#406000001 - Beginning College Survey of Student Engagement

Protocol Information

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<th>Status</th>
<th>Approval Date</th>
<th>Continuing Review Date</th>
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<td>Approved</td>
<td>May 30, 2023</td>
<td>Jun 21, 2020</td>
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Expiration Date: --
Initial Approval Date: Oct 19, 2018
Initial Review Type: Expedited

Feedback

Approval Comment

Amendment A033

This research is approved under the following expedited category:

- Category 7

Protocol Amendment Form

Amendment Request

4000

Select your Protocol Type
Expedited/Full Board
Amendment Number
A033

Select the types of changes being made.
Other changes

Select the appropriate status of the study.
Open to Enrollment – Enrollment continues

Describe the changes being made.
This amendment is in response to F002 as directed by the Board (IRB-01 4/28/23)
Added this statement to the informed consent. "There is a risk of loss of confidentiality - someone outside the study team/institution could get access to your research information from this study. However, security procedures are in place to minimize the risk. We will protect your information and make every effort to keep your personal information confidential."

Why are these changes being made (i.e. what is the rationale for these changes)?
Added the confidentiality clause to the informed consent.

Will any previously enrolled subjects be informed of these changes?
No

Explain why subjects will not be informed.
The project does not have contact information for prior subjects.
General Information

Principal Investigator
Cole, James

Lead Unit
BL-EDUC - EDUCATION

Protocol Title
Beginning College Survey of Student Engagement

Personnel

Person
BrckaLorenz, Allison

Email Address
abrckalo@indiana.edu

Researcher Role
Behavioral/Social Science Researcher - Stage 1 - CITI

Contact Roles

Permissions
Read-Only

Affiliation Type
IU

Training

Behavioral/Social Science Researcher - Stage 1 - CITI 03/21/22 - 03/20/27
## Personnel Attachments

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<td><a href="mailto:brooksjl@indiana.edu">brooksjl@indiana.edu</a></td>
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</table>
Email Address
colejs@indiana.edu

Researcher Role
Principal Investigator (PI)

Home Unit
BL-CPR - CENTER FOR POSTSECONDARY RESEA

Contact Roles
Protocol Editor

Permissions
Full Access

Affiliation Type
IU

Training
Behavioral/Social Science Researcher - Stage 1 - CITI  11/03/22 - 11/03/27

COI Disclosure
Status: Approved

Personnel Attachments

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Name
CV for James Cole

Attachment Type
Person

Dugan, Brendan

Email Address

bjdugan@iu.edu

Researcher Role

Key Personnel

Contact Roles

Permissions

Read-Only

Affiliation Type

IU

Training

Behavioral/Social Science Researcher - Stage 3 - CITI  07/06/22 - 07/05/27

COI Disclosure

Status: Approved

Personnel Attachments

Person

Gonyea, Robert

Email Address

rgonyea@indiana.edu
Researcher Role

Key Personnel

Contact Roles

Permissions
Read-Only

Affiliation Type
IU

Training

COI Disclosure
Status: Approved

Personnel Attachments

Person
Holmes, Bridgette

Email Address
holmesbl@iu.edu

Researcher Role
Other Research Staff

Contact Roles

Permissions
Read-Only
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<td><a href="mailto:anglmill@indiana.edu">anglmill@indiana.edu</a></td>
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<td>Mariano, Gavin</td>
<td><a href="mailto:gmariano@iu.edu">gmariano@iu.edu</a></td>
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Researcher Role

Other Research Staff

Contact Roles

Permissions
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Affiliation Type
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Training

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COI Disclosure
Status: Approved

Personnel Attachments

Person
Priddie, Christen

Email Address
cpriddie@iu.edu

Researcher Role

Other Research Staff

Contact Roles

Permissions
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<td>Sarraf, Shimon</td>
<td><a href="mailto:ssarraf@indiana.edu">ssarraf@indiana.edu</a></td>
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Affiliation Type
IU
Training
Behavioral/Social Science Researcher - Stage 1 - CITI 08/12/20 - 08/11/25
COI Disclosure
Status: Approved
Personnel Attachments
Person
Brandon, Josclynn
Email Address
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Researcher Role
Other Research Staff
Contact Roles
Permissions
Full Access
Affiliation Type
 IU

Training

| Behavioral/Social Science Researcher - Stage 2 - CITI | 12/13/21 - 12/12/26 |

COI Disclosure
Status: Approved

Personnel Attachments

Person
Zhu, Yihan

Email Address
zhuyih@iu.edu

Researcher Role
Other Research Staff

Contact Roles

Permissions
Full Access

Affiliation Type
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Training

| Behavioral/Social Science Researcher - Stage 2 - CITI | 08/13/22 - 08/12/27 |

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**Permissions**

- Full Access

**Affiliation Type**

- IU

**Training**

- Behavioral/Social Science Researcher - Stage 1 - CITI 02/22/21 - 02/21/26

**COI Disclosure**

- Status: Not Disclosed

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Akyuz, Filiz

Email Address
fakyuz@iu.edu

Researcher Role
Other Research Staff

Contact Roles

Permissions
Read-Only
Affiliation Type
IU

Training

COI Disclosure
Status: Not Disclosed

Personnel Attachments

Person
Bacigalupa Albaum, Toni

Email Address
tbacigal@iu.edu

Researcher Role
Other Research Staff

Contact Roles

Permissions
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Affiliation Type
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Training

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**Protocol Type**

**Select Protocol Type**

Expedited
### Participant Information

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### Organizations

#### Organizations List

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### Funding

Will the study be funded, fully or partly by, any of the following sources (this includes pass through funding)? *Select all that apply.*

No external funding

### Conflicts of Interest
Are any of the investigators listed in the personnel section aware of an institutional conflict of interest which could affect or be affected by this research?
No

Do any of the investigators listed in the Personnel section (or their immediate family members) have a significant financial interest which could affect this research?
No

Does the Principal Investigator affirm all investigators listed as personnel on this protocol have agreed to participate in this project, are aware of their status and role, and have been adequately trained to participate in the project?
Yes

Does any research activity in this study present more than minimal risk to human subjects?
No. The research may qualify for Expedited review if all research procedures fall into one of the categories below.

