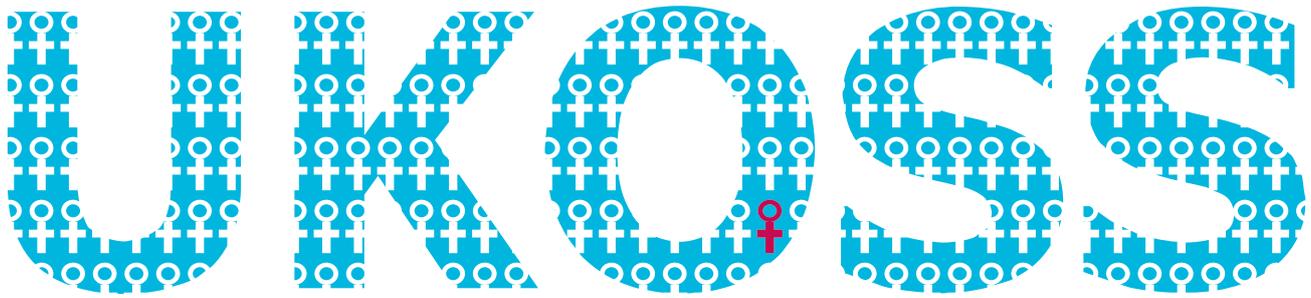


ID Number:



UK Obstetric Surveillance System

# Spontaneous Haemoperitoneum in Pregnancy (SHiP) Study 01/16

Data Collection Form - CASE

**Please report any woman delivering between 1<sup>st</sup> Jan 2016 and 31<sup>st</sup> Dec 2017.**

## Case Definition:

Any woman 20 weeks or more gestation with sudden intra-abdominal haemorrhage requiring surgery (CS, laparotomy, laparoscopy), without preceding trauma

EXCLUDE: women with uterine rupture, trauma.



Royal College of  
Obstetricians  
and Gynaecologists

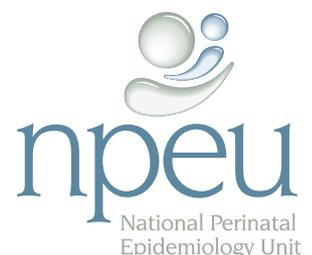
Bringing to life the best  
in women's health care

Please return the completed form to:

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

**Fax: 01865 617775**  
**Phone: 01865 289714**

**Case reported in:** \_\_\_\_\_



## 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 the woman has not yet delivered, please complete the form as far as you are able, excluding delivery and outcome information, and return to the UKOSS Administrator. We will send these sections again for you to complete two weeks after the woman's expected date of delivery.
- 8. If you do not know the answers to some questions, please indicate this in section 7.**
9. 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:<sup>1\*</sup> (enter code, please see back cover for guidance)

1.3 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:

---

1.4 Height at booking:

   cm

1.5 Weight at booking:

   .  kg

1.6 Smoking status:

never  gave up prior to pregnancy   
current  gave up during pregnancy

## Section 2: Previous Obstetric History

### 2.1 Gravidity

Number of completed pregnancies beyond 24 weeks:

Number of pregnancies less than 24 weeks:

If no previous pregnancies, please go to section 3

2.2 Did the woman have any previous pregnancy problems?<sup>2\*</sup> Yes  No

If Yes, please specify: \_\_\_\_\_

\_\_\_\_\_

2.3 Has this woman had any previous deliveries by caesarean section? Yes  No

## Section 3: Previous Medical History

3.1 Does this woman have a known history of endometriosis? Yes  No

If Yes, what year was it diagnosed:

OR tick if not known

Was it histologically confirmed? Yes  No  Not known

3.2 Has this woman had prior abdominal surgery? Yes  No

If Yes, please specify surgery undertaken: \_\_\_\_\_

and indication: \_\_\_\_\_

Was this surgery for endometriosis? Yes  No  Not known

3.3 Did the woman have any other pre-existing medical problems<sup>3\*</sup>? Yes  No

If Yes, please give details: \_\_\_\_\_

\_\_\_\_\_

## Section 4: This Pregnancy

4.1 Final Estimated Date of Delivery (EDD):<sup>4\*</sup>  /  /

4.2 Was this a multiple pregnancy? Yes  No

If Yes, please specify number of fetuses:

4.3 What was the planned mode of delivery for this pregnancy prior to diagnosis of SHiP? Vaginal (including trial of labour)

Abdominal (elective caesarean section)

4.4 Did the woman receive any anticoagulation in this pregnancy? Yes  No

If Yes, please indicate anticoagulants received and timing relative to diagnosis

Anticoagulant used (please tick all that apply)	Was this for prophylaxis (P) or Treatment (T)?	Was this anticoagulation received prior to diagnosis?
Aspirin <input type="checkbox"/>	<input type="checkbox"/>	Yes <input type="checkbox"/> No <input type="checkbox"/>
Heparin (inc. LMWH) <input type="checkbox"/>	<input type="checkbox"/>	Yes <input type="checkbox"/> No <input type="checkbox"/>
Warfarin <input type="checkbox"/>	<input type="checkbox"/>	Yes <input type="checkbox"/> No <input type="checkbox"/>
Thrombolysis <input type="checkbox"/>	<input type="checkbox"/>	Yes <input type="checkbox"/> No <input type="checkbox"/>

4.5 What date was SHiP was first diagnosed? / /

4.6 What were the symptoms prior to diagnosis? (please tick all that apply)

abdominal pain  altered uterine contractions   
haematuria  vaginal bleeding   
fetal heart rate abnormality  Other

