

Examining the Impact of Targeted Training and Technical Assistance Efforts on the Implementation of  
Comprehensive Cancer  
March 23, 2016

**Institutional Review Board**

***Agreement to Comply with Human Subject Protection Requirements***

The following project has been found by the Institutional Review Board (IRB) to be in compliance with the human subject protection requirements as specified in 45 CFR 46.

**Project Title:** Examining the Impact of Targeted Training and Technical Assistance Efforts on the Implementation of Comprehensive Cancer  
**Principal Investigator/Project Director(s):** Sarah O'Dell  
**ICF Project Number:** 121772.0.009  
**Approval Date:** March 23, 2016  
**Next Continuous Review Date:** **March 23, 2017**

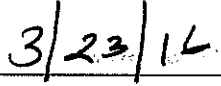
As the responsible principal investigator/Project Director for this project, I agree to adhere to the human subject protection procedures that were approved by the IRB and to inform the chair of the IRB when any changes are made in the approved procedures. The approved procedures include all of the following:

- Subject selection and recruitment procedures
- Data collection procedures
- Informed consent procedures
- Protection of privacy and confidentiality procedures
- Data security procedures
- Additional safeguards specified by the IRB.

If you have any questions regarding changes in procedures that are subject to IRB review, please contact the IRB Chair, Janet D. Griffith ([Janet.Griffith@icfi.com](mailto:Janet.Griffith@icfi.com)), to discuss your concerns.

Also, as the responsible principal investigator or project director, I agree to cooperate with the IRB continuous annual review(s) of this project. Several weeks prior to the next annual review date listed above, the IRB Administrator will send the IRB Project Continuous Review Form or identify where to obtain the form, to complete and submit to the IRB before the annual review date. The purposes of the IRB Project Continuous Review Form are 1) to provide the IRB with updated information on the procedures used to protect the human subjects who are involved in this project, and 2) to help the IRB determine if the project is in compliance with the requirements in 45 CFR 46.

  
\_\_\_\_\_  
(signature)

  
\_\_\_\_\_  
(date)

Please email an original signed copy of this form to the IRB at [IRB@icfi.com](mailto:IRB@icfi.com). A copy of the signed form should also be maintained with your study files.

(Revised-07/18/2014)