Check all category(ies) which apply to this research.
Category 7 - Research on individual or group characteristics or behavior, or research employing survey, interview, oral history, focus group, program evaluation, human factors evaluation, or quality assurance methodologies
Would identification of subjects and/or their responses reasonably place them at risk for any of the following (check any that apply):
None

B-Lay Summary Research Design

Describe the purpose of this study in lay terms, including research question(s) and hypothesis.
Since 2007, the Beginning College Survey of Student Engagement (BCSSE) has investigated, on an annual basis, entering first-year college students’ high school experiences and their expectations for engagement in educational practices during their first year of college. Starting in 2019, the survey will also include entering transfer and delayed entry students. Most of the items in the BCSSE instrument represent empirically-confirmed “good practices” in undergraduate education. That is, they reflect behaviors by students that are associated with desired outcomes of college. Results from BCSSE are also linked with data from the National Survey of Student Engagement (NSSE) (0709000079R008). Thus, many of the items are adapted from NSSE to allow for valid comparisons by institutions that participate in both projects. The results of the administration of the BCSSE will provide an estimate of how students entering college spent their time in high school and how they expect to be engaged in educationally effective practices in college. The study takes place on each individual college campus that registers to participate. There are two versions of the survey (paper and web). Officials at each campus are responsible for administering the survey in an effective and appropriate manner with respect to human subject compliance.
List and describe all research interactions and/or interventions, including the frequency and duration of procedures, and length of participation for individual subjects.

All participants are asked to complete the BCSSE core survey. The BCSSE survey takes about 15 minutes to complete. Participants only respond to the survey one time. There are three versions of the BCSSE survey instrument. Each version is described below, with any differences between the instruments also explained. The three different core survey versions are: 1. Paper survey in U.S. English 2. Web survey in U.S. English, 3. Web survey for English-speaking students at Canadian institutions. All survey versions are designed to measure the same educational activities consistent with the way these questions are asked at U.S. institutions. In the case of the Canadian version of the survey, some terms were changed and previously approved by IRB to more accurately reflect the Canadian educational system. Three versions of BCSSE survey 1. U.S. English version. [Paper]. BCSSE surveys in a print format are sent to each participating institution for local administration. 2. U.S. English version. [Web]. BCSSE survey in a Web-based format. Students log on using provided institutional codes. 3. English-speaking students at Canadian institutions. [Web-only]. Most survey items are the same as the U.S. version, but the wording was changed in a few cases to reflect cultural differences (e.g., conceptions of race and ethnicity are different) and differences between the two post-secondary systems (e.g., college generally refers to a community college or technical school, so references to college are replaced with university).

Will any non-English study documents be uploaded?
No

Is this research funded by, or has a funding application been submitted to, a federal agency? This includes federal pass-through funding.
No
List inclusion criteria - eligibility criteria for subjects.
Eligible students are defined by each institution. Generally entering first-year, transfer, or delayed-entry students at colleges and universities that register for the annual survey administration.

List exclusion criteria (any criteria which would exclude otherwise acceptable subjects).
Institutions may decide to exclude other accidental subjects. For instance, a university may want to only survey their honors college first year students, therefore excluding all other first year students.

Will subjects be paid for their participation in the study? Payment includes reimbursement of expenses (other than compensation for injury).
Yes

Describe the payment arrangement, including amount and timing of disbursement.
Not all institutions use incentives. Institutions choosing to offer survey incentives are responsible for the arrangement, timing, and disbursement. Incentives for survey participation vary based on institutional decisions about what they believe will be effective for their campus. Some institutions offer a small incentive for each participant, such as a bookstore gift certificate, with values ranging from $5-10.
Justify the proposed payment arrangement described above, specifically why payment does not provide undue influence for subject participation.

Incentives for participation are not allowed that would withhold from students rights and privileges to which they would otherwise be entitled if the survey was not being administered (e.g., holds on registration or housing sign-ups are not allowed). The value of incentives is limited so that non-participation would not be perceived as a hardship by students who would prefer to decline participation.

Will partial payment be provided if the subject withdraws prior to completion of the study?
Yes

Explain the plan for providing partial payment.
Depends on how eligibility for the incentive is determined by the institution. For instance, an institution might determine that students who only partially complete survey are eligible, where another institution might determine that only those that submit the final screen are eligible.

Does this research involve (choose all that apply):
- the STUDY of any of the following products (regardless of FDA approval status). “The study of” means at least one objective of the study is related to obtaining data about the product
- USE of any of the following products which have not been cleared or approved by the FDA for use in the US
- USE of any of the following products for open label extension, treatment, or compassionate use

NONE

This research involves (check all that apply):
None of the Above
Is this research considered a prospective clinical study?
No

Is this community-engaged research?
No

C-Sites and Collaborations

Are there additional locations of research, not already listed?
Yes

Provide the name of the site, including city and state.
About 120 and 150 colleges and universities participate in BCSSE each year. Because the specific list is subject to change, this URL is the best source for current information about participating institutions: http://nsse.indiana.edu/html/participants.cfm

Is any research taking place outside the United States?
Yes

List each country:
Canada (occasionally, not every year)

Are you requesting that IU provide IRB approval for any researchers who are NOT IU affiliates?
No
Is this a multi-center study or multi-site clinical trial?
No

D-Recruitment Methods

Describe how potential subjects will be initially identified.
Each institution determines who is eligible to participate. The institution then choose how to administer the survey, either via email or in a group administration.

Check any of the following sources of information which will be used to identify potential subjects.
Student data
Other

Explain the Other source(s) of information which will be used to identify potential subjects.
enrollment records of each institution. Institutions are solely responsible for identifying potential subjects.
Describe how potential subjects will be initially contacted.
School officials administer the survey to incoming students as part of their orientation or welcoming process. Institutions may promote the survey indirectly through email, newspaper advertisements, flyers, or class announcements that notify students that the survey is being administered and that the results are important for institutional improvement. Staff at each campus will contact the students regarding participation in BCSSE by either an email invitation (see uploaded in Notes and Attachments tab) or the survey will be distributed in a in-person group format. An Informed Consent Statement will be given to students either in paper form (for the paper survey) or via the first screen of the web survey for those completing the online version. Institutions administering in a group format (paper or web), typically administer the survey in a classroom, auditorium, or computer lab. Students are provided the URL (either via their email account or in person) for access to the web survey or the student is provided the paper version of the survey with the informed consent. Instructions for group administration are included in the both the paper and web administration instructions (see uploaded in Notes and Attachments tab). These instructions include, among other things, that "Campus staff administering the survey must not convey any message, implicitly or explicitly, to the student that they are required to complete the survey or that there is any penalty for non-completion. Instructions to the student cannot contradict the informed consent".

