

UKOSS

UK Obstetric Surveillance System

Therapies or Prophylaxis for Peripartum Haemorrhage Study 04/07

Data Collection Form - CASE

Please report all women delivering after 1st September 2007 and before 1st October 2008

Case Definition:

A woman treated with **any** of the following therapies for management of peripartum haemorrhage:

EITHER Factor VIIa

OR B-Lynch or other brace suture

OR Arterial ligation or embolisation or intra-arterial ballons (including prophylactic catheter placement prior to delivery)

Please return the completed form to:

**UKOSS
National Perinatal Epidemiology Unit
University of Oxford
Old Road Campus
Oxford
OX3 7LF**

**Fax: 01865 289701
Phone: 01865 289714**

Case reported in: _____



Royal College of
Obstetricians and
Gynaecologists

Instructions

1. Please do not enter any personally identifiable information (e.g. name, address or hospital number) on this form.
2. Please record the ID number from the front of this form against the woman's name on the Clinician's Section of the blue card retained in the UKOSS folder.
3. Fill in the form using the information available in the woman's case notes.
4. Tick the boxes as appropriate. If you require any additional space to answer a question please use the space provided in section 7.
5. Please complete all dates in the format DD/MM/YY, and all times using the 24hr clock e.g. 18:37
6. If codes or examples are required, some lists (not exhaustive) are included on the back page of the form.
7. If you do not know the answers to some questions, please indicate this in section 7.
8. If you encounter any problems with completing the form please contact the UKOSS Administrator or use the space in section 7 to describe the problem.

Section 1: Woman's details

1.1 Year of birth

1.2 Ethnic group^{1*} (enter code, please see back cover for guidance)

1.3 Marital status

single married cohabiting

1.4 Was the woman in paid employment at booking?

Yes No

If Yes, what is her occupation

If No, what is her partner's (if any) occupation

1.5 Height at booking (cm)

1.6 Weight at booking (kg)

1.7 Smoking status

never gave up prior to pregnancy
current gave up during pregnancy

Section 2: Previous Pregnancies

2.1 Gravidity

Number of completed pregnancies 24 weeks and beyond

Number of pregnancies less than 24 weeks

If no previous pregnancies, please go to section 3.

If the woman has had previous pregnancies please indicate whether any of the following were present:

Pregnancy or delivery problems^{2*}

Yes No

If Yes, please specify

Any previous deliveries by caesarean section

Yes No

If Yes, please specify number in total

Was the immediately preceding delivery by caesarean section?

Yes No

*For guidance please see back cover

Section 3: Previous Medical History

3.1 Did the woman have any previous or pre-existing medical problems?^{3*} Yes No

If Yes, please specify _____

3.2 Previous uterine surgery Yes No

If Yes, please specify type and number of operations

Evacuation of retained products of conception (ERPC) Yes Number

Dilatation and curettage Yes Number

Surgical termination of pregnancy Yes Number

Myomectomy Yes Number

Manual removal of placenta Yes Number

Other^{4*} Yes Number

If Other, please specify _____

3.3 Previous uterine perforation Yes No

If Yes, please specify treatment of perforation, if any _____

Section 4: This Pregnancy

4.1 Final Estimated Date of Delivery (EDD)^{5*} DD / MM / YY

4.2 Was this pregnancy a multiple pregnancy? Yes No

If Yes, please specify number of fetuses _____

4.3 Were there problems in this pregnancy?^{2*} Yes No

If Yes, please specify _____

4.4 Was placenta praevia diagnosed prior to delivery? Yes No

If Yes, please specify grade _____

4.5 Was placenta accreta/increta/percreta suspected prior to delivery? Yes No

If Yes, how was it diagnosed?

