Clinical Practice Guidelines

Administration Evaluation: The WCMC follows applicable institutional guidelines on harmonization good

The WCMC expands the EPs supported in V-A, to all research reviewed by the WCMC, regardless of the

The Chariperson (or designee) does not have a potential for conflict of interest on this study.

- Survey – Spanish and English version
- Interventional Schedule – Spanish and English version
- Informed Consent – Spanish version (on version date)
- Informed Consent – English version (version date 04/27/2011)
- Informed Consent – Revised date 04/28/2011
- Intended Process Flow: (WCMC Record date 01/2012)

The approval

The above referenced research study has been given approval for the period of 05/05/2011 to 05/05/2012. The approval

Title: Expired category Research Study

RE: May 01, 2011

DATE: From: To:

Ryan J. Cooper, Graduate Student, ECU, Department of Geography

EAST CAROLINA UNIVERSITY
Office 272-447-9291 • Fax 252-447-2344 • wwww.ecu.edu
11-09 Moody Medical Science Building 600 Moye Boulevard Greenville, NC 27834
Acuerdo de Compromiso

Este acuerdo es una declaración de confidencialidad entre la empresa y el investigador. Se expresa en forma de contrato para garantizar la confidencialidad de la información que se intercambia.

La empresa calibra el acuerdo en función de la naturaleza y el contenido de la información comparte. Los términos y condiciones del acuerdo incluyen restricciones de competencia, uso y divulgación de la información.

El investigador, por su parte, se compromete a respetar estrictamente las condiciones del acuerdo. En caso de incumplimiento, puede ser sancionado con medidas disciplinarias o legales.

Este acuerdo se firmará en fecha [fecha de firma] y entrará en vigor a partir de dicha fecha.
INFORMED CONSENT DOCUMENT

\textbf{INTRODUCTION}
For Reporting Unanticipated Problems Involving Risks to Participants or Others, you are required to report to the IRB all unanticipated problems that have occurred in your research within the time frame specified in the UMCIRB rule.

As Principal Investigator, you are required to report to the IRB all unanticipated problems that affect Participants or Others.

Reporting Unanticipated Problems to the IRB That Affect Participants or Others

Inform IRB of the changes as soon as possible via a protocol deviation form. If these changes are to eliminate or minimize an immediate apparent hazard to the participant, the IRB must immediately undertake to prevent a hazard to the participant. If there are no changes, the protocol deviation form must be submitted to the IRB.

Required Approval for Any Changes to IRB-Approved Research

You have any questions regarding your role or requirements with continuing review, please contact the UMCIRB office at 252-744-2914. If meeting submission deadlines, please contact the UMCIRB office at 252-744-2914 for assistance.

Review for Research Involving Human Subjects, Research on Research Participants, and Graduate Studies. Your approval will be reported to the Vice Chancellor for Research and Graduate Studies. This information is required to summarize all research activities that are allowed to continue or cease within the period of your approval. You must submit this form even if there has been no activity, no participants enrolled or...

Provide a retrospective approval during a period of 12 months. If the IRB approved period was less than 12 months, the requirement to report any research activities outside of the approved period will still be in effect.

As Principal Investigator, you are required to submit a continuing or final review form to...

Continuing and Final Review Obligations

***********IMPORTANT INFORMATION***********