

the collection of information; and to transmit or otherwise disclose the information. The total annual burden

hours estimated for this Information Collection Request are summarized in the table below.

Total Estimated Annualized burden hours:

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response in hours)	Total burden hours
Rural Outreach Benefits Counseling Grant Program Measures .....	10	1	10	1.5	15
Total .....	10	1	10	1.5	15

HRSA specifically requests comments on: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**Jackie Painter,**  
 Director, Division of the Executive Secretariat.  
 [FR Doc. 2015-13088 Filed 5-29-15; 8:45 am]  
**BILLING CODE 4165-15-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:

*Ryan Asherin, Oregon Health Authority:* Based on the report of an investigation conducted by the Oregon Health Authority (OHA) and analysis conducted by ORI in its oversight review, ORI found that Ryan Asherin, former Surveillance Officer and Principal Investigator, OHA, Public Health Division engaged in research misconduct in research supported by the Centers for Disease Control and Prevention (CDC) Emerging Infections Program Grant 5U01CI00306-05.

ORI found that the Respondent engaged in research misconduct by falsifying and/or fabricating data that were included in the CDC research record, a manuscript submitted to *JAMA Intern Med* in January 2013, a published CDC report (*CDC Morbidity and Mortality Weekly Report* 61(09):157-162, March 2012), and presentations in

2012 to CDC and at the 11th Biennial Congress of the Anaerobe Society.

ORI found that the Respondent falsified and/or fabricated fifty-six (56) case report forms (CRFs) while acquiring data on the incidence of *Clostridium difficile* infections in Klamath County, Oregon. Specifically, the Respondent (1) fabricated responses to multiple questions on the CRFs for patient demographic data, patient health information, and *Clostridium difficile* infection data, including the diagnoses of toxic megacolon and ileus and the performance of a colectomy, with no evidence in patient medical records to support the responses; and (2) falsified the CRFs by omitting data on the CRFs that clearly were included in patient medical records.

Mr. Asherin has entered into a Voluntary Settlement Agreement (Agreement) and has voluntarily agreed for a period of two (2) years, beginning on May 12, 2015:

(1) To have his research supervised; Respondent agrees that prior to 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 agrees that he will 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 him must 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; and

(3) to exclude himself 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 FURTHER INFORMATION CONTACT:**  
 Acting Director, Office of Research Integrity, 1101 Wootton Parkway, Suite 750, Rockville, MD 20852, (240) 453-8200.

**Donald Wright,**  
 Acting Director, Office of Research Integrity.  
 [FR Doc. 2015-13054 Filed 5-29-15; 8:45 am]  
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Meeting of the 2015 Hurricane Sandy Conference: Translating Research Into Practice**

**AGENCY:** Department of Health and Human Services, Office of the Secretary.  
**ACTION:** Notice.

**SUMMARY:** The Department of Health and Human Services (HHS) Office of the Assistant Secretary for Preparedness and Response (ASPR) is hereby giving notice that ASPR will convene a Hurricane Sandy Conference: Translating Research into Practice public meeting on August 10-11, 2015. The purpose of the meeting is to broadly share, with interested stakeholders, outcomes of Hurricane Sandy recovery science research and training projects awarded under ASPR FOAs EP-HIT-13-001 and EP-HIT-14-001, Centers for Disease Control and Prevention (CDC) FOAs TP13-001 and OH13-002, and National Institute of Environmental Health Sciences (NIEHS) FOAs RFA-ES-13-008 and NOT-ES-13-003. Meeting participants will discuss opportunities to build a community of practice around Hurricane Sandy recovery research and the path forward for translating Hurricane Sandy recovery science research results into practice; highlight Hurricane Sandy recovery science grants as a model for disaster research science preparedness; and demonstrate the benefit of