



CERTIFICATE OF APPROVAL

COA: 182/2025

Protocol No: EC 25159-18

Title The Application of Knowledge Management and Oral Tradition in
Synthesizing the Maha Thaksa Canon

Principal investigator: Assoc. Prof. Dr.Kowit Nambunmee

School: Health Science

Funding support: XIAN CHIANG CO., LTD.

Approval:

- | | |
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| 1) Research protocol | Version 2 Date August 22, 2025 |
| 2) Information sheet and informed consent documents | Version 2 Date August 22, 2025 |
| 3) Interview guide | Version 1 Date July 1, 2025 |
| 4) Advertising Information Sheet | Version 2 Date August 22, 2025 |
| 5) Principal investigator and Co-investigators | |
| - Assoc. Prof. Dr.Kowit Nambunmee | - Miss Tharinya Kaviya |

The aforementioned documents have been reviewed and approved by the Mae Fah Luang University Ethics Committee on Human Research in compliance with international guidelines such as Declaration of Helsinki, the Belmont Report, CIOMS Guidelines and the International Conference on Harmonization of Technical Requirements for Registration of Pharmaceuticals for Human Use - Good Clinical Practice (ICH - GCP)

Date of Approval: September 22, 2025


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(Jullapong Achalapong, M.D.)

Chairperson of the Mae Fah Luang Ethics Committee on Human Research



For research protocol approved by the Mae Fah Luang University Ethics Committee on Human Research (MFU EC), the investigators must comply with the followings:

1. Strictly conduct the research as required by the protocol.
2. Use only the information sheet, consent form, questionnaire, case record form and advertisement bearing the MFU EC stamp of approval.
3. Submit a progress report (AP 05/2024) for continuing review and for renewing the approval within 30 days before expiration date.
4. When there are changes of the protocol, the investigator must submit an amendment report (AP 06/2024) with amended protocol for MFU EC approval before implementing any changes in the research (unless those changes are required urgently for the safety of the research subjects).
5. When there is any unanticipated problem or serious adverse event, the investigator must submit a safety report (AP 07/2024) as set forth in the ICH-GCP.
6. When there is any deviation or non-compliance with the approved protocol, the investigator must submit a protocol deviation/non-compliance report (AP 08/2024).
7. When the research is complete or terminated, the investigator must submit a closing report (AP 09/2024).

Please go to <https://ethic.mfu.ac.th> to download MFU EC forms for reporting.

I, as an investigator, agree to comply with the above obligation.

(Assoc. Prof. Dr.Kowit Nambunmee)

Date