

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

Title of Study: Evaluating Knowledge and Practices Related to Safer Use of Medications During Pregnancy

RTI Project Number: 0214176.000.001 RTI Proposal Number (if no Project Number)

Project Leader: Molly Lynch

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

Source of Funding for this Study: CDC

Date Submitted to IRB: July 30, 2014

Level of Review (check one):

Full , IRB Meeting Date: \_\_\_\_\_

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

Type of Review (check one):

Preliminary review (For DHHS grants where RTI is prime, 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: 08-07-2015

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



\_\_\_\_\_  
Signature - IRB Member or Chair

08-07-2014

\_\_\_\_\_  
Date of IRB Approval

Ina Wallace, Ph.D.

\_\_\_\_\_  
Name - IRB Member or Chair (print or type)

Copy sent to project leader on

Entered into MIS

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