

# ENTRY FORM

PLEASE COMPLETE BEFORE RANDOMISING THE WOMAN

## ABOUT YOUR HOSPITAL *(please ensure all information below is contained in the medical records)*

<b>1.</b> Country	
<b>2.</b> Hospital code <i>(in your Study File)</i>	

## ABOUT THE PATIENT

<b>3.</b> Patient's initials <i>(first name/last name)</i>		<b>4.</b> Patient Hospital identification number				
<b>5.</b> Do you know her date of birth? <i>(circle)</i>	a) YES	day	month	year	b) NO – approximate age	years

## ABOUT THE DELIVERY

<b>6.</b> Delivery of the baby in this hospital? <i>(circle)</i>	YES	NO				
<b>7.</b> Type of delivery <i>(circle)</i>	VAGINAL	CAESAREAN SECTION				
<b>8.</b> Date of delivery	day	month	year			
<b>9.</b> Time of delivery <i>(24-hour clock)</i>	hours	minutes				
<b>10.</b> Placenta fully delivered? <i>(circle)</i>	YES	NO				
<b>11.</b> Primary cause of haemorrhage? <i>(circle)</i>	UTERINE ATONY	PLACENTA PRAEVIA/ACCRETA	SURGICAL TRAUMA/TEARS	OTHER	UNKNOWN	
<b>12.</b> Systolic Blood Pressure	mmHg <i>(most recent measurement prior to randomisation)</i>					
<b>13.</b> Estimated volume of blood loss	Millilitres <i>(estimated from delivery of baby to immediately prior to randomisation)</i>					
<b>14.</b> Any uterotonic prophylaxis given?	YES	NO	UNKNOWN			
<b>15.</b> Clinical signs of haemodynamic instability?	YES	NO	Haemodynamic instability assessment based on clinical signs <i>(e.g. low BP, tachycardia, falling urine output)</i> that requires an intervention <i>(e.g. intravenous fluids)</i>			

## RANDOMISATION INFORMATION

<b>16.</b> Eligible? <i>(circle)</i>	<b>YES</b> <i>(Get the lowest available number treatment pack and follow instructions)</i>		<b>NO</b> <i>(Do not randomise, record on screening log)</i>				
<b>17.</b> Consent obtained from? <i>(circle)</i>	Woman	Relative	Other representative	Waiver			
<b>18.</b> Insert treatment pack number here	<b>BOX</b>				<b>PACK</b>		
<b>19.</b> Date of randomisation	day	month	year	<b>20.</b> Time of randomisation <i>(24-hour clock)</i>	hours	minutes	
<b>21.</b> a) Name of person randomising patient				b) Signature			

**PLEASE SEND THIS FORM TO THE COORDINATING CENTRE IMMEDIATELY AFTER RANDOMISATION**

[SEE GUIDANCE OVERLEAF](#)

## DATA FORMS GUIDANCE

### AFTER COMPLETING THIS PAPER FORM, YOU CAN:

- ❖ Enter these data directly into the trial database (username and password required)  
[www.thewomantrial.Lshtm.ac.uk](http://www.thewomantrial.Lshtm.ac.uk)
- ❖ Complete an Electronic Data Form (EDF) and send by email or upload to the trial intranet at  
[www.thewomantrial.Lshtm.ac.uk](http://www.thewomantrial.Lshtm.ac.uk)
- ❖ Send as a secure scanned document by email to [woman.data@Lshtm.ac.uk](mailto:woman.data@Lshtm.ac.uk)
- ❖ Fax to +44 20 7299 4663
- ❖ Store original form in the Investigator's Study File
- ❖ **PLEASE GIVE A COPY OF THIS COMPLETED FORM TO THE PERSON RESPONSIBLE FOR COMPLETING THE OUTCOME FORM AT YOUR HOSPITAL**

<b>NOTES:</b>

<b>FOR ADVERSE EVENTS, UNBLINDING AND OTHER URGENT ENQUIRIES PLEASE TELEPHONE +44(0)7768 707500</b>
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**PLEASE NOTE: IF YOUR QUERY IS NOT URGENT PLEASE USE THE NORMAL CONTACT DETAILS IN THE INVESTIGATOR'S STUDY FILE.**