specifically the INBRE program of the National Center for Research Resources (NCRR), National Institutes of Health (NIH), grant P20 RR016475.

Specifically, ORI found that Respondent engaged in research

misconduct by intentionally and knowingly plagiarizing large amounts of text from other writers' published papers without attribution or citation in the following three (3) papers and one (1) abstract. The specific published documents as well as the relevant source documents are:

- Visvanathan, M., Adagarla, B., Lushington, G., Sittampalam, S., *Proceedings of the 2009 International Joint Conference on Bioinformatics, Systems, Biology and Intelligent Computing*, 2009, 494-497. Greater than half (50%) of the total text was obtained from (1) Yang, C.-S., Chuang, L.-Y., Ke, C.-H., Yang, C.-H., *International Journal of Computer Science, International Association of Engineers*, August 2008 35(3), (2) Goffard, N. and Weiller, G., *Nucleic Acids Research*, 2007, 35L:W176- W181, and (3) Chuang, L.-Y., Yang, C.-H., Tu, C.-J., Yang, C.-H., *Proceedings of the Joint Conference on Information Sciences*, Atlantis Press, October 2006.

Retracted: Retracted administratively by IEEE on Jan 5, 2011 http://ieeexplore.ieee.org/xpl/freeabs_all.jsp?arnumber=5260432.

- Vijayan, A.; Skariah, B. E., Nair, B.; Lushington, G., Subramanian, S., Visvanathan, M., *Proceedings of the IEEE International Conference on Bioinformatics and Biomedicine Workshop*, 2009, BIBMW2009, 267-271. Approximately 15% of the text was plagiarized from Goffard, N. and Weiller, G., *Nucleic Acids Research*, 2007, 35L:W176-W181.

Retracted: Retracted administratively by IEEE on Jan 5, 2011 <http://www.computer.org/portal/web/csdl/doi/10.1109/BIBMW.2009.5332106>.

- Visvanathan, M., Netzer, M., Seger, M., Adagarla, B. S., Baumgartner, C., Sittampalam, S., Lushington, G., *International Journal of Computational Biology and Drug Design*, 2009, 2,236-251. A complete paragraph of the text was plagiarized from Goffard, N. and Weiller, G., *Nucleic Acids Research*, 2007, 35L:W176- W181.

- Adagarla, B., Lushington, G., Visvanathan, M., ISMB International Conference, January 2009; the entire abstract for this poster was obtained by plagiarizing text from Pihur, V., Datta, S., Datta S., *Genomics*, 2003, 92:400-403.

Dr. Visvanathan has entered into a Voluntary Settlement Agreement (Agreement) and has voluntarily agreed for a period of two (2) years, beginning on December 20, 2011:

(1) To have any PHS-supported research supervised; ORI acknowledges that Respondent's research is currently being supervised by KU; Respondent

shall ensure that a plan for supervision of his PHS-related duties is submitted to ORI for approval either within two weeks of this Agreement becoming final or prior to receiving or applying for PHS funds if such support is not current at the time this Agreement is completed; the supervision plan must be designed to ensure the scientific integrity of his research contribution; because of the ongoing review of Respondent's research by KU, ORI will only require a summary report on the first and second anniversary of the Agreement detailing how KU has ensured that Respondent's research and language in PHS grant applications and reports of PHS-supported research have been verified to be his own and accurately reported; Respondent agrees to maintain responsibility for compliance with the agreed upon supervision plan;

(2) That this annual summary, provided by any institution employing him, shall provide assurance that each application for PHS funds, or report, manuscript, or abstract involving PHS-supported research in which Respondent was involved, was based on actual experiments or was otherwise legitimately derived, that the data, procedures, and methodology were accurately reported in the application, report, manuscript, or abstract, and that the text in such submissions was his own or properly cited the source of copied language and ideas; and

(3) To exclude himself 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 FURTHER INFORMATION CONTACT:

Director, Division of Investigative Oversight, Office of Research Integrity, 1101 Wootton Parkway, Suite 750, Rockville, MD 20852. (240) 453-8800.

John Dahlberg,

Director, Division of Investigative Oversight, Office of Research Integrity.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-D-0916]

Draft Guidance for Industry and Food and Drug Administration Staff; Medical Device Classification Product Codes; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.