Annual Review Form

The Chair, IRB, c/o Registrar; Institute for Clinical Social Work 200 N. Michigan Ave., Suite 407; Chicago, IL 60601

Progress reports must be completed annually to comply with federal regulations. Complete the following form and return it to the address above. Fill out whether study is active or inactive.

| Principal Researcher: |
|--|
| Dissertation Chair: |
| Research Title: |
| Status of Study |
| ACTIVE: still recruiting subjects or follow-up continuing INACTIVE: recruitment completed and analysis/publication in process |
| Since this study was last approved by IRB (check all that apply) |
| There have been no changes made concerning use of human subjects in this project There have been no changes in research protocol |
| New participants are no longer being enrolled |
| Data are still being collected and/or analyzed |
| Modifications to Research Protocol (check all that apply) |
| A change in principal researcher or supervisor |
| A change in treatment procedures/protocolA change in the type or number of subjects used |
| Any other changes |
| Informed Consent |
| Written consent/assent is being obtained from all human participants |
| Participants have been given ample opportunity to ask questions, be debriefed Participants are being paid for their participation |
| Progress Report: Describe results achieved since last approval period. Include the following |
| Number of subjects enrolled Number of subjects who dropped out |
| 3. Progress made: approximate time to complete project |
| |
| Adverse Events: Describe in detail any adverse events. Include the following: |
| 1. Any accidents or unexpected events |
| 2. Any negative responses to research protocol 3. How protocol/debriefing was changed to eliminate risks of above |
| J. How protocondenticing was changed to chimitate tisks of above |
| Signature: Date: |
| Phone: Email: |

Revised 19 Dec 2006