Check any of the following recruitment materials which will be used to contact potential subjects.
- Direct Mail/Email
- Flyers/Brochures
- Website
- Social Media
- Other

Explain the Other recruitment material(s) which will be used to contact potential subjects.
Any of those checked above could be used by an institution to build awareness of the survey.
Select any of the following circumstances which apply to this research.
None of the above.

Would participation in this study preclude subjects from participating in other research studies?
No

List and describe (in lay terms) the potential risks to which subjects may be exposed as a result of their participation in the research.
BCSSE collects, stores, and reports back to the institution student self-reported personally identifiable data. Students are explicitly instructed on the paper and web versions of the survey not to report their social security number. BCSSE has programmed the web survey to recognize numbers could be a social number and to delete it from all records. When paper surveys are scanned, any students identifiers can could possibly be a social security is deleted from the records. The only anticipated risk due to participation would be the possible loss of confidentiality of the responses as part of the data distribution process. The consequences of such loss of confidentiality would be minimal due to the nature of the questions asked on the BCSSE survey, which do not cover topics that would generally be considered of a sensitive nature. Survey questions are not of a nature that would tend to embarrass students or place them at risk of physical, psychological, social, or legal harm.
Describe procedures for protecting against, or minimizing, the potential risks listed above. Include any procedures that are already being performed on subjects for diagnostic, treatment, or standard purposes. BCSSE makes every effort to maintain the confidentiality of student data, and the institutions that receive data files with identifiable student data are bound to protect student privacy under the terms of the Family Educational Rights and Privacy Act and other privacy requirements. Please see the next question for how confidentiality will be protected.

Explain how research data will be protected so that only approved persons have access to subjects’ identifiable data (i.e. confidentiality of data).

BCSSE entered into an agreement to license the Qualtrics CX Dashboard. The use of the Qualtrics survey (XM Platform) is already licensed by the Indiana University System. As such, UITS previously completed the Higher Education Cloud Vendor Assessment Tool (HECVAT) for licensing with Qualtrics (see description of HECVAT below). In addition, UITS reviewed and approved the BCSSE SSSP Request for licensing the Qualtrics CX dashboard. IU Center for Post Secondary Research staff access to BCSSE data is limited to investigators listed in this application. Up to three individual staff at each participating institution are authorized as BCSSE contacts. Any requests by any non-project investigators or authorized institutional staff for institutionally identifiable data transfer must be authorized by the BCSSE PI and the institutionally authorized staff. HECVAT description "In order to protect the institution and its systems, vendors whose products and/or services will access and/or host institutional data must complete the Higher Education Cloud Vendor Assessment Tool. Throughout this tool, anywhere where the term data is used, this is an all-encompassing term including at least data and metadata. Answers will be reviewed by Institution security analysts upon submittal. This process will assist the institution in preventing breaches of protected information and comply with Institution policy, state, and federal law. This is intended for use by vendors participating in a Third Party Security Assessment and should be completed by a vendor."
Explain how subjects’ physical privacy will be protected, both during recruitment/screening and during participation in the research.
Given the minimal risk nature of this study and that the survey questions are not of a nature that would place a subject in physical harm, there is no specific efforts being made to protect the physical privacy associated with the survey.

Is there a potential for subjects to benefit directly from participation in the study?
No

State the potential benefits or information which may accrue to SCIENCE or SOCIETY in general as a result of this work.
At the national level, information produced by BCSSE complements other sources of information about the quality of undergraduate institutions. BCSSE provides prospective students, parents, alumni, institutional officials, state policy makers, governing board members, faculty, and others information about how students expect to spend their time, which is an important predictor of the extent to which they will benefit from attending college. Participating institutions have a basis for evaluating the performance of their students and the extent to which students engage in activities consistent with good practice in undergraduate education. Participating institutions are able to use the information to identify areas in which additional institutional effort is needed to improve the undergraduate experience.

Describe the provisions for monitoring the data to ensure the safety of subjects.
Subject withdrawals and any complaints will be monitored to ensure that study procedures do not result in unanticipated distress to participants.
Select the category below which best applies.
45 CFR 46.404: Research not involving greater than minimal risk to children.

Will you be enrolling foster children or children who are considered wards (i.e. who have been placed in the legal custody of the State or other agency, institution per local, state, or federal law)?
No

Will all or some subjects consent to participate in the research?
All subjects (or their legally authorized representative) will consent to participate in the research.

For those subjects who will consent to participate, explain how subjects (or subjects’ legally authorized representative) will be presented with the information needed to decide to participate, including all elements of informed consent.
They will be presented with the informed consent as the first screen prior to the start of the online survey (web version only). For those completing the paper version of the survey, the informed consent will be provided along with each copy of the survey.

Describe any informed consent tools which will be used to present information to potential subjects (i.e. consent documents, videos, brochure, drug/device information, etc) and how they will be used.
The informed consent is the first screen prior to the start of the online survey (web version only).
Describe the timing of the informed consent process, including how you will ensure potential subjects have sufficient opportunity to discuss and consider participation before agreeing to participate in the research. Depending on the institution, students will be asked to complete the survey during various times including orientation, welcome week, or via email. Students are instructed that their participation is voluntary.

Will you include all required elements of consent in your consent process?
No

Choose the elements which will be omitted from your consent process.
Description of benefits (to subjects or others) that may reasonably be expected from the research
Statement that the study data or biospecimens may/may not be de-identified and used for future research

The IRB must approve a modification to the consent process. Explain how the research involves no more than minimal risk to the subject.
The only risk is a loss of confidentiality.

Explain how the modification will not adversely affect the rights and welfare of subjects.
As currently worded, the Informed Consent Statement described protections against the sole risk of participation (loss of confidentiality). The direct benefits of completing the survey are generally known to be minimal while the indirect benefits of helping an institution improve are also described. Maintaining a shorter statement of informed consent may increase the likelihood that students will read all the relevant information. Research on college students indicates the need for brevity in order to maintain their attention, and a longer statement may deter them from reading important information.
Explain how the research could not be practicably carried out if all elements of informed consent were required.
As noted above, research indicates the need for short, direct messages when communicating with students, and the burden of reading lengthy recruiting messages and the longer Informed Consent Statement may deter participation in the study at a time when survey response rates are declining. The intent of these modifications is to communicate the critical information that students need to know in a way that is easily digestible. Lengthy statements of informed consent may not be read fully by prospective participants.

