

BARNARD COLLEGE  
Termination of a Research Protocol Involving Human Subjects

Title:	
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<b>Principal Investigator:</b>					
Department:		Telephone:		FAX:	
Associate Investigators: (list only <i>if changed</i> since last report)					

**Study subjects:**

A total of \_\_\_\_\_ subjects were enrolled since the last report to the IRB. Since initial study approval on \_\_\_\_\_ a total of \_\_\_\_\_ subjects have been enrolled in this study and \_\_\_\_\_ have completed or been withdrawn from the study.

**Unforeseen / Adverse developments (since last report to IRB):**

	No unforeseen / adverse developments have occurred.
	Adverse events affecting the risk/benefit ratio of the study have occurred, either from the literature or from the Investigator's findings ( <i>please detail below or on separate sheet</i> ).

**Status of study:**

The study is <b>terminated</b> as of	
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**Permission to continue the study is requested because:**

	Participant enrollment has not yet begun.
	Participant enrollment will continue.
	Participation is complete but data will continue to be analyzed.

**Study methods/procedures have been modified:**

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**If the consent document has been revised (please attach revised consent form).**

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**Brief summary of the findings:**

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**Signature**

As the Principal Investigator of this research project, I certify the following:

That the information provided in this application is complete and accurate.

That I assume full responsibility for the protection of human subjects and the proper conduct of the research.

That the research is performed according to ethical principles and in compliance with all federal, state and local laws, as well as Barnard College policies regarding the protection of human subjects.

Signature of PI		Date:	
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