

Attachment A-6: RTI Institutional Review Board Approval



IRB ID Number: 13336

Office of Research Protection
Institutional Review Board Notice of Approval
Federalwide Assurance No. 3331

Title of Study: Using Conjoint Analysis to Understand Individual Decisions about Health-Risk Behaviors and Cancer Screening

RTI Project Number: 0211965.025.001 RTI Proposal Number (if no Project Number)

Project Leader: Carol Mansfield

Project Team Member Contact (if different from Project Leader):

Source of Funding for this Study: CDC

Date Submitted to IRB: July 22, 2013

Level of Review (check one):

Full , IRB Meeting Date: _____

Expedited , category: **7: Behavioral - surveys, focus groups, etc.**

Type of Review (check one):

Preliminary review (The grant application/contract proposal and protocol submitted to the IRB are in concordance (45 CFR 46.103(f)). **Do not involve human subjects or data until pretest or full study is approved.**)

Amendment, describe:

Add study site(s): _____

Pretest/Pilot Test _____

Full Implementation _____

Renewal

Study Closure

IRB Approval of Special Conditions (check all that apply to this review):

Waiver of Signed Informed Consent/Parental Permission

Waiver of elements of Informed Consent or requirement for Informed Consent/Parental Permission

Participation of Pregnant Women (**Worksheet B** submitted by project team)

Participation of Prisoners (**Worksheet C** submitted by project team)

Participation of Prisoners in DHHS-funded studies (OHRP acknowledgement required)

Participation of Minors (**Worksheet D** submitted by project team)

IRB Agreement of Nonsignificant Risk Device Study Determination

HIPAA Waiver of Authorization

Please note the following requirements:

- If **unexpected problems** or **adverse events** occur, the project team must notify the IRB.
- If there are **changes** in study procedures or protocol or any data collection materials (brochures, letters, questionnaires, etc.) the project team must notify the IRB before they are implemented.
- The project team is required to apply for **continuing review** as long as the study is active, which includes participation of human subjects or possession of human data or specimens.

Expiration Date of IRB Approval: May 8, 2014

(No human subjects research can occur after this date without continuing review and approval.)

July 26, 2013

Signature - IRB Member or Chair

Date of IRB Approval

Ina Wallace, PhD

Name - IRB Member or Chair (print or type)

Copy sent to project leader on: 8-9-13

Entered into MIS

OHRP acknowledgement received for participation of prisoners in DHHS-funded studies on: _____



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RTI Project Number: 0211965.025.001 RTI Proposal Number (if no Project Number)

Project Leader: Carol Mansfield

Project Team Member Contact (if different from Project Leader):

Source of Funding for this Study: CDC

Date Submitted to IRB: November 13, 2013

Level of Review (check one):

Full [], IRB Meeting Date: _____

Expedited [x], category: M: Minor changes in approved research

Type of Review (check one):

[] Preliminary review (The grant application/contract proposal and protocol submitted to the IRB are in concordance (45 CFR 46.103(f)). Do not involve human subjects or data until pretest or full study is approved.)

[x] Amendment, describe: revisions to survey questions

[] Add study site(s): _____

[] Pretest/Pilot Test _____

[] Full Implementation _____

[] Renewal

[] Study Closure

IRB Approval of Special Conditions (check all that apply to this review):

- [] Waiver of Signed Informed Consent/Parental Permission
[] Waiver of elements of Informed Consent or requirement for Informed Consent/Parental Permission
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Ina Wallace

November 13, 2013

Signature - IRB Member or Chair

Date of IRB Approval

Ina Wallace, PhD

Name - IRB Member or Chair (print or type)

[x] Copy sent to project leader on: 11-14-13

[] Entered into MIS

[] OHRP acknowledgement received for participation of prisoners in DHHS-funded studies on: _____