Explain why the research could not be practicably carried out without identifiers.
The research requires merging responses with institutional records.

Explain how subjects will be informed of pertinent results at the conclusion of the study, if appropriate. If subjects will not be informed, enter N/A.
NA

Indicate in what language(s) the consent conversation will be conducted.
English

Explain how you will ensure potential subjects understand the information you have presented to them before they agree to participate in the study.
The ICS is written in plain, simple English that should be understandable to college students.
Briefly describe any training provided to investigators who are obtaining informed consent.
Those responsible for administering the survey on each campus are provided with instructions on how to administer the survey either via email or in a group format. The instructions stress to the proctors that participation in the survey is voluntary.

Does the research include any minimal risk procedures to which subjects will not consent?
No

For those subjects who will consent to participate, choose whether the consent process will be documented by a written signature from subjects.
No subjects will provide a written signature as documentation of consent.

Since no subjects will provide a written signature as documentation of consent, a waiver of documentation of consent is required. Choose the option which most appropriately applies to your study.
The research presents no more than minimal risk of harm to subjects AND involves no procedures for which written consent is normally required outside of the research context.

Can children provide legal consent for themselves to participate in this study?
No children can provide legal consent for themselves.
Will consent be obtained from subjects’ parents/guardians? Select both options if consent will be obtained from some parents but not all.
No. I am requesting a waiver of parental consent.

Choose the option which most appropriately applies to your study.
Parental/guardian consent cannot be practicably obtained.

Explain how the research involves no more than minimal risk to the subject.
Student responses to questions asked on BCSSE pose no risk to children. The only anticipated risk due to participation would be the possible loss of confidentiality of the responses as part of the data distribution process. The consequences of such loss of confidentiality would be minimal due to the nature of the questions asked on the BCSSE survey, which do not cover topics that would generally be considered of a sensitive nature. Survey questions are not of a nature that would tend to embarrass students or place them at risk of physical, psychological, social, or legal harm.

Explain how the waiver will not adversely affect the rights and welfare of subjects.
Given the protocols and procedures in place, the rights and welfare of subjects will not be adversely affected in any way.

Explain how the research could not be practicably carried out if informed consent were required.
BCSSE is administered to incoming first-year college students during new student orientations, welcome week activities, or at other times prior to or at the onset of their first term/semester on campus. Parents are not usually present with their child during the time of administration of the survey.
Explain why the research could not be practicably carried out without identifiers.

Data collected by BCSSE are used by institutions for at least three reasons. One is for academic advising. Without identifiers, the institutions would not be able to provide the advising reports to divisors for use with their advisees. Two is that many participating institutions do both BCSSE and NSSE. The identifiers allows institutions to link the data at the student-level, which allows institutional assessment of first year and other programs. Third, is that institutions link the data to other institutional records such as persistence data, which allows the institution to then analyze/better understand student persistence on their campus.

Explain how subjects will be informed of pertinent results at the conclusion of the study, if appropriate. If subjects will not be informed, enter N/A.

N/A

In order to describe the process for obtaining assent from children, check all that apply.

Some/all children will provide assent.

Describe the timing of the assent process, including how you will ensure children have sufficient opportunity to discuss and consider participation before agreeing to participate in the research.

The timing for the assent process will be identical as the process for adult subjects providing consent.
Describe any assent tools which will be used to present information to potential subjects (i.e. assent document, study information sheet, videos, brochure, drug/device information, etc) and how they will be used.

The assent tools will be identical as for adult subjects providing consent.

Will children indicate their consent with a signature on an assent or consent statement?

Subjects will not provide a signature

Are you part of a covered entity (health care provider that transmits health information electronically) or are you receiving information from a covered entity as part of your research?

No

Attachments

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<td>Informed Consent Statement</td>
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Data Collection Instrument

Attachment

BCSSE 2023 EXP ITEMS - FINAL.DOCX

Name
BCSSE 2023 Exp items - FINAL

Comments

End of Protocol Form

KC IRB History

approvalDate
June 21, 2004

FOR HSO OFFICE USE ONLY

Action History

Description
Annual Reminder Generated - Expedited or Full Board

Date
October 19, 2020

Action Date
October 19, 2020
Comments
Annual Reminder

Updated By
kc

Update Time
2020-10-19T12:00:00.000Z

Description
Expedited Approval

Date
August 3, 2020

Action Date
August 3, 2020

Comments
Amendment-028: Approved

Updated By
apneel

Update Time
2020-08-03T12:00:00.000Z

Description
Assigned to Agenda

Date
August 3, 2020
Amendment-028: Submitted to IRB

Updated By
apneel

Update Time
2020-08-03T12:00:00.000Z

Description
Amendment Created
Date
August 3, 2020

Action Date
August 3, 2020

Comments
Amendment-028: Created

Updated By
apneel

Update Time
2020-08-03T12:00:00.000Z

Description
Expediting Approval

Date
April 2, 2020

Action Date
April 2, 2020

Comments
Amendment-027: Approved

Updated By
apneel

Update Time
2020-04-02T12:00:00.000Z
Description
Assigned to Agenda

Date
April 2, 2020

Action Date
April 2, 2020

Comments
Amendment-027:

Updated By
apneel

Update Time
2020-04-02T12:00:00.000Z

Description
Submitted to IRB

Date
April 1, 2020

Action Date
April 1, 2020

Comments
Amendment-027: Submitted to IRB

Updated By
colejs

Update Time
2020-04-01T12:00:00.000Z
Amendment Created

Date
April 1, 2020

Action Date
April 1, 2020

Comments
Amendment-027: Created

Updated By
colejs

Update Time
2020-04-01T12:00:00.000Z

Expeditied Approval

Date
February 3, 2020

Action Date
February 3, 2020

Comments
Amendment-026: Approved

Updated By
apneel

Update Time
2020-02-03T12:00:00.000Z
**Description**

Assigned to Agenda

**Date**

February 3, 2020

**Action Date**

February 3, 2020

**Comments**

Amendment-026:

