has taken final action in the following case:

Meleik Goodwill, Ph.D., Wadsworth Center, N.Y.S. Department of Health:

Based on the Wadsworth Center report and the oversight review conducted by the Office of Research Integrity (ORI), the U.S. Public Health Service (PHS) found that Meleik Goodwill, Ph.D., former postdoctoral fellow, Wadsworth Center, N.Y.S. Department of Health, engaged in research misconduct in research supported by National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health (NIH), grant R21 ES013269–02.

Specifically, PHS found that the Respondent engaged in research misconduct by the fabrication of data for growth curves presented in Figure 1 in the 2007 Journal of Neuroimmunology article (Goodwill, M.K., Lawrence, D.A., & Seegal, R.F. “Polychlorinated biphenyls induce proinflammatory cytokine release and dopaminergic dysfunction: Protection in interleukin-6 knockout mice.” Journal of Neuroimmunology 183(1–2):125–132, 2007), and by the use of composite images of Western-blot bands from unrelated experiments done in 2005 that were falsely labeled as if from different experiments to construct Figure 4A in the 2007 Journal of Neuroimmunology article. Figure 4B of the article also was falsified by use of identical sets of images of Western-blot bands from unrelated experiments done in 2005 that were falsely labeled as if from different experiments to construct Figure 4A in the 2007 Journal of Neuroimmunology article. The falsification of data in the 2007 Journal of Neuroimmunology article was retracted in J. Neuroimmunol. 197(1):197, 2008.

Dr. Goodwill has entered into a Voluntary Settlement Agreement in which she has voluntarily agreed, for a period of three (3) years, beginning on January 21, 2011:

(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 her in any capacity on PHS-supported research, or that submits a report of PHS-funded research in which she is involved, must concurrently submit a plan for supervision of her duties to ORI for approval; the supervisory plan must be designed to ensure the scientific integrity of her research contribution; Respondent agrees that she will not participate in any PHS-supported research until such a supervisory plan is submitted to ORI;

(2) That any institution employing her, in conjunction with each application for PHS funds, or report, manuscript, or abstract involving PHS-funded research in which she was involved, must certify to ORI that the data provided 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 herself voluntarily from service 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. 2011–2975 Filed 2–9–11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Assistant Secretary for Planning and Evaluation; Medicare Program; Meeting of the Technical Advisory Panel on Medicare Trustee Reports

AGENCY: Assistant Secretary for Planning and Evaluation, HHS.

ACTION: Notice of meeting.

SUMMARY: This notice announces public meetings of the Technical Advisory Panel on Medicare Trustee Reports (Panel). Notice of these meetings is given under the Federal Advisory Committee Act (5 U.S.C. App. 2, section 10(a)(1) and (a)(2)). The Panel will discuss the long-term rate of change in health spending and make recommendations to the Medicare Trustees on how the Trustees might more accurately estimate the long term rate of health spending in the United States. The discussion is expected to focus on highly technical aspects of estimation involving economics and actuarial science. Panelists are not restricted, however, in the topics that they choose to discuss.

Procedure and Agenda: This meeting is open to the public. The Panel will likely hear presentations by HHS staff presentations regarding long range growth. After any presentations, the Panel will deliberate openly on the topic. Interested persons may observe the deliberations, but the Panel will not hear public comments during this time. The Panel will also allow an open public session for any attendee to address issues specific to the topic.

Authority: 42 U.S.C. 217a; Section 222 of the Public Health Services Act, as amended. The panel is governed by provisions of Public Law 92–463, as amended (5 U.S.C. Appendix 2), which sets forth standards for the formation and use of advisory committees.


Sherry Glied,

Assistant Secretary for Planning and Evaluation.

[FR Doc. 2011–3009 Filed 2–9–11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Meeting of the Presidential Commission for the Study of Bioethical Issues

AGENCY: The Presidential Commission for the Study of Bioethical Issues, Office of the Assistant Secretary for Health, Department of Health and Human Services.

ACTION: Notice of Meeting.