## Consent

Waiver of informed consent  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 egally authorized representative is present.	
Opt out from in-hospital data collection	
Did the participant or participant representative opt out from being included in the study?	
Date when participant decided to opt out from the study	
Please upload a scanned copy of the signed opt-out form	
Opt in consent for out of hospital follow up	
Who was approached for consent to out of hospital follow-up?	<ul> <li>Patient participant</li> <li>Legally authorised representative</li> <li>The patient is included under the waiver of informed consent</li> </ul>
Why was a representative of the participant approached for consent instead of the participant?	<ul> <li>The participant is incapacitated because of the trauma</li> <li>The participant is younger than 18 years</li> </ul>
Did the participant or participant representative consent to out of hospital follow-up?	
Did the participant assent to out of hospital follow-up?	<ul><li>Yes</li><li>No</li><li>The participant is incapacitated because of the trauma</li></ul>
When did the participant or participant representative consent to out of hospital follow-up?	
How did the participant or participant representative consent to out of hospital follow-up?	<ul><li>○ In writing</li><li>○ Verbally</li></ul>
Please upload a scanned copy of the signed consent	

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29-03-2024 3:37am