**Updated By**

apneel

**Update Time**

2020-02-03T12:00:00.000Z

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**Description**

Submitted to IRB

**Date**

January 30, 2020

**Action Date**

January 30, 2020

**Comments**

Amendment-026: Submitted to IRB

**Updated By**

colejs

**Update Time**

2020-01-30T12:00:00.000Z
Amendment Created

Date
January 27, 2020

Action Date
January 27, 2020

Comments
Amendment-026: Created

Updated By
colejs

Update Time
2020-01-27T12:00:00.000Z

Annual Reminder Generated - Expedited or Full Board

Date
October 19, 2019

Action Date
October 19, 2019

Comments
Annual Reminder

Updated By
kc

Update Time
2019-10-19T12:00:00.000Z
Submitted to IRB

January 28, 2019

Amendment-025: Submitted to IRB

apneel

2019-01-28T12:00:00.000Z

Returned To PI

January 28, 2019

Amendment-025:

apneel

2019-01-28T12:00:00.000Z
Description
Submitted to IRB

Date
January 24, 2019

Action Date
January 24, 2019

Comments
Amendment-025: Submitted to IRB

Updated By
colejs

Update Time
2019-01-24T12:00:00.000Z

Description
Amendment Created

Date
January 24, 2019

Action Date
January 24, 2019

Comments
Amendment-025: Created

Updated By
colejs

Update Time
2019-01-24T12:00:00.000Z
Expedited Approval

Date
October 19, 2018

Action Date
October 19, 2018

Comments
Renewal-007: Approved

Updated By
apneel

Update Time
2018-10-19T12:00:00.000Z

Assigned to Agenda

Date
October 19, 2018

Action Date
October 19, 2018

Comments
Renewal-007: Research complies with and is subject to 45 CFR 46 effective January 21, 2019 (i.e. Revised Common Rule or 2018 Requirements).

Updated By
apneel
Submit to IRB

October 19, 2018

Returned to PI

October 19, 2018

Renewal-007: Submitted to IRB

Renewal-007:
Submitted to IRB

October 19, 2018

Returned To PI

October 19, 2018
Renewal-007: Submitted to IRB

Date
October 1, 2018

Action Date
October 1, 2018

Comments
Renewal-007: Submitted to IRB

Updated By
colejs

Update Time
2018-10-01T12:00:00.000Z
Action Date
October 1, 2018

Comments
Renewal/Amendment-007: Created

Updated By
colejs

Update Time
2018-10-01T12:00:00.000Z

Description
Renewal Reminder Generated

Date
September 29, 2018

Action Date
September 29, 2018

Comments
Renewal Reminder Letter #2

Updated By
kc

Update Time
2018-09-29T12:00:00.000Z

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| Description          | Expedited Approval          |
| Date                 | May 11, 2018                |
| Action Date          | May 11, 2018                |
| Comments             | Amendment-024: Approved     |
| Updated By           | apneel                      |
| Update Time          | 2018-05-11T12:00:00.000Z    |
Assigned to Agenda

Date
May 11, 2018

Action Date
May 11, 2018

Comments
Amendment-024:

Updated By
apneel

Update Time
2018-05-11T12:00:00.000Z

Submitted to IRB

Date
May 10, 2018

Action Date
May 10, 2018

Comments
Amendment-024: Submitted to IRB

Updated By
colejs

Update Time
2018-05-10T12:00:00.000Z
Returned To PI

May 9, 2018

Amendment-024:

Updated By: apneel

2018-05-09T12:00:00.000Z

Submitted to IRB

May 1, 2018

Amendment-024: Submitted to IRB

Updated By: colejs

2018-05-01T12:00:00.000Z
Returned To PI

May 1, 2018

Amendment-024: Open for incomplete submission

morans

2018-05-01T12:00:00.000Z

Submitted to IRB

May 1, 2018

Amendment-024: Submitted to IRB

colejs

2018-05-01T12:00:00.000Z
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| Description       | Expedited Approval                                                      |
| Date              | April 25, 2018                                                         |
| Action Date       | April 25, 2018                                                         |
| Comments          | Amendment-023: Approved                                                  |
| Updated By        | apneel                                                                 |
| Update Time       | 2018-04-25T12:00:00.000Z                                                |
Assigned to Agenda

Date
April 25, 2018

Action Date
April 25, 2018

Comments
Amendment-023:

Updated By
apneel

Update Time
2018-04-25T12:00:00.000Z

Description
Submitted to IRB

Date
April 24, 2018

Action Date
April 24, 2018

Comments
Amendment-023: Submitted to IRB

Updated By
apneel

Update Time
2018-04-24T12:00:00.000Z
Returned To PI

Date
April 24, 2018

Action Date
April 24, 2018

Comments
Amendment-023: Submitted to IRB

Updated By
colejs

Update Time
2018-04-20T12:00:00.000Z
Amendment Created

Date
April 20, 2018

Action Date
April 20, 2018

Comments
Amendment-023: Created

Updated By
colejs

Update Time
2018-04-20T12:00:00.000Z

Expeditied Approval

Date
February 20, 2018

Action Date
February 20, 2018

Comments
Amendment-022: Approved

Updated By
apneel

Update Time
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Date
February 20, 2018

Action Date
February 20, 2018

Comments
Amendment-022:

Updated By
apneel

Update Time
2018-02-20T12:00:00.000Z

Submitted to IRB

Date
February 20, 2018

Action Date
February 20, 2018

Comments
Amendment-022: Submitted to IRB

Updated By
colejs

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February 5, 2018

Action Date
February 5, 2018

Comments
Amendment-021:

Updated By
apneel

Update Time
2018-02-05T12:00:00.000Z

 Submitted to IRB

Date
February 2, 2018

Action Date
February 2, 2018

Comments
Amendment-021: Submitted to IRB

Updated By
colejs

Update Time
2018-02-02T12:00:00.000Z
Amendment Created

Date
February 2, 2018

Action Date
February 2, 2018

Comments
Amendment-021: Created

Updated By
colejs

Update Time
2018-02-02T12:00:00.000Z

Administrative Correction

Date
October 18, 2017

Action Date
October 18, 2017

Comments
SSS amended in February 2017. Approved version not on the Notes and Attachments tab.

