

Institutional Review Board Office

Suite 203, MC-419 528 East Green Street Phone: 217.333.2670 Fax: 217. 333.0405

E-mail: <u>irb@uiuc.edu</u>
Web: <u>www.irb.uiuc.edu</u>

Research Amendment

For Submitting Changes to Previously Approved Human Subjects Research

All applications must be completed, signed by the RPI, and submitted either electronically or single-sided hard copy. Please - No Staples

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|---|--|--|--|---|---|--|
| All modifications to human subjects | s resear | ch must be reviewed | and ar | proved prior to imp | Form Version 1.05 lementation. | |
| All modifications to human subjects research must be reviewed and approved prior to implementation. Minor modifications Minor modifications to previously approved projects include those that do not alter the risk-benefit assessment for the research. Examples include changes in the investigators; minor changes in the consent form(s), recruiting materials, measures, or procedures; minor changes in compensation, time of participation, or subject recruitment; or the use of a new site that is not materially different from a previously approved site. Minor modifications may also include changes to other parameters, whereby the investigator provides the subjects with more accurate information as a result of additional experience with the protocol. | | | | | | |
| Major modifications Major modificat activities not previously approved; or to being of participants (including the los alteration of the risk-benefit assessment measures that significantly differ from informed consent process, and change | hat involues of confi ent for the those cu | ve an increased level identiality); or that inverse research. For examperently approved, cha | of risk to olve a d ple, add nging in | o the physical, emotic ecreased benefit; or t ing a new subject pop clusion or exclusion o | onal, or psychological well- hat otherwise result in oulation, adding new criteria, changing the | |
| 1. DATE THIS REPORT WAS COMPLETED 2. RESPONSIBLE PROJECT INVESTIGAT | <u> </u> | | AMEN | IDMENT NUMBER (ST. | ART WITH 01): 10015 | |
| Last Name: Winkelmes | <u> </u> | First Name: Mary-An | n | Academic Degree(| s): PhD. MA | |
| Dept. or Unit: Provost | | Office Address: 580 II | | 1 | Mail Code: MC 317 | |
| Street Address: 807 S Wright St | | City: Champaign | | State: IL | Zip Code: 61820 | |
| Phone: 217 244 5108 | Fax: 21 | 7 265 4183 | | E-mail: mawink@illir | <u> </u> | |
| UIUC Status: Non-visiting member of (Mark One) ☐ Faculty ☐ Staff | | | | | | |
| 3. PROJECT TITLE Illinois Initative on Transparency in Le | arning ar | IRB PROTOCOL nd Teaching | NUMBE | R: 10015 | | |
| 4. MAJOR OR MINOR MODIFICATION? I | the RPI | 's judgment, which ca | tegory | of modification is this | ? | |
| ☐ Minor ☐ Major ☒ Uncertain | | | - / | | | |
| 5. REVISED MATERIALS: For revision procedures, measures, etc.), or to add rapplication for Exemption incorporating modification of consent forms, assent for | new proc the revis | edures that were not place in the decirion of the decirions as appropriate the decirion of the | previoùs rrougho | sly approved, please out the form. Amendm | resubmit the IRB-1 or | |
| PLEASE SUPPLY THE FOLLOWING with this Research Amendment: a) A marked up version of the IRB-1 Application or Application for Exemption and any modified attachments or consent documents. NOTE: If your computer does not allow "strike-throughs" or other editing on the IRB-1 or Application for Exemption, it is acceptable to cross off deleted sections with a pen and use a highlighter to emphasize changes. b) The entire IRB-1 Application or Application for Exemption reflecting the revisions. Revised consent documents and other relevant attachments that have changed as a result of the amendment Mark One: Changes marked versions and final versions are: Attached Will Follow. | | | | | | |
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6. DESCRIBE THE AMENDMENT. Describe the requested change(s) and clearly reference materials submitted with this form. Provide a clear rationale for the proposed change(s). Explain whether the risk-benefit assessment for the research is likely to change as a result of the proposed amendment(s). Justify changes that will affect risks, benefits, informed consent, inclusion or exclusion criteria, the subject population(s), research sites, or the confidentiality of private, identifiable subject information. The risk-benefit assessment for this research is not likely to change as a result of any of the proposed changes. Minor adjustments have been made to the self-declared nationality question on the online survey (attached). The change was made in response to analysis provided by Illinois STEM Initiative staff. The existing information and consent statement for participating students was revised in minor ways to agree with new templates on IRB website, and was joined by an additional, similar information and consent statement for participating instructors. The additional information and consent statement for instructors will allow for future reporting of anonymous data in the aggregate about how instructors describe their teaching when they register online to participate and when they participate in any optional conversations online via Elluminate, via email, or even in person or on the telephone via oral interview. Elisa Mustari (ISTEM) and Judith Sunderman (College of Engineering) join the project as investigators. The attached documents include: 1) approved online survey form indicating in red the proposed change to the self-declared nationality question (#31) 2) original IRB-1 application with amendments indicated by "track changes" function; 3) clean copy of amended IRB-1 application 4) proposed interview protocol for possible future interviews with instructors 5) revised information and consent statement for students with changes indicated by "track changes" function and new information and consent statements for instructors 6) online registration form for instructors (beginning of term and end of term) If additional Item 6 information is attached, check here: 7. INVESTIGATOR ASSURANCES The original, inked signature of the Responsible Project Investigator is required before this form can be processed. Other investigators are also responsible for these assurances and are encouraged to sign. Neither stamps nor proxy signatures are accepted in this section. I certify that the information supplied in this form, with attachments, is complete and correct, that the modified protocol has not yet been used with any human subject, and that it will not be implemented until IRB approval has been obtained. NOTE: The signature of the RPI must be submitted before IRB Review (scanned or faxed signatures are acceptable). Responsible Principal investigator Date Investigator Date

