

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

**Title of Study:** Survey of Prison Inmates  
**RTI Project Number:** 0213181 **RTI Proposal Number** (if no Project Number)  
**Project Leader:** Tim Smith  
**Project Team Member Contact** (if different from Project Leader):  
**Source of Funding for this Study:** Bureau of Justice Statistics  
**Date Submitted to IRB:** November 16, 2015

**Level of Review** (check one):

Full , IRB Meeting Date:

Expedited , category: **M: Minor changes in approved research**

**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: revisions to sample size, questionnaire, consent text and consent processes

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

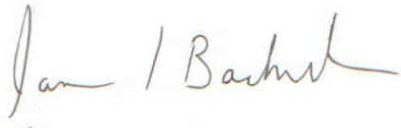
HIPAA Waiver of Authorization

**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:** April 22, 2016

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



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

**November 23, 2015**

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

Jamia Bachrach, JD

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

Copy sent to project leader on: November 23, 2015

Entered into MIS

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