Updated By
apneel
Update Time
2017-10-18T12:00:00.000Z

Description
Amendment Deleted

Date
October 18, 2017

Action Date
October 18, 2017

Comments
Amendment-020: Deleted - AMD not needed

Updated By
apneel

Update Time
2017-10-18T12:00:00.000Z

Description
Amendment Created

Date
October 16, 2017

Action Date
October 16, 2017

Comments
Amendment-020: Created
Updated By
colejs

Update Time
2017-10-16T12:00:00.000Z

Description
Expedited Approval

Date
February 21, 2017

Action Date
February 21, 2017

Comments
Amendment-019: Approved

Updated By
apneel

Update Time
2017-02-21T12:00:00.000Z

Description
Assigned to Agenda

Date
February 21, 2017

Action Date
February 21, 2017
Amendment-019: Submitted to IRB

February 20, 2017

Returned To PI

February 20, 2017
Action Date
February 20, 2017

Comments
Amendment-019: open for pre review

Updated By
apneel

Update Time
2017-02-20T12:00:00.000Z

Description
Submitted to IRB

Date
February 15, 2017

Action Date
February 15, 2017

Comments
Amendment-019: Submitted to IRB

Updated By
colejs

Update Time
2017-02-15T12:00:00.000Z

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Date
December 13, 2016

Action Date
December 13, 2016

Comments
Amendment-018:

Updated By
apneel

Update Time
2016-12-13T12:00:00.000Z

Submitted to IRB

Date
December 7, 2016

Action Date
December 7, 2016

Comments
Amendment-018: Submitted to IRB

Updated By
colejs

Update Time
2016-12-07T12:00:00.000Z
Amendment Created

Date
December 7, 2016

Action Date
December 7, 2016

Comments
Amendment-018: Created

Updated By
colejs

Update Time
2016-12-07T12:00:00.000Z

Expedited Approval

Date
November 15, 2016

Action Date
November 14, 2016

Comments
Renewal-006: Approved

Updated By
morans

Update Time
2016-11-15T12:00:00.000Z
Description
Assigned to Agenda

Date
November 15, 2016

Action Date
November 14, 2016

Comments
Renewal-006:

Updated By
morans

Update Time
2016-11-15T12:00:00.000Z

Description
Submitted to IRB

Date
November 8, 2016

Action Date
November 8, 2016

Comments
Renewal-006: Submitted to IRB

Updated By
colejs

Update Time
2016-11-08T12:00:00.000Z
Renewal with Amendment Created

Date
November 7, 2016

Action Date
November 7, 2016

Comments
Renewal/Amendment-006: Created

Updated By
colejs

Update Time
2016-11-07T12:00:00.000Z

Renewal Reminder Generated

Date
November 7, 2016

Action Date
November 7, 2016

Comments
Renewal Reminder Letter #2

Updated By
kc

Update Time
2016-11-07T12:00:00.000Z
Renewal Reminder Generated

Date
October 23, 2016

Action Date
October 23, 2016

Comments
Renewal Reminder Letter #1

Updated By
kc

Update Time
2016-10-23T12:00:00.000Z

Expedited Approval

Date
February 19, 2016

Action Date
February 19, 2016

Comments
Amendment-017: Approved

Updated By
kmumaw

Update Time
2016-02-19T12:00:00.000Z
Amendment-017: Submitted to IRB

Submitted to IRB

Date
February 9, 2016

Action Date
February 9, 2016

Comments
Amendment-017: Submitted to IRB

Updated By
colejs

Update Time
2016-02-09T12:00:00.000Z

Amendment-017: Withdrawn - returning to researcher for editing

Amendment Withdrawn

Date
February 9, 2016

Action Date
February 9, 2016

Comments
Amendment-017: Withdrawn - returning to researcher for editing

Updated By
morans

Update Time
2016-02-09T12:00:00.000Z
Withdrawn

February 9, 2016

Amendment-017: returning to researcher for editing

morans

2016-02-09T12:00:00.000Z

Submitted to IRB

February 9, 2016

Amendment-017: Submitted to IRB

colejs

2016-02-09T12:00:00.000Z
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Date
March 13, 2015

Action Date
March 12, 2015

Comments
Amendment-016: Approved

Updated By
morans

Update Time
2015-03-13T12:00:00.000Z

Description
Assigned to Agenda

Date
March 13, 2015

Action Date
March 12, 2015

Comments
Amendment-016:

Updated By
morans

Update Time
2015-03-13T12:00:00.000Z
Submitted to IRB

March 4, 2015

Amendment-016: Submitted to IRB

colejs

2015-03-04T12:00:00.000Z

Returned To PI

March 4, 2015

Amendment-016: Returned per colejs request

millsa

2015-03-04T12:00:00.000Z
Submitted to IRB

February 27, 2015

Amendment-016: Submitted to IRB

colejs

2015-02-27T12:00:00.000Z

Amendment Created

February 27, 2015

Amendment-016: Created

colejs

2015-02-27T12:00:00.000Z
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Comments
Renewal-005: Upload nal appendices

Updated By
millsa

Update Time
2014-12-23T12:00:00.000Z

Description
Administrative Correction

Date
December 22, 2014

Action Date
December 22, 2014

Comments
Renewal-005: Uploading approved Prompt Report pdf to replace word version.

Updated By
millsa

Update Time
2014-12-22T12:00:00.000Z
Description
Submitted to IRB

Date
December 2, 2014

Action Date
December 2, 2014

Comments
Renewal-005: Submitted to IRB

Updated By
colejs

Update Time
2014-12-02T12:00:00.000Z

Description
Renewal Created

Date
December 1, 2014

Action Date
December 1, 2014

Comments
Renewal/Amendment-005: Created

Updated By
colejs

Update Time
2014-12-01T12:00:00.000Z
Record Committee Decision

Date
September 18, 2014

Action Date
September 18, 2014

Comments
FYI-001:

Updated By
slbenken

Update Time
2014-09-18T12:00:00.000Z
FYI-001: Need to determine if report represents an unanticipated problem involving risk to subjects or others

Updated By
slbenken

Update Time
2014-09-13T12:00:00.000Z

FYI-001: Submitted to IRB

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slbenken
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Date
April 18, 2014

Action Date
April 18, 2014

Comments
Amendment-014:

Submitted to IRB

Date
April 18, 2014

Action Date
April 18, 2014
Amendment-014: Submitted to IRB

Date
April 11, 2014

Action Date
April 18, 2014

Comments
Amendment-014:

Updated By
kmumaw

Update Time
2014-04-18T12:00:00.000Z

Description
Submitted to IRB

Date
April 18, 2014
Action Date
April 11, 2014

Comments
Amendment-014: Submitted to IRB

Updated By
chwrober

Update Time
2014-04-11T12:00:00.000Z

Description
Amendment Deleted

Date
April 11, 2014

Action Date
April 11, 2014

Comments
Amendment-015: Deleted - Erroneous copy of A014

Updated By
chwrober

Update Time
2014-04-11T12:00:00.000Z

Description
Amendment Withdrawn
Amendment-015: Withdrawn - Merging with A014

Updated By
chwrober

Update Time
2014-04-11T12:00:00.000Z

Description
Returned To PI

Amendment-014: Merging with A015

Updated By
chwrober

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Updated By
kmumaw

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Date
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February 27, 2014