Ultrasound MRI Other

If Other, please specify _____

Section 5: Delivery

5.1 Was delivery induced? Yes No

If Yes, please state indication _____

Was vaginal prostaglandin/misoprostol used? Yes No

5.2 Did the woman labour? Yes No

If Yes, please state date and time of diagnosis of labour DD / MM / YY hh : mm

Was syntocinon used during labour? Yes No

Duration of syntocinon during labour [] hrs [] mins

5.3 Was delivery by caesarean section? Yes No

If Yes, please state grade of most senior operator _____

What was the indication for caesarean section? _____

Method of anaesthesia (tick all that apply)

Epidural Single-shot spinal Continuous spinal CSE General

5.4 What was used for third stage prophylaxis? (tick all that apply)

None Syntocinon Syntometrine Other (please specify) _____

Haemorrhage

5.5 What was the estimated blood loss (total mls)?

5.6 Please indicate what treatments were undertaken

	Tick all that apply	Please rank the therapies in the order in which they were first used (1, 2, 3 etc)	Was this therapy used for prophylaxis (P) or treatment (T)*. Please tick (P) or (T)	
			(P)	(T)
Syntocinon infusion	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Ergometrine	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Prostaglandin F2α	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Misoprostol	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Intra-abdominal packing	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Intrauterine balloons	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Intrauterine packing	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Recombinant factor VIIa	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Vessel embolisation/ligation	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Intra-arterial balloons	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
B-Lynch or other brace suture	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Hysterectomy (please tick) Total <input type="checkbox"/> Subtotal <input type="checkbox"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>
Other (please specify) _____	<input type="checkbox"/>	<input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="checkbox"/>

5.7 Was a B-Lynch or other compression suture used?

Yes No

If No, please go question 5.8

If Yes, what technique was used (please tick)

- Unclear
- Classical B-Lynch
- Modified B-Lynch (Two vertical sutures, uterus not opened)
- Multiple vertical compression sutures
- Square sutures
- Systematic devascularisation of uterus
- Other

If Other, please specify _____

Date and time of procedure / / :

How many units of blood had been transfused by the start of the procedure?

What was the time between the decision to use a compression suture and the time the suture was placed (min)

*For guidance please see back cover

5.8 Was recombinant factor VIIa (Novoseven) used?

Yes No

If No, please go question 5.9

If Yes, please indicate total number of units transfused before rf VIIa was given

Blood

Fresh Frozen Plasma (FFP)

Platelets

Cryoprecipitate

Date and time of VIIa first used? / / :

Dose of rf VIIa given (first) (mg) .

What was the time between the decision to use rfVIIa and the time it was first given? (min)

Total number of doses Total dose given (mg) .

5.9 Is arterial embolisation (interventional radiology) routinely available in your unit?

Yes No

If Yes, when and where is it available (tick all that apply)

Same hospital Different hospital

During normal working hours Out of hours

5.10 Was major blood vessel catheterisation or ligation carried out on this woman?

Yes No

If No, please go to question 5.11

If Yes, which technique was used? Balloon Embolisation Ligation

When were catheters placed? / / :

Which vessels were occluded? _____

If embolisation was carried out, what material was used? _____

Where was the procedure performed? Same hospital Different hospital

Date and time of procedure? / / :

How many units of blood had been transfused by the start of the procedure?

What was the time between the decision to perform embolisation/ligation and the time the procedure was performed (min)

5.11 What was the primary underlying cause of haemorrhage (please tick one only)

Uterine atony

Placenta praevia

Placenta accreta/increta/percreta

Placental abruption

Uterine infection

Uterine rupture

If Yes, please specify pre-labour during labour traumatic

Extension of incision at time of caesarean section

Extension of previous caesarean section scar at the time of caesarean section

Genital tract trauma/tears

Other cause

If Other, please specify _____

5.12 Did the woman refuse transfusion of blood products?

Yes No

5.13 Please record the amounts of blood products received in total by this woman (units)

	Total (units)
Whole blood or packed red cells	<input type="text"/> <input type="text"/>
Fresh Frozen Plasma (FFP)	<input type="text"/> <input type="text"/>
Platelets	<input type="text"/> <input type="text"/>
Cryoprecipitate	<input type="text"/> <input type="text"/>
Cell salvaged blood (ml)	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>

5.14 Was the woman actively warmed during treatment for haemorrhage?

Yes No

5.15 Were the intravenous fluids actively warmed?

Yes No

5.16 Was the woman's temperature monitored?

Yes No

If Yes, what was her lowest recorded temperature? .

