IRB ID Number: 13011

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

**Title of Study:** Medical Device Decision Analysis: A Risk Tolerance Pilot Study
**RTI Project Number:** 0302907 **RTI Proposal Number** (if no Project Number)
**Project Leader:** Juan Gonzalez
**Project Team Member Contact** (if different from Project Leader): Valerie Tower
**Source of Funding for this Study:** FDA
**Date Submitted to IRB:** April 5, 2012
**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 with regard to the scientific conduct of the study, informed consent content, and all other issues pertaining to the protection of human subjects. (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: January 27, 2013**
(No human subjects research can occur after this date without continuing review and approval.)

Ina Wallace

**Signature - IRB Member or Chair**

**April 9, 2012**
**Date of IRB Approval**

Ina Wallace, PhD
**Name - IRB Member or Chair (print or type)**

☒Copy sent to project leader on: 4-10-12
☐Entered into MIS
☐ OHRP acknowledgement received for participation of prisoners in DHHS-funded studies on:_____

Office of Research Protection, Institutional Review Board
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