



LUNG CENTER OF THE PHILIPPINES

INSTITUTIONAL ETHICS REVIEW BOARD

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INFORMED CONSENT EVALUATION FORM

Submission Date	
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LCP-IERB CODE		Sponsor Protocol Number	
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Protocol Title	
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	Name	Mobile / Phone Number
Principal Investigator		
Sponsor		

INSTRUCTIONS:	Please evaluate how the elements outlined below have been appropriately addressed by the informed consent form (ICF), as applicable, and by confirming the submitted information and putting your comments in the space provided under "COMMENTS." In your comments, ensure that <u>vulnerability, recruitment process, and process of obtaining informed consent</u> are always assessed in the context of the study protocol and the participant. Finalize your review by indicating your decision, recommendations and signing in the space provided for the primary reviewer.	
1	Does the Informed Consent document state that the procedures are primarily intended for research? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
2	Are procedures for obtaining Informed Consent appropriate? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
3	Does the Informed Consent document contain comprehensive and relevant information? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
4	Is the information provided in the protocol consistent with those in the consent form? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:

5	Are study related risks mentioned in the consent form? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
6	Is the language in the Informed Consent document understandable? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
7	Is the Informed Consent translated into the local language/dialect? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
8	Is there adequate protection of vulnerable participants? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
9	Are the different types of consent forms (assent, patient representative) appropriate for the types of study participants? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
10	Are names and contact numbers from the research team and the IERB in the informed consent? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
11	Does the ICF mention privacy & confidentiality protection? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
12	Is there any inducement for participation? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
13	Is there provision for medical / psychosocial support? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
14	Is there provision for treatment of study-related injuries <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:
15	Is there provision for compensation? <input type="checkbox"/> Yes <input type="checkbox"/> No	Comments:

Decision	<input type="checkbox"/> Approval <input type="checkbox"/> Minor Revision <input type="checkbox"/> Major Revision / Resubmission <input type="checkbox"/> Disapproval
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Recommendations: (Identify items for revision)	
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Primary Reviewer	Signature	Date