Investigator

Date

Investigator

Date



University of Illinois at Urbana-Champaign

IRB Application for Exemption

Institutional Review Board Suite 203, MC-419 528 East Green Street

Phone: 217.333.2670 Fax: 217.333.0405 irb@uiuc.edu www.irb.uiuc.edu

All forms must be completed, signed by the RPI, and submitted by FAX. Email, or single-sided hard copy.

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| | | Verson 1.01 | | |
| Project Title: Illinois Initiative on Tra | nsparency in Learning and Teaching | 5 | | |
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| | . The RPI must be a non-visiting memb | | | |
| will serve as project supervisor at UIUC. | | | | |
| other campuses may not serve as RPI, bu | | | | |
| Last Name: Winkelmes | First Name: Mary-Ann | Academic Degrees: PhD, MA, | | |
| Dept. or Unit: Office of the Provost an | dOffice Address: 534 Illini Union | Mail Code: 317 | | |
| Chancellor for Academic Affairs | Bookstore,807 S. Wright St | | | |
| Street Address: 807 S. Wright St | City: Champaign | Zip Code: 61820 | | |
| Phone: 217 244-5108 | Fax: 217 265-4183 | E-mail: mawink@illinois.edu | | |
| UIUC Status (please mark one): Non-vis | iting member of Faculty | ⊠ Staff | | |
| institutions. Include all persons who will consent process, data collection, data ana | | | | |
| Last Name: Mustari | First Name: Elisa | Academic Degrees: Ph.D. | | |
| Dept. or Unit: I-STEM | Office Address: 704 S Sixth St | Mail Code: MC 308 | | |
| Street Address: 704 S Sixth St | City: Champaign, IL | Zip Code: 61820 | | |
| Phone: (217) 333-0291 | Fax: | E-mail: | | |
| | | emustari@illinois.edu | | |
| · · · · · · · · · · · · · · · · · · · | | tudent | | |
| ☐ Visiting Scholar | Non-UIUC Affiliate of (Institution | 1) | | |
| | | | | |
| | a list of Additional Investigators, if appl | | | |
| Judith Sunderman, Ph.D. | | | | |
| College of Ed EOL M/C 708 | Champaign, IL 61820 | | | |
| jsunderm@illinois.edu | Grampary 12 Globs | | | |
| 1.3 Please review the 6 categories of exemples | ption listed below and indicate the catego | ry or categories that apply to your | | |
| research. (Note: Exemptions do NOT appl | | | | |
| cognitively impaired or persons who are ed | | - | | |

- 1. Research conducted in established or commonly accepted educational settings, involving normal educational practices, such as: research on regular and special education instructional strategies, or research on the effectiveness of or the comparison among instructional techniques, curricula, or classroom management methods.
- 2. Research involving the use of educational tests (cognitive, diagnostic, aptitude, achievement), survey procedures, interview procedures, or observation of public behavior, unless:
 - information obtained is recorded in such a manner that human subjects can be identified, directly or through identifiers linked to the subjects; and

