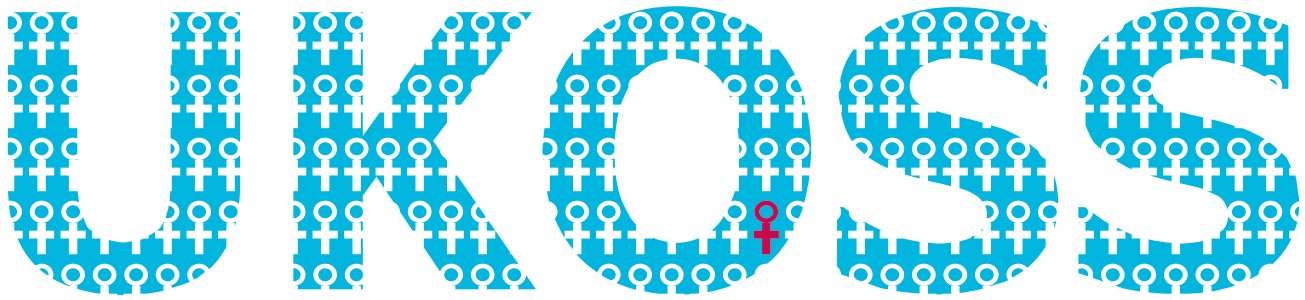


ID Number:



UK Obstetric Surveillance System

HELLP Syndrome Study 02/11

Data Collection Form - Control

Instructions for selecting control women

1. Identify the date and time of delivery for the woman you have reported who has had HELLP Syndrome. This woman is the CASE.
2. From the delivery suite/operating theatre records identify the two pregnant women delivering immediately **BEFORE** the woman who has had HELLP Syndrome (these women should NOT have had HELLP Syndrome). These women will act as the CONTROLS.
3. Please retrieve the hospital case notes for these control women from medical records.
4. Please complete a Control Data Collection Form (this form) for each of the women you have identified as the controls.
5. You will also have been sent a Case Data Collection Form. Please complete the case form with information about the woman who has had HELLP Syndrome.



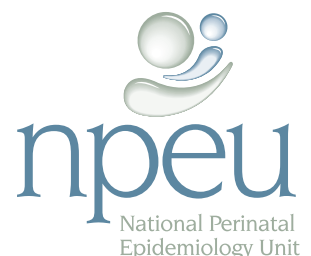
Royal College of
Obstetricians and
Gynaecologists

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:^{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 and weight at booking:** cm AND kg
- 1.6 Smoking status:** never gave up prior to pregnancy
current gave up during pregnancy

*For guidance please see back cover

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 What was the date of delivery/termination/miscarriage in the most recent previous pregnancy:

/ /

2.3 Please indicate if any of the following were present in previous pregnancies: (Please tick all that apply)

Pregnancy induced hypertension (PIH) Pre-eclampsia Eclampsia

HELLP syndrome Gestational diabetes

2.4 Did the woman have any other previous pregnancy problems?^{2*}

Yes No

If Yes, please specify: _____

Section 3: Previous Medical History

3.1 Did the woman have essential hypertension at booking or prior to pregnancy?

Yes No

If Yes, was she receiving anti-hypertensive medication at booking or prior to pregnancy?

Yes No

3.2 Does the women have pre-existing diabetes mellitus? Type 1 Type 2 Neither

3.3 Did the women have any other previous or pre-existing medical problems?^{3*}

Yes No

If Yes, please specify: _____

Section 4: This Pregnancy

4.1 Final Estimated Date of Delivery (EDD):^{4*}

/ /

4.2 Was this a multiple pregnancy?

Yes No

If Yes, please specify number of fetuses:

4.3 Date of booking:

/ /

4.4 What was the platelet count at booking?

x10⁹/L

*For guidance please see back cover

4.5 Was the woman diagnosed with any of the following in this pregnancy?

	Yes	No	Date of diagnosis
Pregnancy induced hypertension (PIH)	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY
Pre-eclampsia	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY
Eclampsia	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY
Gestational diabetes	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY

