Human Subjects Form (4A)

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

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

Student's Name _____

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 for this research:

2) Describe and assess any potential risk (physical, psychological, social, legal or other): (see risk description, p. 16-17 of the Rules)

3) Describe how informed consent will be obtained (attach a sample of completed **Form 4B**):

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, and invasion of privacy.)

☐ Minimal risks involved . . . . . **Informed Consent (Form 4B)** is strongly recommended for all subjects, but is not required. (See clarification on p. 16-17 of the International Rules.)
(See risk section on p. 16-17)

☐ More than minimal risks involved . . . . . **Qualified Scientist** and **Informed Consent (Forms 2, 4B)** are required for all subjects. **Designated Supervisor (Form 3)** may be required.

☐ Unacceptable risks involved . . . . . Project is rejected and must be revised.

(must check appropriate decision)

MINIMUM OF THREE SIGNATURES REQUIRED*

1) Medical Professional: (a psychologist, psychiatrist, medical doctor, physician's assistant, or registered nurse) (circle one)

Chester C. Wood, M.D.		
Member of IRB's Printed Name	Signature	Date of Approval

2) Teacher:

Arthur Amend, M.S.		
Member of IRB's Printed Name	Signature	Date of Approval

3) School Administrator:

Joseph Ellis, Ed.D.		
Member of IRB's Printed Name	Signature	Date of Approval

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