| any disclosure of the human subjects' responses outside the research could reasonably place the subjects at risk of criminal or civil liability or be damaging to the subjects' financial standing, employability, or reputation. Note: This category does not apply to the following types of research involving children: surveys, interviews, and observations of public behavior when the investigator is a participant in the activities being observed. |
|--|
| 3. Research involving the use of educational tests (cognitive, diagnostic, aptitude, achievement), survey procedures, interview procedures, or observation of public behavior that is not exempt in category 2 above if: human subjects are elected or appointed public officials or candidates for public office; <u>or</u> federal statute(s) require(s) without exception that the confidentiality of the personally identifiable information will be maintained throughout the research and thereafter. |
| 4. Research involving the collection or study of existing data, documents, records, pathological specimens, or diagnostic specimens, if these sources are publicly available or if the information is recorded by the investigator in such a manner that subjects cannot be identified, directly or through identifiers linked to the subjects. Note: In order to be eligible for this exemption, all data, documents, records or specimens must exist prior to IRB review and must have been collected for purposes other than the proposed research. (To qualify for an exemption in this category, the proposed research must be strictly retrospective.) |
| 5. Research and demonstration projects that are conducted by or subject to the approval of department or agency heads. The program under study must deliver a public benefit or service (e.g., Social Security Act or Older Americans Act). Such research or demonstration projects must be conducted pursuant to specific federal statutory authority; there must be no statutory requirement that the project be reviewed by an Institutional Review Board (IRB); and the project must not involve significant physical invasions or intrusions upon the privacy of participants. |
| ☐ 6. Taste and food quality evaluation and consumer acceptance studies: if wholesome foods without additives are consumed; or if a food is consumed that contains a food ingredient at or below the level and for a use found to be safe, or agricultural chemical or environmental contaminant at or below the level found to be safe by the Food and Drug Administration (FDA) or approved by the Environmental Protection Agency (EPA) or the Food Safety and Inspection Service of the US Department of Agriculture (USDA). |
| If the proposed research does not qualify in any of these categories, please complete the IRB 1 application form at www.irb.uiuc.edu . |
| 2. Research Summary. In layman's language, please summarize the objectives and significance of the research. |

2. Research Summary. In layman's language, please summarize the objectives and significance of the research The Illinois Initiative on Transparency in Learning and Teaching provides a structure for faculty who wish to contribute to a large and significant research project on students' learning, while relying on others to provide the education-research expertise and administrative support. It also brings teachers and students into dialogue about the processes of teaching and learning.

The premise of the research is that students learn more deeply and retain information longer when they understand how and why they are learning course content in particular ways. Further, students' critical thinking abilities and their capacity to monitor and take responsibility for their own learning will increase, both during "transparent" courses and long afterward. Faculty satisfaction and effectiveness should increase as a result of this purposeful dialogue about their students' particular learning styles and procedures. Students' satisfaction with their courses and their teachers will likely be enhanced, along with their performance in "transparent" courses.

Ultimately, this research will identify which small changes to teaching and learning practices produce the greatest beneficial impact on students' learning, with results specific to: the past experience of the student, the size of the course, the level of the course (beginning college through advanced degree) and the discipline. Longer-term results may include higher retention and graduation rates for undergraduate students, including community college students who transfer into four-year institutions, and greater participation of diversely prepared students in Masters and PhD degree programs.

| Please check here and attach additional Research Summary information, if applicable. |
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| 3. Participants. Describe who will participate in this research and how these persons will be recruited. |
| Participants will be students enrolled in the courses of voluntarily participating instructors at the |
| University of Illinois, University of Chicago, and other institutions, including Semarang State |
| University, Gustavus Adolphus College, Franklin University, El Colegio de Mexico, University of |
| Glasgow, University of Sunderland. |
| An amendment will be filed to indicate participating courses/institutions each year. |
| |
| Please check here and attach additional Participant information, if applicable. |
| 4. Research Procedures. Specifically describe what the participants will do and where the activities will take |
| place. Outline the approximate dates and durations for specific activities, including the total number of |
| treatments, visits, or meetings required and the total time commitment. Please include a copy of each of your |
| measures as attachments. |
| All potential student participants will receive an electronic invitation from the principal investigator |
| and/or course instructor during the last week their participating course is in session. Clicking on a link |
| provided in the invitation will lead a student to an Information and Consent form (attached). Clicking "I |
| agree" will lead to the online survey. Students who agree to participate will complete an online survey |
| (attached) of approximately thirty multiple-choice questions about their perceptions of their learning |
| experience in the course. The survey takes approximately four or five minutes to complete. |
| Participating instructors register online (registration form attached) to indicate their voluntary |
| participation. Information and Consent forms for participating instructors (attached) allow for future |
| reporting of anonymous data in the aggregate about how instructors describe their teaching on the |
| registration form, or in any optional conversations online, via Elluminate, via email, or even in person |
| or on the telephone via oral interview. |
| |
| At the three-year, five-year and ten-year marks (when possible), some of these student and instructor |
| participants will receive an electronic invitation to complete online surveys or oral interviews about the |
| longer-term impact of their past participation on their learning and/or teaching practices. The follow- |
| up surveys will consist of approximately fifteen questions (to be filed by the principal investigator in an |
| amendment) and will take approximately six minutes to complete. The follow-up interview protocol |
| |
| (attached) will take approximately forty-five minutes. |
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| Please check here and attach additional Research Procedures information, if applicable. |

5. Data Collection. Please explain how confidentiality will be maintained during and after data collection. If appropriate, address confidentiality of data collected via e-mail, web interfaces, computer servers and other networked information.

