**EC-2.1 Form**

(For Faculty of Public Health Staff)

# Project Proposal

*“This project will be conducted after receiving the approval from ethical review committee for human research.”*

1. **Title of Project:** ………………………………………………………………………...….

…………………………………………………………………………………………...….

1. **Investigators:**

**2.1 Principal Investigator :** ……………………..……………………………..………....

Contact address: ……………………..…………………………………………………….

……………………………………………………………………………………………..

Tel: ….………………….... Fax:….……………..….. E-mail: …...……….……….……..

**2.2** **Co-Investigator 1:** ……………………..……………………………..………………

Contact address: ……………………..…………………………………………………….

……………………………………………………………………………………………..

Tel: ….………………….... Fax:….……………..….. E-mail: …...……….……….……..

**2.3** **Co-Investigator 2:** ……………………..……………………………..………………

Contact address: ……………………..…………………………………………………….

……………………………………………………………………………………………..

Tel: ….………………….... Fax:….……………..….. E-mail: …...……….……….……..

**2.4** **Co-Investigator 3:** ……………………..……………………………..………………

Contact address: ……………………..…………………………………………………….

……………………………………………………………………………………………..

Tel: ….………………….... Fax:….……………..….. E-mail: …...……….……….……..

1. **Funding source:**………………………...……**Budget:** ………………….……………….

**4. Research category:**

[ ] Biomedical Research

[ ] Social science Research

**5. Research area:**

[ ] Infectious Disease

[ ] Non-infectious Disease

[ ] Public Health Technology

[ ] Public Health Administration and Health System Management

[ ] Environmental Health and Occupational Health

[ ] Health Promotion

[ ] Community Health

[ ] Public Health Nursing

[ ] Food and Nutrition

[ ]Others *(please indicate)* ……………………………………………….……………

**6. Rationale and background (clear, comprehensive and concise):**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**7. Objective(s):**

**7.1 General objective**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

…………………………..……………………………………………………………..……

**7.2 Specific objective**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**8. Reasons for conducting this study in human subjects:**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**9. Benefits/usefulness to the subjects and society in general:**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10. Methodology:**

**10.1 Research design and Data collection method**

**10.1.1 Research design**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.1.2 Data collection method**

* Self-administered questionnaires
* Interviews
* Physical examination
* Others *(please indicate)* ……………………………………………………...…

**10.2 Reference population, Samples, Sample size and Sampling technique**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.3 Inclusion criteria**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.4 Exclusion criteria**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.5 Discontinuation criteria**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.6 Duration of data collection**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**10.7 Data Collection (describe in detail, particularly the activities dealing with human subjects)**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**11. Risks and/or undesirable consequences that may happen to subjects and actions to be taken to prevent or reduce them:**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**12. Ethical consideration:**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**12.1 Incentive present**

[ ] No

[ ] Yes *(please indicate):* ……………………………….…….……….….

**13. Are there any negative impacts that may happen to institutions, society, culture**

**or environment?**

[ ] No

[ ] Yes *(please indicate the impacts):* ……………………………….…….……….….

…..……………………………………………...……...…………………………………………….………………………………………………………………………………...………………….…………………………………………………………………………...……

*Measures to prevent or confront such impacts:*………….………….....……...

……..………………………………………………...………………………………...…………….…………………………………………………………………………………...……………….……………………………………………………………………………...…

**14. Consent approval requirements:**

[ ] Subject or representative

[ ] Head or authorized person of the designated institute

**15. Research presentation:**

[ ] Without declaring names, but with disclosing occupation of subjects

and/or site of study

[ ] Without declaring names, and site of study, but with disclosing occupation of

subjects

[ ] Without declaring names, site of study, and occupation of subjects

**16. References (Vancouver style):**

………………………………………………………………………………………..…

…………………………..……………………………………………………………..……

……………………………………………..…………………………………………..……

**This project will be performed in accordance with the Declaration of the Helsinki Ethical Principles**.

Signature …………..…………….…….…….… (Principal Investigator)

(………….……………….…………..)

Date………………………