Consent V10 Dated22mar24

Section I: Consent Wavier	
This only applies to patients who are unconscious or otherwise unable to provide consent and do not have a legally authorized representative. Patient participants who are included under the waiver of informed consent have to be approached and receive information about the study, the option to opt out, and consented for follow up when they regain consciousness or a legally authorized representative is present.	
Is the patient included under the waiver of informed consent?	
Reason for waiver of consent	 Whether the patient is unconscious or not able to provide consent Do not have a legally acceptable representative (LAR) Both
Please fill LAR attempt form to contact LAR	
Section II: Opt in consent for follow up data collection	
Who gave opt-in consent for follow-up data collection?	Patient participantLegally authorised representativeOpt-in consent not given
Please fill opt-out form	
Why was participant representative approached for consent instead of the participant?	☐ The participant is incapacitated because of the trauma☐ The participant is younger than 18 years
When did the participant or participant representative consented for follow-up data collection?	
How did the participant or participant representative consented for follow-up data collection?	○ In writing○ Verbally
Section III: Assent form	
Did the participant representative consented for follow-up data collection?	
Did the participant gave assent for follow-up?	YesNoThe participant is incapacitated because of the trauma

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