## Connecticut College

## RESEARCH APPLICATION REVIEW FORM

For Human Subjects Institutional Review Board (IRB)

Researcher Name	Committee Action:
E-mail:	Date:
Box No.:	IRB Comments:
Telephone No.:	
Date Submitted:	
Signature of Faculty Supervisor:	
TYPE OF RESEARCH:	
<ul> <li>☐ Honors Study</li> <li>☐ Masters Thesis</li> <li>☐ Individual Study</li> <li>☐ Course-related project</li> <li>☐ Faculty project</li> <li>☐ Other</li> </ul>	
<b>COMMITTEE MEMBERS</b> (2010-2011):	
Jason Nier (Chair)	-
Michel Belt	
Santiba Campbell (Spring 2011)	
Michelle Dunlap	
Cherise Harris	
Audrey Zakriski	

Please fill out the following form for research plans that involve the use of human participants and **submit** the completed form to the *Human Subjects Institutional Review Board* (Professor Nier, IRB Chair) before the research is begun. The purpose of the form is to bring to the attention of the **IRB** research plans that may involve ethical issues in the use of human participants.

## 1. Title of Research Project:

dure, inclu concernin	uding exactly w g ethical issues	hat the subjects what may be invo	be included in an Appendix. A complete description of the proce- will experience, should be included. Include an explicit statement olved in the research plan. Also submit an outline of your plan for efing statement to subjects.		
3. Other	investigators (	(including name	e, position, and department):		
4. Projec	et period:				
5. Name	of external sp	onsoring agenc	y (if applicable):		
Accounta		PAA)? YES	any data covered by the Health Insurance Portability and NO If yes, what steps are being taken to de-individuate		
7. Projec	et application s	status: New	Renewal		
8. Does	your research	project involve	any of the following:		
	Yes	No			
			participants under the age of 18		
			covert observation		
studies of ethnic and othe			studies of ethnic and other group differences		
	intervention research				
use of deception			use of deception		
i			invasion of privacy		
			aversive (noxious) stimulation		
			induction of mental or physical stress or deprivation (e.g., food, water, sensory, sleep)		
			invasive procedures (e.g., drugs, blood samples, surgery)		
			potentially embarrassing situations		
			consent forms		
			other ethical issues concerning the dignity and welfare of the participants (express below)		

2. Attach an overview of your research proposal, including your hypotheses. All instruments (e.g., surveys,

For every item marked YES or for "consent forms", a NO, provide a brief description of the precise procedure you plan to follow (unless already clearly stated in Item 2).

(NOTE: Items 9-12 po	ertain <i>only</i> to	Psy 101,102	Subject Poo	l Users.)		
9. Estimated number of minutes the experiment will take:				# Minutes		
10. Number of participation # Pa		archer would	like to have a	and why (i	.e., justification):	
11. Number of sessions	the researche	er expects to ru	ın:	#	‡ Sessions	
12. Whether group is:  Average number						
13. If required, training of that that training is a	-		•	has been o	completed, and certificati	on
14. Other comments to	the IRB:					
15. Certifications:						
I certify that the statem College IRB should the protocol. I accept respo pants, and I agree to m	re be any cha onsibility for t	nges in the res	search proto this research	col or if pi h, the supe	roblems arise from this rvision of human partici	-
	(printed)	Investi	gator's Signa	<u>ture</u>	Date	

**PLEASE SUBMIT TO:** Professor Nier (janie@conncoll.edu or Box 5305 — **electronic submissions are preferred**). The Committee Chair will inform you as soon as possible whether it has approved your project or whether additional information or revision is required.