

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?	<input type="radio"/> Yes <input type="radio"/> No
Reason for waiver of consent	<input type="radio"/> Whether the patient is unconscious or not able to provide consent <input type="radio"/> Do not have a legally acceptable representative (LAR) <input type="radio"/> 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?	<input type="radio"/> Patient participant <input type="radio"/> Legally authorised representative <input type="radio"/> Opt-in consent not given
Please fill opt-out form	
Why was participant representative approached for consent instead of the participant?	<input type="checkbox"/> The participant is incapacitated because of the trauma <input type="checkbox"/> 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?	<input type="radio"/> In writing <input type="radio"/> Verbally

Section III: Assent form

Did the participant representative consented for follow-up data collection?	<input type="radio"/> Yes <input type="radio"/> No
Did the participant gave assent for follow-up?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> The participant is incapacitated because of the trauma