

**COUNCIL FOR SCITIFIC AND INDUSTRIAL RESEARCH-INSTITUTIONAL REVIEW BOARD (IRB)
APPLICATION FORM FOR PROTOCOL SUBMISSION**



(Please complete and submit together with full Protocol for IRB Consideration)

PART 1: ADMINISTRATIVE INFORMATION OF PRINCIPAL INVESTIGATOR (S)

1.1 Title of study:

1.2 Full name of Principal Investigator (s) (PI) (Surname first):
(please specify if more than one)

i. Surname:		First Name:
ii. Institutional Affiliation:		Others:
iii. Full Postal Address of Principal Investigator:		
iv. Telephone:	v. Mobile:	v. Email:

PART 11: INFORMATION ON COLLABORATOR (S)/ CO-PRINCIPAL INVESTIGATORS
(please specify if more than one)

1.6(a) Name of first Collaborator:	Institutional Affiliation (s):	
	Full Address:	
	Telephone:	Mobile:
	Email:	
1.6(b) Name of second Collaborator:	Institutional Affiliation:	
	Full Address:	
	Telephone:	
	Email:	

1.6(c) Name of third Collaborator	Institutional Affiliation:	
	Full Address:	
	Telephone:	Mobile:
	Email:	

PART III: INFORMATION ON SPONSOR (S)

1.7 Name of Sponsoring Agency (please specify name of lead person)	Institutional Affiliation (s):	
	Full Address	
	Telephone:	Mobile:
	Email:	

PART IV: INFORMATION ON PROPOSED STUDY

1. NATURE OF PROTOCOL (Please tick appropriate column)

i) Institutional Protocol <input type="checkbox"/>	ii) Academic Protocols <input type="checkbox"/>	iii) Individual <input type="checkbox"/>	iv. Others(specify) <input type="checkbox"/>
	PhD <input type="checkbox"/> MSc <input type="checkbox"/> MPhil <input type="checkbox"/> MPH <input type="checkbox"/> Undergraduate <input type="checkbox"/>		

2.TYPE OF STUDY (Please tick appropriate column)

Type A	B. Type B	Type C
i. Clinical Trial <input type="checkbox"/>	iv. Social Science <input type="checkbox"/> (please tick which of the below is applicable to your study)	v. Implementation Research <input type="checkbox"/>
ii) Biomedical Study/Epidemiological Study <input type="checkbox"/>	a. Economic studies <input type="checkbox"/>	
iii) Others (specify): <input type="checkbox"/>	b. Policy Studies <input type="checkbox"/>	
	c. Exploratory studies <input type="checkbox"/>	
	d. Monitoring and Evaluation studies <input type="checkbox"/>	
	h. Other (Specify) <input type="checkbox"/>	

3. STUDY SITE (S)			
i. Name of Study Site:	Region (s): Greater Accra		
	District (s): Accra		
	Sub-District (s)		
ii. Name of Study Site:	Region (s):		
	District (s):		
	Sub-District (s)		
iii. Name of Study Site:	Region (s):		
	District (s):		
	Sub-District (s)		
iv. Name of Study Site:	Region (s):		
	District (s):		
	Sub-District (s)		
4. DATE OF INITIAL SUBMISSION TO CSIR-IRB: DD/MM/YY			
5. DURATION OF STUDY:	Start of Study:	End of Study:	
PART V: TYPE OF REQUEST (please tick appropriately)			
A) New Submission:	1. Yes <input type="checkbox"/>	2. No <input type="checkbox"/>	
B) Request for Amendment:	1. Yes <input type="checkbox"/>	2. No <input type="checkbox"/>	
C. Type of Amendment: <input type="checkbox"/>			
i. Additional information to protocol <input type="checkbox"/>	ii. Change of study site <input type="checkbox"/>	iii. Additional study site <input type="checkbox"/>	iv. Additional PI (s) <input type="checkbox"/>
v. Change of PI(s) <input type="checkbox"/>	vi. Work on Informed Consent <input type="checkbox"/>	vii. Work on Questionnaire (s) <input type="checkbox"/>	
viii. Others (please specify):			

PART VI: FOR IRB OFFICIAL USE ONLY

1. Protocol ID No: Date Received: Month received: Year received:

2. Process of Protocol Consideration:

i. Underwent Full General Review

ii Recommended for Expedited Review

3. Date of Initial Review:

4. Outcome of Initial Review:

i. Outright approval

ii. Conditional approval

iii. Redesign document

iv. Pending

Other (specify:

5. Date of Re-submission:

6. Outcome of Resubmission Review:

i. Approved

ii. Conditional

iii. Pending

Others (specify):

7. Date of Final Approval:

8. Status of Approved Study:

i. Started

ii. Ongoing

iii. Completed

iv. Yet to Start

vi. Suspended

vi. Discontinued

vii. Others (specify)

9. Request for Presentation of Study by PI (s) (1) Yes (2) No

10. Submission of Reports:

A. Submitted

B. Not Submitted

Type of Report Submitted:

i. Initial/ Inception report

ii. Interim Report

iii. Progress report

iv. Adverse Report/Serious Adverse report

v. Final /Completion report

Adapted from- GHS