**Attachment 12**

**ICF Macro IRB Approval**

**Institutional Review Board Findings Form**

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| --- |
| **Name of Project Director(s):** Michelle Revels**Title of Project:** Impact Evaluation of CDC’s Colorectal Cancer Control Program-annual review**ICF Project Number:** 635243.0.007.00.000 |
| **Type of Review:** New X Renewal |
| **Findings of the Board:** Project is exempt from IRB review (Please see IRB exemption form) X Project complies with all of the requirements of 45 CFR 46, "Protection of HumanSubjects" Project does not comply with all of the requirements of 45 CFR 46 |
| **Project Approved Until:** September 30, 2015**Next Annual Review Date:** November 22, 2013 |
| October 30, 2012*Chair, Institutional Review Board Date* |

(Revised-06/26/12)

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**ICF**

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: Impact Evaluation of CDC's Colorectal Cancer Control

Program-annual review.

PrincipalInstigator/Project Dlrector(s): Michelle Revels

ICF Project Number: 635243.0.007.00.000

Approval Date:

Next Continuous Review Date:

October 30, 2012

November 22, 2013

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), 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

quirem ts in 45 CFR 46.

*ttl*

Please email an original signed copy of this form to the IRB at IRB@icfi.com. A copy of the signed form should

also be maintained with your study files.

(Revtsed-()6/26112)

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