TRANSACTION GRANTED EARLY TERMINATION—Continued

ET date	Trans No.	ET req status	Party name
		G	Powersport Auctioneer Holdings, LLC.
		G	Powersport Auctioneer Holdings, LLC.
	20110269	G	Religare Enterprises Limited.
		G	Landmark Partners, Inc-CT.
		G	Landmark Partners, LLC.
	20110273	G	ICV Partners II, L.P.
		G	Cargo Airport Services U.S.A. LLC.
		G	Cargo Airport Services U.S.A. LLC.
	20110275	G	TPG Star, L.P.
		G	ZS VPSI, L.P.
		G	VPSI, Inc.
		G	VPSI, L.L.C.
	20110276	G	Alliant Techsystems Inc.
		G	Sentinel Capitai Partners IV, L.P.
	20110290	G	North American Rescue, LLC.
		G	Ascent Media Corporation.
		G	ABRY Partners IV, LP.
		G	Monitronics International, Inc.
	20110298	G	Charlesbank Equity Fund VII, Limited Partnership.
		G	Behrman Capital III L.P.
		G	Peacock Holding Company, Inc.

FOR FURTHER INFORMATION CONTACT: Sandra M. Peay, Contact Representative or Renee Chapman, Contact Representative, Federal Trade Commission, Premerger Notification Office, Bureau of Competition, Room H–303, Washington, DC 20580, (202) 326–3100.

By Direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2010–31092 Filed 12–10–10; 8:45 am]

BILLING CODE 6750–01–M

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:

Sagar S. Mungekar, PhD, New York University School of Medicine: Based on the Respondent's written admission and set forth below, the New York University School of Medicine (NYUSOM) and the Office of Research Integrity (ORI) found that Sagar S. Mungekar, PhD, former MD/PhD student in the Sackler Institute of Graduate Biomedical Sciences at NYUSOM, engaged in research misconduct in research supported by National Institute of General Medical Sciences (NIGMS), National Institutes of Health (NIH), grants R01 GM35769, R01 GM55624, and T32 GM07308, and National Institute of Allergy and Infectious Diseases (NIAID), NIH, grant T32 AI007180.

Dr. Mungekar admitted that in his PhD thesis he "increased statistical significance of the calculated means and standards of deviation [*sic*] of the UV spectrophometic [*sic*] data presented by discarding certain experimental data and thus presented data that was falsified. In addition, as the repression ratios calculated and conclusions reached based on these data that included falsified data, those values and conclusions are fabricated. Approximately, 60–75 of the [Respondent's] PhD research data was changed or falsified." Dr. Mungekar also admitted "while doing these experiments, I did not sequence all of the constructs that I constructed, thus, I could not be certain of the exact identity of the plasmids in question."

ORI found that Dr. Mungekar engaged in research misconduct (42 CFR 93.103) by fabricating and falsifying data. Specifically, ORI found that Dr. Mungekar falsified five tables and five figures (Tables 2–1, 2–2, 3–1, 4–1, 4–2 and Figures 2–3, 3–1, 3–2, 4–3, and 4–4) in his Ph.D. thesis entitled "Autoregulation of Ribonuclease E," by discarding certain spectrophotometric data, to increase statistical significance, used to calculate repression ratios and RNA decay rates. Dr. Mungekar also claimed to have constructed 53 different reporter plasmids with RNase E mutants, when sequencing data did not exist to support this claim.

Dr. Mungekar has entered into a Voluntary Settlement Agreement in which he has voluntarily agreed, for a period of three (3) years, beginning on November 22, 2010:

(1) That any institution that submits an application for PHS support for a research project on which the Respondent's participation is proposed or that uses him in any capacity on PHS-supported research, or that submits a report of PHS-funded research in which he is involved, must concurrently submit a plan for supervision of his duties to ORI for approval; the supervisory plan must be designed to ensure the scientific integrity of his research contribution; Respondent agrees that he will not participate in any PHS-supported research until such a supervision plan is submitted to ORI;

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

(3) to exclude himself voluntarily from serving in any advisory capacity to the U.S. Public Health Service (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. 2010–31168 Filed 12–10–10; 8:45 am]

BILLING CODE 4150–31–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Priority Setting for the Children's Health Insurance Program Reauthorization Act (CHIPRA) Pediatric Quality Measures Program—Notice of Correction

On pages 75469 and 75470, Volume 75, Number 232, **Federal Register** notice publication dated December 3, 2010, under **DATES** section, the correct date is: January 14, 2011. Also, on pages 75470 and 75471, under section **SUPPLEMENTARY INFORMATION** all Web links that include the word: "ahrg" should be changed to: "AHRQ".

