



FINAL REPORT

CGHMC RERB Protocol No.	Approval Date: (dd/mm/yyyy)
Protocol Title	
Principal Investigator	Contact Number
Study Site	
Total number of participants	Study Arms
Study materials <input type="checkbox"/> Drug <input type="checkbox"/> Device <input type="checkbox"/> Biologic specimen <input type="checkbox"/> Others	
Summary of Recruitment Accrual ceiling set by RERB New participants accrued since last review Total number of participants accrued since protocol began No. of participants who are lost to follow up No. of participants withdrawn from the study No. of participants who experienced SAEs/ SUSARs Number of participants who completed the study	
Duration of the Study	
Study objectives	
Summary of Findings in the Site	
Signature of Principal Investigator	Date Submitted

To be filled up by RERB

Date Received:	Received by: (Printed Name/Signature)
COMMENTS OF PRIMARY REVIEWER	

RECOMMENDED ACTION: <input type="checkbox"/> Accepted <input type="checkbox"/> Request additional Information: (Specify) <input type="checkbox"/> Recommend Further Action: (Specify) <input type="checkbox"/> Not accepted	Type of Review <input type="checkbox"/> Full Board <input type="checkbox"/> Expedited Meeting Date:
PRIMARY REVIEWER: _____ <div style="display: flex; justify-content: space-around; width: 100%;"> Signature over name Date </div> RERB CHAIR: _____ <div style="display: flex; justify-content: space-around; width: 100%;"> Signature over name Date </div>	