

February 13, 2008

HUMAN RESEARCH PROTECTIONS  
Institutional Review Board

Dr. Christina Falci  
Dr. Julia McQuillan  
717 OLDH  
(0324)

IRB#2008-02-8737 EP

TITLE OF PROJECT: **Faculty Networks and Departmental Climates in STEM at UNL**

Dear Dr. Falci:

This letter is to officially notify you of the approval of your project by the Institutional Review Board (IRB) for the Protection of Human Subjects. It is the Board's opinion that you have provided adequate safeguards for the rights and welfare of the participants in this study. Your proposal seems to be in compliance with this institution's Federal Wide Assurance 00002258 and the DHHS Regulations for the Protection of Human Subjects (45 CFR 46).

Date of EP Review: **02/12/08**

You are authorized to implement this study as of the Date of Final Approval: **02/12/08**

This approval is Valid Until: **02/11/09**

We wish to remind you that the principal investigator is responsible for keeping this Board informed of any changes involved with the procedures or methodology in this study. You should report any unanticipated problems involving risks to the participants or others to the Board. For projects which continue beyond one year from the starting date, the IRB will request continuing review and update of the research project. Your study will be due for continuing review as indicated above. The investigator must also advise the Board when this study is finished or discontinued by completing the enclosed Protocol Final Report form and returning it to the Institutional Review Board.

1. Uploaded on NUgrant is the IRB approved Informed Consent form for this project. Please use this form when making copies to distribute to your participants. If it is necessary to create a new informed consent form, please send us your original so that we may approve and stamp it before it is distributed to participants.

We wish to remind you that the principal investigator is responsible for reporting to this Board any of the following events within 48 hours of the event:

- Any serious event (including on-site and off-site adverse events, injuries, side effects, deaths, or other problems) which in the opinion of the local investigator was unanticipated, involved risk to subjects or others, and was possibly related to the research procedures;
- Any serious accidental or unintentional change to the IRB-approved protocol that involves risk or has the potential to recur;
- Any publication in the literature, safety monitoring report, interim result or other finding that indicates an unexpected change to the risk/benefit ratio of the research;
- Any breach in confidentiality or compromise in data privacy related to the subject or others; or
- Any complaint of a subject that indicates an unanticipated risk or that cannot be resolved by the research staff.

If you have any questions, please contact Shirley Horstman, IRB Administrator, at 472-9417 or email [shorstman1@unl.edu](mailto:shorstman1@unl.edu).

Sincerely,



Dan R. Hoyt, Chair  
for the IRB