

**CHONG HUA HOSPITAL**

Healing with Passion. Caring with Compassion.

**IRB APPLICATION FORM
AND QUESTIONNAIRE**

Form HRP-IRB-010

NAME OF ORGANIZATION/INSTITUTION: _____ REFERENCE NO: _____

TITLE OF RESEARCH:	
Anticipated start date	
Anticipated end date	
Principal Investigator: (Must be a member of the consultant staff of Chong Hua Hospital)	
Specific Role in this project:	
Sub – Investigator: (Must be a member of the consultant staff of Chong Hua Hospital)	
Specific Role in this project:	
Has this protocol been disapproved by another IRB/IEC or hospital? YES ____ NO ____ If YES from which IRB/IEC or HOSPITAL?	

INVESTIGATOR'S ASSURANCE
THE SIGNATURES BELOW SIGNIFY THAT:
<ul style="list-style-type: none">• The information and documents provided is/are accurate, current and valid• The principal investigator has the ultimate responsibility for the protection of rights, welfare and safety of the subjects• The principal investigator has the ultimate responsibility for the ethical conduct of the research• Each individual listed as investigator has received the required training on Good Clinical Practice• Each investigator and member of the team has the necessary experience on how to conduct a research on human subjects and shall abide by the regulations of Chong Hua Hospital in its conduct• The principal investigator has the ultimate responsibility for the prompt management of any adverse reactions or suspected adverse reactions attendant to the conduct of the study.• No research or part of it will commence before the IRB has given its approval• The research will be conducted according to the protocol or its amendments duly approved by the Chong Hua Hospital IRB.

Principal Investigator:	[Printed Name and Signature]	Date
Sub – Investigator:	[Printed Name and Signature]	Date
Received by:	[Printed Name and Signature]	Date Received
		Time Received

**SECTION I: INVOLVEMENT OF HUMAN SUBJECTS**

1. Does the study involve human subjects? ☐ Yes ☐ No

If the answer is No, you may not submit the study for IRB review

SECTION II: PROJECT FUNDING

1. Is the project funded? ☐ Yes ☐ No

If the project is funded, kindly specify the funding source.

SECTION III: CONFLICT OF INTEREST (only required for funded research)

1. Is there any real, potential or apparent conflict of interest on the part of Investigator or any of the study team? ☐ Yes ☐ No

If Yes, please declare and explain.

N.B. NON DISCLOSURE OF ANY CONFLICT OF INTEREST MAY AFFECT IRB APPROVAL.

SECTION IV: TYPE OF RESEARCH STUDY

STUDY TYPE: (Mark "✓" whichever apply to the study)

- | | | | | |
|------------------------------------------|----------------------------------------|---------------------------------------|---------------------------------------------|-------------------------------------------|
| <input type="checkbox"/> Survey | <input type="checkbox"/> Social | <input type="checkbox"/> Medical | <input type="checkbox"/> Community Based | <input type="checkbox"/> Individual Based |
| <input type="checkbox"/> Screening | <input type="checkbox"/> Observational | <input type="checkbox"/> Epidemiology | <input type="checkbox"/> Intervention Study | |
| <input type="checkbox"/> Clinical Trial: | <input type="checkbox"/> Phase I | <input type="checkbox"/> Phase II | <input type="checkbox"/> Phase III | <input type="checkbox"/> Phase IV |
| <input type="checkbox"/> Genetic Study | <input type="checkbox"/> Retrospective | <input type="checkbox"/> Prospective | <input type="checkbox"/> Others _____ | |
| <input type="checkbox"/> Single Center | <input type="checkbox"/> Multicenter | <input type="checkbox"/> Others _____ | | |

**SECTION V: Study Population**

1. Does the study involve healthy volunteers? ☐ **Yes** ☐ **No**
2. Does the study involve patients with disease? ☐ **Yes** ☐ **No**
3. Does the study involve a vulnerable population? ☐ **Yes** ☐ **No**

If Yes, please identify.

4. Will the study exclude a particular group of individuals ☐ **Yes** ☐ **No**

If Yes, please identify.

SECTION VI: Characteristics of Study Population (Mark "✓" whichever apply to the study)

- Age Range \Rightarrow ☐ 0 -17 yrs ☐ 18 - 44 yrs ☐ 45 - 65 yrs ☐ \geq 66 yrs
- Pediatric \Rightarrow ☐ None ☐ < 1 yr ☐ 1-3 yrs ☐ 4 -14 yrs
- Impaired \Rightarrow ☐ None ☐ Physically ☐ Cognitively ☐ Mentally

SECTION VII: Drugs/Devices, Genetic Testing, Radiation and Biological Samples

Does the study involve the use of any of the following?

1. An FDA approved drug or medical device ☐ **Yes** ☐ **No**
2. Unapproved indication for an FDA approved drug ☐ **Yes** ☐ **No**
3. An investigational medical device ☐ **Yes** ☐ **No**
4. A non-medical device ☐ **Yes** ☐ **No**
5. A proprietary product ☐ **Yes** ☐ **No**
6. A biological agent ☐ **Yes** ☐ **No**
7. A genetic testing ☐ **Yes** ☐ **No**
8. Radiation exposure ☐ **Yes** ☐ **No**