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Amendment-013:

Updated By
kmumaw

Update Time
2014-02-27T12:00:00.000Z

Submitted to IRB

Date
February 25, 2014

Action Date
February 25, 2014

Comments
Amendment-013: Submitted to IRB

Updated By
colejs

Update Time
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Amendment Created

Date
February 21, 2014

Action Date
February 21, 2014

Comments
Amendment-013: Created

Updated By
colejs

Update Time
2014-02-21T12:00:00.000Z

Amendment Deleted

Date
January 9, 2014

Action Date
January 9, 2014

Comments
Amendment-A008: Deleted

Updated By

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Description
Administrative Correction

Date
August 20, 2013

Action Date
August 20, 2013

Comments
Uploading current study documents

Updated By

Update Time
2013-08-20T12:00:00.000Z
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Expedited Approval

Date
February 14, 2013

Action Date
February 13, 2013

Comments
Amendment-012: A012 014

Updated By

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Submitted to IRB

Date
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Action Date
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Comments
Amendment-012: Submit to IRB

Updated By

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Amendment Created

Date
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Action Date
February 12, 2013

Comments
Amendment-012 created

Updated By

Update Time
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Description
Notified Committee

Date
January 11, 2013

Action Date
January 11, 2015

Comments

Updated By

Update Time
2013-01-11T12:00:00.000Z

Description
Expedited Approval

Date
January 11, 2013

Action Date
January 11, 2015

Comments
Renewal-004:

Updated By

Update Time
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Description
Submitted to IRB

Date
January 10, 2013

Action Date
January 10, 2013

Comments
Renewal-004: Submit to IRB

Updated By

Update Time
2013-01-10T12:00:00.000Z

Description
Renewal/Amendment Created

Date
December 21, 2012

Action Date
December 21, 2012

Comments
Renewal/Amendment-004 created

Updated By

Update Time
2012-12-21T12:00:00.000Z
Renewal Reminder Generated

Date
December 14, 2012

Action Date
December 14, 2012

Comments
Renewal Reminder Letter #1

Updated By

Update Time
2012-12-14T12:00:00.000Z

Notified Committee

Date
August 29, 2012

Action Date
August 29, 2012

Comments
A011 012

Updated By

Update Time
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Action Date
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Updated By

Update Time
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Notified Committee

Date
March 1, 2012

Action Date
March 1, 2012

Comments
A010 011

Updated By

Update Time
2012-03-01T12:00:00.000Z
Description
Expedited Approval

Date
March 1, 2012

Action Date
March 1, 2012

Comments
Amendment-010: A010 011

Updated By

Update Time
2012-03-01T12:00:00.000Z

Description
Submitted to IRB

Date
March 1, 2012

Action Date
March 1, 2012

Comments
Amendment-010: Submit to IRB

Updated By

Update Time
2012-03-01T12:00:00.000Z
Description
Expedited Approval

Date
February 13, 2012

Action Date
February 13, 2012

Comments
Renewal-003:

Updated By

Update Time
2012-02-13T12:00:00.000Z

Description
Submitted to IRB

Date
February 9, 2012

Action Date
February 9, 2012

Comments
Renewal-003: Submit to IRB

Updated By

Update Time
2012-02-09T12:00:00.000Z
Renewal/Amendment Created

Date
January 18, 2012

Action Date
January 18, 2012

Comments
Renewal/Amendment-003 created

Update Time
2012-01-18T12:00:00.000Z

Renewal Reminder Generated

Date
January 13, 2012

Action Date
January 13, 2012

Comments
Renewal Reminder Letter #2

Update Time
2012-01-13T12:00:00.000Z
Renewal Reminder Generated

Date
December 19, 2011

Action Date
December 19, 2011

Comments
Renewal Reminder Letter #1

Updated By

Update Time
2011-12-19T12:00:00.000Z

Notified Committee

Date
March 9, 2011

Action Date
March 4, 2011

Comments

Updated By

Update Time
2011-03-09T12:00:00.000Z
Expedited Approval

Date
March 9, 2011

Action Date
March 4, 2011

Comments
Amendment-009:

Updated By

Update Time
2011-03-09T12:00:00.000Z

Submitted to IRB

Date
March 2, 2011

Action Date
March 2, 2011

Comments
Amendment-009: Submit to IRB

Updated By

Update Time
2011-03-02T12:00:00.000Z
Amendment Created

Date
February 24, 2011

Action Date
February 24, 2011

Comments
Amendment-009 created

Updated By

Update Time
2011-02-24T12:00:00.000Z

Notified Committee

Date
February 17, 2011

Action Date
February 17, 2011

Comments

Updated By

Update Time
2011-02-17T12:00:00.000Z
Expedited Approval

Date
February 17, 2011

Action Date
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Comments
Renewal-002:

Updated By

Update Time
2011-02-17T12:00:00.000Z

Submitted to IRB

Date
February 8, 2011

Action Date
February 8, 2011

Comments
Renewal-002: Submit to IRB

Updated By

Update Time
2011-02-08T12:00:00.000Z
Renewal Created

Date
February 8, 2011

Action Date
February 8, 2011

Comments
Renewal-002 created

Updated By

Update Time
2011-02-08T12:00:00.000Z

Administrative Correction

Date
February 8, 2011

Action Date
February 8, 2011

Comments
CO- PIs updated with CR

Updated By

Update Time
2011-02-08T12:00:00.000Z
Amendment Created

Date
January 28, 2011

Action Date
January 28, 2011

Comments
Amendment-008 created

Updated By

Update Time
2011-01-28T12:00:00.000Z

Renewal Reminder Generated

Date
January 25, 2011

Action Date
January 25, 2011

Comments
Renewal Reminder Letter #2

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Expedited Approval

Date
July 1, 2010

Action Date
June 30, 2010

Comments
Amendment-007: CAT 7

Updated By

Update Time
2010-07-01T12:00:00.000Z

Submitted to IRB

Date
June 30, 2010

Action Date
June 30, 2010

Comments
Amendment-007: Submit to IRB

Updated By

Update Time
2010-06-30T12:00:00.000Z
Amendment Created

Date
June 30, 2010

Action Date
June 30, 2010

Comments
Amendment-007 created

Updated By

Update Time
2010-06-30T12:00:00.000Z

Notified Committee

Date
May 13, 2010

Action Date
May 13, 2010

Comments
CAT 7

Updated By

Update Time
2010-05-13T12:00:00.000Z
Expedited Approval

Date
May 13, 2010

Action Date
May 13, 2010

Comments
Amendment-006: CAT 7

Updated By

Update Time
2010-05-13T12:00:00.000Z

Submitted to IRB

Date
May 13, 2010

Action Date
May 13, 2010

Comments
Amendment-006: Submit to IRB

Updated By

Update Time
2010-05-13T12:00:00.000Z
Amendment-006 created

Updated By

Update Time
2010-05-12T12:00:00.000Z

CAT 7

Updated By

Update Time
2010-03-11T12:00:00.000Z
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Renewal Created

