

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?

- ☐ Yes
☐ No

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 participant
☐ Legally authorised representative
☐ Opt-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?

- ☐ Yes
☐ No

Did the participant gave assent for follow-up?

- ☐ Yes
☐ No
☐ The participant is incapacitated because of the trauma