Data from online surveys will be stored on the Swanlund Server, maintained by Swanlund System Services in the Provost's Office at the University of Illinois, and on a flash drive kept in a locked drawer in the office of the Campus Coordinator for Programs on Teaching and Learning, Office of the Provost and Vice Chancellor for Academic Affairs, at the University of Illinois.

Students' anonymity will be preserved. Students' identities will never be tracked: no key or other identifier will link students' answers with their identity. Data from the survey will be preserved for the duration of this ten-year study (2009-2019). Students' data will be shared only in aggregate form. Instructors will not receive students' data until after course grades are submitted. Instructors' identities will be known only to the investigators, who will keep these identities confidential. Instructors' names will not be used in any reports, presentations or publications resulting from the Transparency Initiative, nor will names be shared with institutions' Review Boards or other administrators. Instructors' data will be shared only anonymously in aggregate form. Information and Consent Statements are attached.

Please check here and attach additional Data Collection information, if applicable.

6. Consent Process Describe when and where voluntary consent will be obtained, how often, by whom, and from whom. Attach copies of all consent forms (as well as assent forms for those under age 18 if any). All potential student participants will receive an electronic invitation from the principal investigator and/or course instructor during the last week their participating course is in session. Clicking on a link provided in the invitation will lead a student to an Information and Consent form (attached). Clicking "I agree" will lead to the online survey. Students who agree to participate will complete an online survey (attached) of approximately thirty multiple-choice questions about their perceptions of their learning experience in the course. The survey takes approximately four or five minutes to complete. Participating instructors register online (registration form attached) to indicate their voluntary participation. Information and Consent forms for participating instructors (attached) allow for future reporting of anonymous data in the aggregate about how instructors describe their teaching on the registration form, or in any optional conversations online, via Elluminate, via email, or even in person or on the telephone via oral interview.

At the three-year, five-year and ten-year marks (when possible), some of these student and instructor participants will receive an electronic invitation to complete online surveys or oral interviews about the longer-term impact of their past participation on their learning and/or teaching practices. The follow-up surveys will consist of approximately fifteen questions (to be filed by the principal investigator in an amendment) and will take approximately six minutes to complete.

| Please check here and attach additional Consent Process information, if applic | |
|---|---------------------------------------|
| 7. Dissemination of Results. What is (are) the proposed form(s) of dissemination (e.g. | |
| academic paper, conference presentation, sharing within the industry or profession, etc.) Results of students' survey responses in aggregate form (with no individuals iden | |
| with instructors after grades are submitted and will likely be disseminated via con | · · · · · · · · · · · · · · · · · · · |
| white papers, journal publications and perhaps even a book publication. Data from | - |
| instructors' responses to the online registration form and optional communications onlin | · · · · |
| similarly disseminated (with no individuals identified) in aggregate form. | • |
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| Dloggo shook how and stands additional Discoving and CD state in C. | if amplicable |
| Please check here and attach additional Dissemination of Results information. | , if applicable. |
| | |
| 8. Individually identifiable information. Will any individually identifiable information | on, including images of |
| subjects, be published, shared, or otherwise disseminated? Please mark the appropriate | box below. |
| | |
| ☐ Yes No | |
| | |
| Note: If yes, subjects must provide explicit consent or assent for such dissemination. P | rovide appropriate |
| options on the relevant consent documents. | |
| | |
| 9. Funding Information. | |
| Is your research funded or is there a pending funding decision? \(\subseteq \text{Yes} \text{ No} \) | |
| | |
| If "yes", please indicate the funding agency here: Funding will eventually be sought. | |
| be filed by the principal investigator to indicate which agencies have received a f | funding proposal. |
| Diago submit a convertible for diagonary | |
| Please submit a copy of the funding proposal. | |
| | |
| 10. Expected Completion Date: September, 2019 | |
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| INVESTIGATOR ASSURANCES | |
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| I certify that the project described above, to the best of my knowledge, qualifies as an exe | |
| agree that any changes to the project will be submitted to the Institutional Review Board | |
| to implementation. I realize that some changes may alter the exempt status of this project signature of the RPI is required before this application may be processed (scanned or faxed states). | |
| acceptable). | nguaturos are |
| | |
| Mary-Ann Winkelmes | 09/17/2009 |
| Responsible Project Investigator | Date |