5.17 Was CVP monitoring used?

Yes No

5.18 Was intra-arterial monitoring used?

Yes No

Section 6: Outcomes

Section 6a: Woman

6a.1 Please indicate whether any of the following morbidities occurred (tick all that apply)

Adult respiratory distress syndrome

Pulmonary oedema

Disseminated intravascular coagulopathy (DIC)

Renal failure requiring dialysis

Cardiac arrest

Pulmonary embolism

DVT

Other thrombosis

If Other thrombosis, please specify _____

6a.2 Was the woman admitted to ITU?

Yes No

If Yes, duration of stay (days)

Or Tick if woman is still in ITU

Or Tick if woman was transferred to another hospital

6a.3 Did any other major maternal morbidity occur?^{8*}

Yes No

If Yes, please specify _____

6a.4 Did the woman die?

Yes No

If Yes, please specify date of death

/ /

What was the primary cause of death as stated on the death certificate?

Section 6b: Infant 1

NB: If more than one infant, for each additional infant, please photocopy the infant section of the form (before filling it in) and attach extra sheet(s) or download additional forms from the website: www.npeu.ox.ac.uk/ukoss

6b.1 Date and time of delivery

/ / : 24hr

6b.2 Mode of delivery

spontaneous vaginal ventouse lift-out forceps rotational forceps
breech pre-labour caesarean section caesarean section after onset of labour

6b.3 Birthweight (g)

6b.4 Was the infant stillborn?

Yes No

If Yes, was this

Antepartum OR Intrapartum

Please go to section 7

6b.5 Was the infant admitted to the neonatal unit?

Yes No

6b.6 Did this infant die?

Yes No

If Yes, please specify date of death

/ /

What was the primary cause of death as stated on the death certificate?

(please state if not known) _____

Section 7

Please use this space to enter any other information you feel may be important

Section 8:

Name of person completing the form _____

Designation _____

Today's date

/ /

You may find it useful in the case of queries to keep a copy of this form.

If you are unable to make a copy please tick the box

Definitions

1. UK Census Coding for ethnic group

WHITE

01. British
02. Irish
03. Any other white background

MIXED

04. White and black Caribbean
05. White and black African
06. White and Asian
07. Any other mixed background

ASIAN OR ASIAN BRITISH

08. Indian
09. Pakistani
10. Bangladeshi
11. Any other Asian background

BLACK OR BLACK BRITISH

12. Caribbean
13. African
14. Any other black background

CHINESE OR OTHER ETHNIC GROUP

15. Chinese
16. Any other ethnic group

2. Current or previous pregnancy problems, including:

Pre-eclampsia (hypertension and proteinuria)

Eclampsia

Thrombotic event

Amniotic fluid embolism

3 or more miscarriages

Preterm birth or mid trimester loss

Neonatal death

Stillbirth

Baby with a major congenital abnormality

Small for gestational age (SGA) infant

Large for gestational age (LGA) infant

Infant requiring intensive care

Puerperal psychosis

Placenta praevia

Gestational diabetes

Significant placental abruption

Post-partum haemorrhage requiring transfusion

3. Previous or pre-existing maternal medical problems, including:

Essential hypertension

Cardiac disease (congenital or acquired)

Renal disease

Endocrine disorders e.g. hypo or hyperthyroidism

Psychiatric disorders

Haematological disorders e.g. sickle cell disease, diagnosed thrombophilia

Inflammatory disorders e.g. inflammatory bowel disease

Epilepsy

Diabetes

Autoimmune diseases

Cancer

HIV

4. Examples of other previous uterine surgery:

Myomectomy

Endometrial resection/ablation

Septal resection

Polypectomy

5. Estimated date of delivery (EDD):

Use the best estimate (ultrasound scan or date of last menstrual period) based on a 40 week gestation

6. Definition of prophylaxis and treatment:

(P) Prophylaxis or support: following haemorrhage, other treatments given and considered successful, but this therapy add 'just in case' to support other interventions.

(T) Treatment/rescue: following haemorrhage, other treatments given and considered to have failed, so this therapy is given as rescue.

7. Major maternal medical complications, including:

Persistent vegetative state

Cerebrovascular accident

Pulmonary oedema

Mendelson's syndrome

Renal failure

Thrombotic event

Septicaemia

Required ventilation