If Other, please specify: \_\_\_\_\_

4.7 What was the initial presumed diagnosis? \_\_\_\_\_

4.8 How was the haemoperitoneum diagnosed? (please tick all that apply)

Peritoneal lavage  Ultrasound   
CT  CTPA   
MRI  At surgery

4.9 What mode of surgery was used to manage the haemorrhage? (please tick one)

laparoscopy  planned caesarean section or hysterotomy   
(i.e. delivery of baby intended at onset of surgery)  
laparotomy  emergency caesarean section or hysterotomy   
(delivery of baby not planned at the start of laparotomy)

4.10 Were there signs of active endometriosis at the time of surgery for SHiP? Yes  No

4.11 At the time of surgery, was the pregnancy noted to be: intrauterine  extrauterine

4.12 What was the identified source/location of bleeding? \_\_\_\_\_

4.13 What was the estimated total blood loss?  ml

4.14 What was the estimated intraperitoneal blood loss? <500ml  ≥500ml

4.15 What was the woman's lowest measured haemoglobin value?  g/dl

OR tick if not measured

**4.16 Did the woman refuse blood products?**

Yes  No

**If No**, were blood products given?

Yes  No

**If Yes**, please state total units of each: *(enter zero if none given)*

Whole blood or packed red cells

Fresh Frozen Plasma (FFP)

Platelets

Cryoprecipitate

Cell salvaged blood (ml)

**4.17 Were any haemostatic drugs used?**

Yes  No

**If Yes**, please tick all that apply:

fibrinogen  Factor VII

Tranexamic acid  Other

**If Other**, please specify: \_\_\_\_\_

**4.18 Were there any other problems in this pregnancy?<sup>2\*</sup>**

Yes  No

**If Yes**, please specify: \_\_\_\_\_

## Section 5: Delivery

**5.1 Did this woman have a miscarriage?**

Yes  No

**If Yes**, please specify date:

 /  / 

**5.2 Did this woman have a termination of pregnancy (including hysterotomy)?**

Yes  No

**If Yes**, please specify date:

 /  / 

**If Yes to 5.1 or 5.2, please go to sections 6a, 7 and 8**

**5.3 Is this woman still undelivered?**

Yes  No

**If Yes**, will she be receiving the rest of her antenatal care from your hospital?

Yes  No

**If No**, please indicate name of hospital providing future care:

\_\_\_\_\_

Will she be delivered at your hospital?

Yes  No

**If No**, please indicate name of delivery hospital, then go to Section 7

\_\_\_\_\_

**5.4 Was delivery induced?**

Yes  No

**If Yes**, please state indication: \_\_\_\_\_

**5.5 Did the woman labour?**

Yes  No

**5.6 Was delivery by caesarean section?**

Yes  No

**If Yes**, please state:

Grade of urgency:<sup>5\*</sup>

Indication for caesarean section: \_\_\_\_\_

Method of anaesthesia:

Regional  General anaesthetic

**5.7 Was any placental abnormality identified?**

Yes  No

If Yes, please tick which abnormalities were identified:

Praevia

Accreta

Increta/percreta

Other

If Other, please specify: \_\_\_\_\_

## Section 6: Outcomes

### Section 6a: Woman

**6a.1 Was the woman admitted to ITU or level 3 care?**

Yes  No

If Yes, duration of stay:

days

OR Tick if woman is still in ITU or level 3 care:

OR Tick if woman was transferred to another hospital:

**6a.2 Did any other major maternal morbidity occur?<sup>6\*</sup>**

Yes  No

If Yes, please specify: \_\_\_\_\_

**6a.3 Did the woman die?**

Yes  No

If Yes, please specify date and time of death

/  /   :   
24hr

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

(Please state if not known.) \_\_\_\_\_

Was a post mortem examination undertaken?

Yes  No

If Yes, did the examination confirm the certified cause of death/diagnosis?

Yes  No  Not known

### 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](http://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 Sex of infant:**

Male  Female  Indeterminate

**6b.5 Was the infant stillborn?**

Yes  No

If Yes, please go to section 7.

**6b.6 5 min Apgar**



## 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. Previous or current pregnancy problems, including

Thrombotic event  
Amniotic fluid embolism  
Eclampsia  
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  
Surgical procedure in pregnancy  
Hyperemesis requiring admission  
Dehydration requiring admission  
Ovarian hyperstimulation syndrome  
Severe infection e.g. pyelonephritis

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

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  
Autoimmune diseases  
Cancer  
HIV

### 4. Estimated date of delivery (EDD)

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

### 5. RCA/RCOG/CEMACH/CNST Classification for urgency of caesarean section:

1. Immediate threat to life of woman or fetus
2. Maternal or fetal compromise which is not immediately life-threatening
3. Needing early delivery but no maternal or fetal compromise
4. At a time to suit the woman and maternity team

### 6. Major maternal medical complications, including:

Persistent vegetative state  
Cardiac arrest  
Cerebrovascular accident  
Adult respiratory distress syndrome  
Disseminated intravascular coagulopathy  
HELLP  
Pulmonary oedema  
Mendleson's syndrome  
Renal failure  
Thrombotic event  
Septicaemia  
Required ventilation

### 7. Fetal/infant complications, including:

Respiratory distress syndrome  
Intraventricular haemorrhage  
Necrotising enterocolitis  
Neonatal encephalopathy  
Chronic lung disease  
Severe jaundice requiring phototherapy  
Major congenital anomaly  
Severe infection e.g. septicaemia, meningitis  
Exchange transfusion