Date
March 9, 2010

Action Date
March 9, 2010

Comments
Renewal-001 created

Updated By

Update Time
2010-03-09T12:00:00.000Z

Renewal Reminder Generated

Date
March 8, 2010

Action Date
March 8, 2010

Comments

Updated By

Update Time
2010-03-08T12:00:00.000Z
**Description**

**Expired**

**Date**

March 8, 2010

**Action Date**

March 8, 2010

**Comments**

**Updated By**

**Update Time**

2010-03-08T12:00:00.000Z

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**Description**

**Notified Committee**

**Date**

January 20, 2010

**Action Date**

January 19, 2010

**Comments**

CAT 7

**Updated By**

**Update Time**

2010-01-20T12:00:00.000Z
Description
Expedited Approval

Date
January 20, 2010

Action Date
January 19, 2010

Comments
Amendment-005: CAT 7

Updated By

Update Time
2010-01-20T12:00:00.000Z

Description
Renewal Reminder Generated

Date
January 18, 2010

Action Date
January 18, 2010

Comments
Renewal Reminder Letter #2

Updated By

Update Time
2010-01-18T12:00:00.000Z
Renewal Reminder Generated

Date
January 3, 2010

Action Date
January 3, 2010

Comments
Renewal Reminder Letter #1

Updated By

Update Time
2010-01-03T12:00:00.000Z

Submitted to IRB

Date
December 22, 2009

Action Date
December 22, 2009

Comments
Amendment-005: Submit to IRB

Updated By

Update Time
2009-12-22T12:00:00.000Z
**Description**

**Amendment Created**

**Date**

December 22, 2009

**Action Date**

December 22, 2009

**Comments**

Amendment-005 created

**Updated By**

**Update Time**

2009-12-22T12:00:00.000Z

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**Description**

**Notified Committee**

**Date**

June 19, 2009

**Action Date**

June 18, 2009

**Comments**

Cat 7

**Updated By**

**Update Time**

2009-06-19T12:00:00.000Z
Expedited Approval

Date
June 19, 2009

Action Date
June 18, 2009

Comments
Amendment-004: Cat 7

Updated By

Update Time
2009-06-19T12:00:00.000Z

Submitted to IRB

Date
June 18, 2009

Action Date
June 18, 2009

Comments
Amendment-004: Submit to IRB

Updated By

Update Time
2009-06-18T12:00:00.000Z
Amendment Created

Date
June 10, 2009

Action Date
June 10, 2009

Comments
Amendment-004 created

Updated By

Update Time
2009-06-10T12:00:00.000Z

Administrative Correction

Date
May 19, 2009

Action Date
May 19, 2009

Comments
adding co-investigators

Updated By

Update Time
2009-05-19T12:00:00.000Z


**Description**

**Notified Committee**

**Date**

May 7, 2009

**Action Date**

May 6, 2009

**Comments**

**Updated By**

**Update Time**

2009-05-07T12:00:00.000Z

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**Description**

**Expedited Approval**

**Date**

May 7, 2009

**Action Date**

May 6, 2009

**Comments**

Amendment-003:

**Updated By**

**Update Time**

2009-05-07T12:00:00.000Z
Amendment-003: Submit to IRB

2009-05-07T12:00:00.000Z

Amendment-003 created

2009-04-29T12:00:00.000Z
Description
Notified Committee

Date
April 20, 2009

Action Date
April 15, 2009

Comments

Updated By

Update Time
2009-04-20T12:00:00.000Z

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Description
Expedited Approval

Date
April 20, 2009

Action Date
April 15, 2009

Comments
Amendment-002:

Updated By

Update Time
2009-04-20T12:00:00.000Z
Submitted to IRB

Date
April 20, 2009

Action Date
April 20, 2009

Comments
Amendment-002: Submit to IRB

Updated By

Update Time
2009-04-20T12:00:00.000Z

Amendment Created

Date
April 14, 2009

Action Date
April 14, 2009

Comments
Amendment-002 created

Updated By

Update Time
2009-04-14T12:00:00.000Z
Description
Notified Committee

Date
March 4, 2009

Action Date
March 4, 2009

Comments

Updated By

Update Time
2009-03-04T12:00:00.000Z

Description
Expedited Approval

Date
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Action Date
March 4, 2009

Comments
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Updated By

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Action Date
March 3, 2009

Comments

Updated By

Update Time
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Submitted to IRB

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Action Date
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Comments
Submit to IRB

Updated By

Update Time
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Administrative Details Form

Protocol Details

9031
Protocol Type
Expedited

Billing Account #

Study Status

Submission Details

9000
Submission Review Level
Expedited

9030
Expedited Category.
Category 7: Survey, interview, focus groups, human factor, group behavior or characteristics

9002
Criteria for Approval. Select to confirm.
Approved: The criteria for approval of the research are satisfied in accordance with IU HRPP Policies, and applicable federal regulations.

Protocol Determinations
Protocol Level of Risk.
Minimal risk

Is renewal required for this research?
No

Check all determinations that need to be made.
Informed Consent Waiver
Vulnerable Population

Informed Consent Waivers
Alteration of informed consent granted in accordance with IU HRPP Policies
Waiver of documentation of informed consent granted in accordance with IU HRPP Policies.

Identify population(s) involved in this research.
Children

Involvement of Children: Select to confirm.
The involvement of children in the research is appropriate in accordance with IU HRPP Policies.

Children Category.
The involvement of children in the research satisfies the conditions of Category 404 in accordance with IU HRPP Policies.

Parental Consent/Waivers. Choose all that apply.
Waiver of parental/guardian permission granted in accordance with IU HRPP Policies.
Child Assent. Choose all that apply.
The plan for soliciting and documenting assent from children is appropriate in accordance with IU HRPP Policies.

Other Determinations.