

IRB Signature

Signature Jean Taylor Woodbury By MB 10/29/2014
 Jean Taylor-Woodbury, RN, MS, ANP-BC, Chair Mona Bosch, CIP Date

Review and Approval Information

E&I Study Number 14133 - 01 Approval Date Wednesday, October 22, 2014
 Review Process Expedited 5 & 6 Expiration Date Wednesday, October 21, 2015 at 11:59 PM

This document certifies the IRB's approval, per 34 CFR 97 including Subpart D Section 404, of the bolded items identified under "Documents Approved" to be conducted by the named Principal Investigator.

A waiver of the requirement for documentation of informed consent is granted according to 34 CFR 97.117(c)(2).

NOTE: Subjects must be asked for their consent using the most recently approved, stamped version(s). All IRB approved consent documents are version controlled and may not be modified in any way without prior IRB approval. Use of an unapproved document may constitute non-compliance.

Study

Understanding the Relationship between
 Implementation of Texas House Bill 5 and Algebra II
 Completion Rates in High Schools

Client SEDL
 Sponsor U.S. Department of Education

Grant Number and/or Title

ED-IES-12-C-00012 Regional Education Laboratory Southwest

Principal Investigator

Ginger L. Stoker, PhD
 E&I PI Number 16708 - 001

Address

SEDL
 4700 Mueller Blvd.
 Austin, TX 78723

Performance Sites

SEDL, 4700 Mueller Blvd., Austin, TX 78723

Sub-Investigator(s), Approval Date(s)

Lynn T. Mellor, PhD, 10/23/2014
 Verónica Ruiz de Castilla, MA, PhD, 10/23/2014
 Michael Vaden-Kiernan, PhD, 10/23/2014

Documents Approved

Documents Approved	Document #	Version	Date
Protocol		4.10	10/23/2014
Electronic Consent Letter	E&I 10/29/2014		10/23/2014
Survey			10/23/2014

Stipulations of Approval

1. No subjects may be involved in any study procedure prior to the IRB approval date or after the expiration date. Investigators and sponsors are responsible for initiating Continuing Review proceedings.
2. All protocol modifications must be IRB approved prior to implementation. This includes any addition

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E&I Business Office
 14400 East 42nd Street, Suite 240
 Independence, MO 64055
Phone (816) 421-0008
 Fax (816) 356-2227



E&I West Coast Board
 5710 Paradise Drive, Suite 11
 Corte Madera, CA 94925
Phone: (415) 485-0717
 Fax: (415) 485-0328

or change of recruitment materials, change of investigator, or performance site address.
(Exception: If necessary to eliminate apparent immediate hazard to subjects.)

3. Report to E&I within five working days of learning if any of the following occur:
 - Unanticipated problems involving risk to human subjects or others;
 - Unanticipated Serious Adverse Events and Safety Reports;
 - Protocol deviations, violations, and exceptions that impact subject welfare or safety or study integrity including changes intended to reduce immediate risk to subjects;
 - Use of an investigational product in an emergency situation; and
 - Claims for compensation or for medical care for research-related injury.
4. Advertising and recruitment materials must be approved by E&I prior to use or publication.

END

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