

IMPLEMENTATION STUDY CONSENT FORM

Full Title of Project: A Multi-Centre Randomised Controlled Trial of the Clinical and Cost-Effectiveness of Pre-Hospital Whole Blood Administration versus Standard Care for Traumatic Haemorrhage.

Short Title: SWiFT: Study of Whole blood In Frontline Trauma

Chief Investigators (CI): Dr Laura Green, Professor Jason Smith

Co-investigator for the Implementation study: Dr Grazia Antonacci

Please initial box

1. I confirm that I have read and understand the participant information sheet version ____ dated ____ / ____ /20____ for the above study and have had the opportunity to ask questions which have been answered fully.	
2. I understand that my participation is voluntary, and I am free to withdraw at any time, without giving any reason and without my legal rights being affected.	
3. I give permission for Imperial College London to access my research records relevant to this research.	
4. I give consent for information collected about me to be used to support other research in the future.	
5. I give consent to take part in the above study.	
6. I give consent to being contacted about potentially taking part in other research studies.	
7. I understand that data from the interview / focus group (including anonymised quotations) may be used in publications.	
8. I give permission for the interview / focus group meeting to be recorded. I understand that this recording will be deleted once data have been transcribed.	

Name of participant

Signature

Date

Name of person taking consent

Signature

Date