

	คณะกรรมการกลางพิจารณาจริยธรรมการวิจัยในคน Central Research Ethics Committee; CREC โทรศัพท์ 082-258-9529 E-mail: official@crecthailand.org	AO 01-S10
		V. 5.1 date 15 Oct 2024
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แบบประเมินรายงานความก้าวหน้าของการวิจัย

Assessment of Progress Report

PROTOCOL No.:	CREC No:
PROTOCOL TITLE:	
Principal Investigator:	
Institution:	
Sponsor:	
Last Date of Approval: / /	Date of Expiration: / /
Protocol Renewal: ☐ yes (ขอต่ออายุการรับรอง) ☐ No (รายงานความก้าวหน้า ไม่ขอต่ออายุการรับรอง)	
Submission: ☐ More than 30 days before the expiration/ due date ☐ Within 30 days before the expiration/ due date ☐ After expired date	
Type of continuing review	

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☐ Expedited review - Qualified for expedited review at the time of initial review. - Previously approved in full board but at this time of continuing review: (a) Where (i) <i>the research is permanently closed to the enrollment for new subjects.</i> (ii) <i>all subjects have completed all research-related interventions; and</i> (iii) <i>the research remains active only for long-term follow-up of subjects; or</i> (b) Where no subjects have been enrolled and no additional risks have been identified; or (c) Where the remaining research activities are limited to data analysis - Previously approved in full board but meets the following conditions ▪ <i>Research not conducted under and investigational new drug (IND) or investigational device exemption (IDE);</i> ▪ <i>Research involves no greater than minimal risk to the subjects; and</i> ▪ <i>No additional risks have been identified.</i>		☐ Full board review Not eligible for expedited review		
Issues to review				
Trial progress	1. Recruitment rate (difference between the actual and expected rates of enrollment) is appropriate / reasonable. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
	2. The number and reasons of subject withdrawal is reasonable. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Protocol	3. All amendment of the research protocol and related documents have been reviewed by CREC without major concerns <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Risk	4. Risks to subjects are minimized. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA

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	5. Risks to subjects are reasonable in relation to anticipated benefits <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Equity	6. Selection of subjects is equitable. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Informed consent	7. The site is using the most recently approved version of informed consent document (<i>date stamps to indicate an approved version</i>). <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
	8. The informed consent document contains accurate, up-to-date information about the study. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
	9. Informed consent is sought from each subject or the subject's legally authorized representative appropriately. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Others	10. Where appropriate, the research adequately monitors the data collected to ensure the safety of the subjects. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
	11. Where appropriate, there are adequate provisions to protect the privacy of subjects and to maintain the confidentiality of data. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
	12. Appropriate additional safeguards are included to protect vulnerable subjects. <i>If no, please write a comment.</i>	☐ YES	☐ NO	☐ NA
Decision		For decision: approval		
☐ Approval ☐ Request for information ☐ Recommend further action		Frequency of continuing review	☐ 3 months ☐ 6 months ☐ 1 year	

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Comments:

REVIEWER'S SIGNATURE:
DATE:

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