Dated: December 7, 2010.

Carolyn M. Clancy,

Director, AHRQ.

[FR Doc. C1–2010–31110 Filed 12–10–10; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number NIOSH–219]

Implementation of Section 2695 (42 U.S.C. 300ff–131) of Public Law 111–87: Infectious Diseases and Circumstances Relevant to Notification Requirements

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

ACTION: General Notice and Request for Comments.

SUMMARY: The Ryan White HIV/AIDS Treatment Extension Act of 2009 (Pub. L. 111–87) addresses notification procedures for designated officers, medical facilities, and State and community public health officers regarding exposure of emergency response employees (EREs) to potentially life-threatening infectious diseases. The Secretary of Health and Human Services (Secretary) has delegated authority to the Director of the Centers for Disease Control and Prevention (CDC) to issue a list of potentially life-threatening infectious diseases, including emerging infectious diseases, to which EREs may be exposed in responding to emergencies (including a specification of those infectious diseases that are routinely transmitted through airborne or aerosolized means); guidelines describing circumstances in which employees may be exposed to these diseases; and guidelines describing the manner in which medical facilities should make determinations about exposures. CDC is seeking comment on the list of diseases and guidelines contained in this notice.

DATES: Comments must be received by February 11, 2011.

ADDRESSES: Comments on the content of this Notice should be in writing and addressed to:

- *E-mail:* NIOSH Docket Officer, *nioshdocket@cdc.gov.* Include "Infectious Diseases" and "42 U.S.C. 300ff–131" in the subject line of the message.
- *Mail:* NIOSH Docket Office, Robert A. Taft Laboratories, MS–C34, 4676 Columbia Parkway, Cincinnati, OH 45226.
- *Internet:* Federal e-rulemaking portal, *http://www.regulations.gov.* Follow the instructions for submitting comments.

Instructions: All submissions received must include the agency name and docket number for this Notice. All comments will be posted without change to *http://www.cdc.gov/niosh/docket/archive/docket219.html,* including any personal information provided. For detailed instructions on submitting comments and additional information about this process, *see* the "Public Participation" heading of the **SUPPLEMENTARY INFORMATION** section of this document.

Docket: For access to the docket to read background documents or comments received, go to *http://www.cdc.gov/niosh/docket/archive/docket219.html.*

FOR FURTHER INFORMATION CONTACT: Centers for Disease Control and Prevention, Attention: James Spahr, Associate Director, Emergency Preparedness & Response, Office of the Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, 1600 Clifton Road, NE., Mailstop E20, Atlanta, GA 30333. Telephone (404) 498–6185 (this is not a toll-free number).

SUPPLEMENTARY INFORMATION:

Table of Contents

Public Participation

Interested persons or organizations are invited to participate in this request for public comments by submitting written views, arguments, recommendations, and data. Comments are invited on any topic related to this proposal. In particular, CDC invites comment on the list of infectious diseases and both sets of guidelines discussed herein.

Comments submitted by e-mail or mail should be titled "Docket #219 Public Comments," addressed to the "NIOSH Docket Officer," and identify the author(s), return address, and a phone number, in case clarification is needed. Comments can be submitted by e-mail to *nioshdocket@cdc.gov* as e-mail text or as a Microsoft Word file attachment. Printed comments can be sent to the NIOSH Docket Office at the address above. All communications received on or before the closing date for comments will be fully considered by CDC in developing a final list of infectious diseases and guidelines which will be published in the **Federal Register.**

Introduction

The Ryan White HIV/AIDS Treatment Extension Act of 2009 (Pub. L. 111–87) amended the Public Health Service Act (PHS Act, 42 U.S.C. 201–300ii), including the addition of a Part G to Title XXVI, which addresses notification procedures and requirements for medical facilities, State public health officers and their designated officers regarding exposure of EREs to potentially life-threatening infectious diseases. (*See* Title XXVI, Part G of the PHS Act, codified as amended at 42 U.S.C. 300ff–131 to 300ff–140.)

For purposes of these notification requirements, Section 2695 [42 U.S.C. 300ff–131] requires the Secretary of Health and Human Services (Secretary) to develop and disseminate:

(1) A list of potentially life-threatening infectious diseases,