FORM C
HUMAN SUBJECTS FORM

Required for all research involving humans. IRB approval required before experimentation

Deadline Date: February 15, 2009

The Adult Sponsor, Qualified Scientist or Designated Supervisor cannot serve on the SCR/IRB for their particular project.

Student's Name: _____
First Last

Title of Project: _____

To be completed by Student Researcher: (All questions are applicable and must be answered)

1. Explain why human subjects are proposed or necessary to this research.

2. Describe and assess any potential risk (physical, psychological, social, legal or other).

3. Describe how informed consent will be obtained.

4. Describe procedures to minimize risk.

5. Describe benefits to the individual or society.

6. Compare the benefits of this research to the risks.

To be completed by Institutional Review Board (IRB) prior to experimentation:
(Risk includes but is not limited to exercise, ingestion, physical and emotional stress, invasion of privacy)

Please check appropriate decision:

- ☐ Minimal risks involved
- ☐ More than minimal risks involved…..Qualified Scientists and Informed Consent are required for all subjects
- ☐ Unacceptable risks involved……Project must be revised

(see reverse side)

YORK COLLEGE
The City University of New York

For JSHS National Guidelines, visit www.jshs.org
Please reproduce **3 copies** of this form for each of your student applicants

Please send this form and the Student Registration Forms to:
Dr. Panayiotis Meleties, Program Director,
York College – CUNY
94-20 Guy R. Brewer Blvd., Jamaica, NY 11451

Minimum of three (3) signatures required □ Specify position:

IRB Member's Printed Name	Signature	Psychologist/M.D./Nurse (circle one)	Date of Approval
IRB Member's Printed Name	Signature	Science Teacher	Date of Approval
IRB Member's Printed Name	Signature	School Administrator	Date of Approval

□ **ATTENTION:** 1. When project concerns behavioral research, the IRB must include a psychologist, psychiatrist, or individual with human behavioral training.
2. Tests or questionnaires of any type must be attached to the Research plan and be reviewed by the IRB.

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