

**TOUGALOO COLLEGE INSTITUTIONAL REVIEW BOARDS REVIEW  
DOCUMENTAITON OF REVIEW AND APPROVAL**

IRB STUDY NUMBER: \_\_\_\_\_  
*(IRB Office will assign)*

**SECTION 1: INVESTIGATOR INFORMATION**

**Principal Investigator:** \_\_\_\_\_ **Department:** \_\_\_\_\_  
(Last, First, Middle Initial – must have faculty/staff status or faculty sponsor must sign)

Building/Room No.: \_\_\_\_\_ Phone: \_\_\_\_\_ E-mail: \_\_\_\_\_

**Contact Information:**

Name: \_\_\_\_\_ Address: \_\_\_\_\_ Phone: \_\_\_\_\_

Fax: \_\_\_\_\_ E-Mail: \_\_\_\_\_

If this is a Student Protocol, List Name of Student: \_\_\_\_\_ Phone: \_\_\_\_\_

Protocol Title: \_\_\_\_\_

Sponsor/Funding Agency: \_\_\_\_\_ PI on Grant: \_\_\_\_\_

Sponsor Protocol #/Grant#: \_\_\_\_\_ Period: From \_\_\_\_\_ to \_\_\_\_\_

Sponsor Type:  Federal  State  Industry  Not-for-Profit  Unfunded  Internally Funded

Grant Title (If different from project title): \_\_\_\_\_

**SECTION II: TYPE OF REVIEW**

Expedited Review

Full Board Review

**SECTION III: SPECIAL SUBJECT POPULATIONS**

Research to Include:  Minors  Pregnant Women  Cognitively Impaired  Prisoners

Economically or Educationally Disadvantaged  Fetuses (Or Fetal Tissue)

**SECTION IV: RESEARCH SUBMISSION**

Included with Research Submission  Informed Consent, dated\*:

Authorization, dated\*\*:

Summary Safeguard Statement, dated\*\*:

Protocol, dated\*\*:

Drug Brochure, dated\*\*:

Advertisement, dated\*\*:

Other: Description: \_\_\_\_\_, dated\*\*:

*\*version dates are required on the informed consent statements*

*\*\*dates are optional and only necessary if required by the investigator or sponsor*

**SECTION V: INVESTIGATOR STATEMENT OF COMPLIANCE**

I assure the Board that all procedures performed under the project will be conducted in strict accordance with those federal regulations and Tougaloo College policies that govern research involving human subjects. I agree to submit any deviation from the project (e.g. change in principal investigator, research methodology, subject recruitment procedures, etc.) to the Board in the form of an amendment for IRB approval prior to implementation. By signing this form, I am certifying that all co-investigators listed on the study are aware of the research and are agreeing to participate.

Note: This form and any additional material requested by the Board will not be processed unless they are neatly typed and legible, properly prepared, and signed personally by the principal investigator.

Signature of Investigator: \_\_\_\_\_

Date: \_\_\_\_\_

**SECTION VI: IRB COMMITTEE APPROVALS**

\_\_\_\_\_  
Committee Member                      \_\_\_\_\_  
Date                      Approved: \_\_\_\_\_                      Disapproved: \_\_\_\_\_  
Reason for Disapproval: \_\_\_\_\_

\_\_\_\_\_  
Committee Member                      \_\_\_\_\_  
Date                      Approved: \_\_\_\_\_                      Disapproved: \_\_\_\_\_  
Reason for Disapproval: \_\_\_\_\_

\_\_\_\_\_  
Committee Member                      \_\_\_\_\_  
Date                      Approved: \_\_\_\_\_                      Disapproved: \_\_\_\_\_  
Reason for Disapproval: \_\_\_\_\_

\_\_\_\_\_  
Committee Member                      \_\_\_\_\_  
Date                      Approved: \_\_\_\_\_                      Disapproved: \_\_\_\_\_  
Reason for Disapproval: \_\_\_\_\_

\_\_\_\_\_  
Committee Member                      \_\_\_\_\_  
Date                      Approved: \_\_\_\_\_                      Disapproved: \_\_\_\_\_  
Reason for Disapproval: \_\_\_\_\_

**SECTION VII: IRB APPROVAL**

This protocol, informed consent statement, authorization, and/or waiver of authorization for use of human subjects in research has been reviewed and approved by the Tougaloo College Institutional Review Board for a maximum of a one-year period beyond the final approval date unless otherwise indicated as follows:

\_\_\_\_\_  
\_\_\_\_\_

Authorized IRB Signature: \_\_\_\_\_                      IRB Approval Date: \_\_\_\_\_