

**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 Client SEDL  
 Implementation of Texas House Bill 5 and Algebra II Sponsor U.S. Department of Education  
 Completion Rates in High Schools

**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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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.

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**END**

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