

[Hospital Contact Details]  
Name of PI  
Name of Hospital  
Hospital address  
Contact phone number  
Email



## CONSENT FORM THE I'M WOMAN TRIAL

**Title of research: Intramuscular tranexamic acid to prevent heavy bleeding after childbirth in women at higher risk**

|                                |  |                                     |  |
|--------------------------------|--|-------------------------------------|--|
| Site ID Number                 |  | Name of Site Principal Investigator |  |
| Participant Hospital ID number |  | Screening ID Number                 |  |
| Name of Participant            |  |                                     |  |

### STATEMENT OF PERSON GIVING CONSENT:

1. I confirm that I have read/have had read to me the information sheet for the above study in a language I understand.
2. I have discussed with the doctor to my satisfaction, and I have had the opportunity to ask questions.
3. I understand that my participation is voluntary. I have been given enough information about the research study to judge that I want to take part in it.
4. I understand that I am free to withdraw at any time, without giving any reason and without my medical care or legal rights being affected.
5. I understand that I will be given a copy of this consent form and the information sheet to keep for myself.
6. I understand that the study staff may look at sections of my medical notes and those of my baby/ies. I give permission for these individuals to have access to these records.
7. I understand that my data (with all personal information removed) will be made freely available for the public.
8. I give permission for a copy of this consent form, which contains my personal information, to be made available to the Trial Coordinating Centre in London for monitoring purposes only.
9. I agree to take part in the above study, the I'M WOMAN trial.

\_\_\_\_\_  
Name of woman

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature / Thumbprint or other mark (if unable to sign)

\_\_\_\_\_  
Name of witness

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature

*(A witness is needed if a patient cannot read or write)*

### STATEMENT OF PERSON OBTAINING INFORMED CONSENT:

I have fully explained this research to this participant and have given sufficient information, including about risks and benefits, to make an informed decision.

\_\_\_\_\_  
Name

\_\_\_\_\_  
Date

\_\_\_\_\_  
Signature