

STIU Institutional Review Board's Human Research Ethics Checklist Form (STIU_IRB)	Date:
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Project Name:

Principle Investigator:

Please check below

Lecturer from Department/Faculty/Program _____

Doctorate Student Master's degree Student Bachelor's degree Student

Other (please specify) _____

Contact number: _____ Email: _____

Please complete this checklist to determine the risk of your research project and attach this checklist with the Human Research Ethics Application Form if any of the following topics is checked.

A "YES" answer may indicate that your research is not negligible risk, and then the Human Research Ethics Application Form will need to be completed.

Please in the box, if any of the following research topics covered in some part of your project.

1. Parenting issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
2. Sensitive - Personal issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
3. Sensitive - Cultural issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
4. Grievances, Death or Serious or Traumatic Loss	<input type="checkbox"/> Yes	<input type="checkbox"/> No
5. Depression, Anxiety, Suicide	<input type="checkbox"/> Yes	<input type="checkbox"/> No
6. Gambling	<input type="checkbox"/> Yes	<input type="checkbox"/> No
7. Drugs	<input type="checkbox"/> Yes	<input type="checkbox"/> No
8. Sexuality / Abuse	<input type="checkbox"/> Yes	<input type="checkbox"/> No
9. Race / Ethnic / Gender	<input type="checkbox"/> Yes	<input type="checkbox"/> No

10. Disease / Health problem	<input type="checkbox"/> Yes	<input type="checkbox"/> No
11. Criminal issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
12. Sensitive Cultural / Social issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
13. Religious issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
14. Political issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No
15. Mentally ill patients	<input type="checkbox"/> Yes	<input type="checkbox"/> No
16. Children under 18 years old of age	<input type="checkbox"/> Yes	<input type="checkbox"/> No
17. Elderly people with memory handicapped	<input type="checkbox"/> Yes	<input type="checkbox"/> No
18. Dementia patients	<input type="checkbox"/> Yes	<input type="checkbox"/> No
19. Nightlife waiters and massage parlors	<input type="checkbox"/> Yes	<input type="checkbox"/> No
20. Intellectual disabilities/ADHD	<input type="checkbox"/> Yes	<input type="checkbox"/> No
21. Serious infectious disease patients	<input type="checkbox"/> Yes	<input type="checkbox"/> No
22. Ethnic groups minority of different nationalities and/or religions	<input type="checkbox"/> Yes	<input type="checkbox"/> No
23. Inmates/accuses/ defendants in criminal cases	<input type="checkbox"/> Yes	<input type="checkbox"/> No
24. Gamblers/waiters in gambling establishments	<input type="checkbox"/> Yes	<input type="checkbox"/> No
25. Disabled persons	<input type="checkbox"/> Yes	<input type="checkbox"/> No
26. Sex workers	<input type="checkbox"/> Yes	<input type="checkbox"/> No
27. LGBT people	<input type="checkbox"/> Yes	<input type="checkbox"/> No
28. Pregnant women	<input type="checkbox"/> Yes	<input type="checkbox"/> No
29. Immigrants/people exile/migrant workers	<input type="checkbox"/> Yes	<input type="checkbox"/> No
30. Soldiers	<input type="checkbox"/> Yes	<input type="checkbox"/> No
31. Drug addicts	<input type="checkbox"/> Yes	<input type="checkbox"/> No

Please in the box, if any of the following procedures used in your project.

1. Use personal data obtained from the government offices/agency	<input type="checkbox"/> Yes	<input type="checkbox"/> No
2. Use personal data obtained from the non-government offices/agency	<input type="checkbox"/> Yes	<input type="checkbox"/> No
3. Deception of participants	<input type="checkbox"/> Yes	<input type="checkbox"/> No
4. Concealing the purpose of the research	<input type="checkbox"/> Yes	<input type="checkbox"/> No
5. Covert Observation	<input type="checkbox"/> Yes	<input type="checkbox"/> No
6. Audio, visual recording without consent	<input type="checkbox"/> Yes	<input type="checkbox"/> No
7. Recruitment of a third party or agency	<input type="checkbox"/> Yes	<input type="checkbox"/> No
8. Psychological treatment	<input type="checkbox"/> Yes	<input type="checkbox"/> No
9. Special method of teaching	<input type="checkbox"/> Yes	<input type="checkbox"/> No
10. Administration of drugs	<input type="checkbox"/> Yes	<input type="checkbox"/> No
11. Use of medical records where participants can be identified or liked	<input type="checkbox"/> Yes	<input type="checkbox"/> No
12. Research conducted in unsafe environment	<input type="checkbox"/> Yes	<input type="checkbox"/> No
13. Interview children without parental or guardian consent	<input type="checkbox"/> Yes	<input type="checkbox"/> No
14. Research involves sensitive cultural, social, political or religious issues	<input type="checkbox"/> Yes	<input type="checkbox"/> No

Research proposal should be submitted with details, including:

	Require Document/Contents	Yes	No
1	Project Name:		
2	Background of problem		
3	Objectives of research		
4	Hypothesis and/or conceptual framework		
5	Review literature		
6	Research methodology		
7	Detail of population/research participants e.g. number, qualifications, inclusion/exclusion criteria, sample size determination		
8	Details of how to approach potential participant/access to medical record, recruitment method, informed consent process etc.		
9	Benefits and risks to research participant or to public including risk management		
10	How to protect welfare and safety of participants, how to keep privacy & confidentiality		
11	References		

I (researcher) am willing to comply with the conditions of the Board of Directors as follows:

1. The researcher is aware that it is a violation of human research ethics. If research data was collected prior to approval or approval from the Human Research Ethics Committee.
2. When the research project certificate expires but the data collection in-person/person/ person has not yet been completed. The researcher had to stop the data collection process and request a new authorization and must be done at least 30 days before the certificate expires along with submitting research progress reports. If the research process is left with data processing analyze data or write a research report, the researcher does not need to apply a new ethical certification.
3. The researcher will carry out the research process as indicated in the research proposal/thesis outline strictly.
4. The researcher will use the data sheet for the sample/participant in the research, consent document of the sample/person participating in the research, and research methods only as the committee's approved.

5. If a serious adverse event occurs in a committee-approved data collection place, the researcher must report to the committee within 5 working days.
6. If there are adjustments/changes in research operations, the researcher will send a list of requests for adjustments/changes to the committee for consideration and/or approval/certification before proceeding.
7. Research projects undertaken not more than 1 year, the researcher must submit the research project completion report and abstract of research results within 30 days upon completion of the research project. For a research project that is a student thesis/dissertation must submit an abstract of research results within 30 days upon completion of the research project and the thesis/dissertation has been submitted.
8. I wish to obtain a certificate.
 - Thai language (specify the name-surname Thai language.....)
 - English (specify the name-surname English.....)

..... (.....) Advisor/...../..... (.....) Main Researcher/...../.....
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Endorsement of Human Research Ethics Review Request

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 (.....)
Dean/Associate Dean
/...../.....