



PROGRESS REPORT

CGHMC RERB Protocol No.	Sponsor Protocol No.
Protocol Title	
Investigator:	Contact Details:
Study Site	Approval Date: (dd/mm/yyyy)
Annual Progress Report: (Please indicate inclusive period): Total number of screened subjects: Total number of randomized subjects: Total number of withdrawn patients: Total number of SAEs: Total number of lost to follow up: Total number of completed subjects:	
Summary of Amendments	
Summary of protocol deviations/ non-compliance	
Is there any new information in recent literature or similar research that may change the risk/ benefit ratio for participants in this study? Summarize.	
Name and Signature of PI:	Date Submitted: (dd/mm/yyyy)

To be filled up by RERB

Date Received:	Received by: (Printed Name/Signature)	
Primary Reviewers/Signature:		Date
Recommendations <input type="checkbox"/> Approved <input type="checkbox"/> Request further information <input type="checkbox"/> Suspend or terminate the study <input type="checkbox"/> Others: _____		Type of review: <input type="checkbox"/> Expedited review <input type="checkbox"/> Full board review Date of meeting: _____
RERB Final Decision:		



Certified by: Name of CGHMC RERB Chair	Signature	Date
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CONTINUING REVIEW APPLICATION FORM (FORM 3.3B)

CGHMC RERB Protocol No.	Sponsor Protocol No.
Protocol Title	
Investigator	Contact details
Study Site	Approval Date: (dd/mm/yyyy)
ACTION REQUESTED: <input type="checkbox"/> Renew – New participant accrual to continue <input type="checkbox"/> Renew – Enrolled participant follow up only <input type="checkbox"/> Terminate – Protocol discontinued	
Any amendment since last review? <input type="checkbox"/> Yes <input type="checkbox"/> No (Discuss briefly)	
Any change in participant population recruitment or selection criteria since the last review? (Explain the changes if any) <input type="checkbox"/> Yes <input type="checkbox"/> No	
Any change in the Informed Consent process or documentation since the last review? (Please explain, if yes.) <input type="checkbox"/> Yes <input type="checkbox"/> No	



<p>Is there any new information in recent literature or similar research that may change the risk/ benefit ratio for participants in this study? (Discuss and attach a narrative.)</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Any unexpected complication or side effect noted since the last review? (Discuss and attach a narrative.)</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Did any participant withdraw from this study since the last approval? (Reasons for withdrawal)</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Any new investigator that has been added to or removed from the research team since the last review? (Please identify them and submit the CVs of new investigators.)</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>
<p>Are there any new collaborating sites that have been added or deleted since the last review? Please identify the sites and note the addition or deletion.</p>
<p>Summary of protocol participants:</p> <p><input type="checkbox"/> Reached the target number of participants approved by CGHMC RERB</p> <p><input type="checkbox"/> New participants recruited/enrolled since last review</p> <p><input type="checkbox"/> Total participants screened since protocol began</p> <p><input type="checkbox"/> Total participants enrolled and ongoing since the last review</p> <p><input type="checkbox"/> Total number of SAEs since the last review</p> <p><input type="checkbox"/> Total number of participants withdrawn or terminated</p>
<p>ACCRUAL EXCLUSIONS</p> <p><input type="checkbox"/> None</p>



- ☐ Male
- ☐ Female
- ☐ Others (Specify) _____

Impaired participants

- ☐ None
- ☐ Physically
- ☐ Cognitively
- ☐ Both

Name and Signature of PI	Date
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To be filled up by RERB

Date Received:	Received by: (Printed Name/Signature)	
Primary Reviewers/Signature:		Date:
<p>Recommendations</p> <p><input type="checkbox"/> Approved</p> <p><input type="checkbox"/> Request amendment to the protocol or the consent form</p> <p><input type="checkbox"/> Request further information</p> <p><input type="checkbox"/> Suspend or terminate the study</p> <p><input type="checkbox"/> Others: _____</p>		<p>Type of review:</p> <p><input type="checkbox"/> Expedited review</p> <p><input type="checkbox"/> Full board review</p> <p>Date of meeting: _____</p>
RERB Final Decision:		
Certified by: Name of CGHMC RERB Chair	Signature	Date