4.6 What were the levels of the following in this pregnancy or tick if not recorded:

	Tick if booking level not recorded	Level at booking	Tick if highest level not recorded	Highest Level	Date of highest level recorded
Systolic BP (mmHg)	<input type="checkbox"/>	□□□	<input type="checkbox"/>	□□□	DD / MM / YY
Diastolic BP (mmHg)	<input type="checkbox"/>	□□□	<input type="checkbox"/>	□□□	DD / MM / YY
Proteinuria (please indicate units)	<input type="checkbox"/>	_____	<input type="checkbox"/>	_____	DD / MM / YY

4.7 Were there any other problems in this pregnancy?^{2*}

Yes No

If Yes, please specify: _____

4.8 Was the woman given corticosteroids?

Yes No

If Yes, please specify:

Agent	Dose	Units	Indication	Date started
_____	_____	_____	_____	DD / MM / YY

4.9 Was any antihypertensive medication commenced/continued in this pregnancy (antenatally or postnatally)?

Yes No

If Yes, please specify:

Name of drug	Date treatment started
_____	DD / MM / YY
_____	DD / MM / YY

4.10 Were any of the following treatments commenced/continued in this pregnancy (antenatally or postnatally)?

	Yes	No	Date treatment started
Magnesium sulphate	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY
Aspirin	<input type="checkbox"/>	<input type="checkbox"/>	DD / MM / YY

*For guidance please see back cover

4.11 Was any other medication commenced/continued in this pregnancy (antenatally or postnatally)?

Yes No

If Yes, please specify:

Name of medication	Indication	Date treatment started
_____	_____	<input type="text" value="DD"/> <input type="text" value="DD"/> / <input type="text" value="MM"/> <input type="text" value="MM"/> / <input type="text" value="YY"/> <input type="text" value="YY"/>
_____	_____	<input type="text" value="DD"/> <input type="text" value="DD"/> / <input type="text" value="MM"/> <input type="text" value="MM"/> / <input type="text" value="YY"/> <input type="text" value="YY"/>

4.12 Did the women refuse blood products?

Yes No

If No, were blood products given?

Yes No

4.13 Were any of the following used for thromboprophylaxis? (please tick all that apply)

	Antenatally	Postnatally
TED Stockings	<input type="checkbox"/>	<input type="checkbox"/>
Low molecular weight heparin	<input type="checkbox"/>	<input type="checkbox"/>

4.14 Did the woman develop any overt clinical signs of coagulopathy (non-obstetric bleeding)? E.g. petechiae, haematuria, bleeding gums

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?

Yes No

If Yes, please specify date:

/ /

5.3 Was delivery induced?

Yes No

If Yes, please state indication: _____

5.4 Did the woman labour?

Yes No

5.5 Was delivery by caesarean section?

Yes No

If Yes, please state:

Grade of urgency:^{5*} _____

Indication for caesarean section: _____

Method of anaesthesia: Regional General anaesthetic

Section 6: Outcomes

Section 6a: Woman

6a.1 Was the woman admitted to ITU (critical care level 3) or obstetric HDU? Yes No

If Yes, duration of stay: days

OR Tick if woman is still in ITU/HDU:

OR Tick if woman was transferred to another hospital:

6a.2 Did the woman require ventilation? Yes No

6a.3 Did the woman require haemodialysis? Yes No

If Yes, for how long was she dialysed? days

6a.4 Did the woman have hepatic encephalopathy Yes No

6a.5 Was the woman transferred to a liver unit? Yes No

6a.6 Did any other major maternal morbidity occur?^{6*} Yes No

If Yes, please specify: _____

6a.7 Has the woman been discharged from hospital? Yes No

If Yes, what was the date of the woman's discharge from hospital? / /

Was the woman readmitted after discharge? Yes No Not known

If Yes, what was the reason for readmission? _____

6a.8 Did the woman die? Yes No

If Yes, please specify date and time of death / / :

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 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

6b.1 Date and time of delivery: / / :

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

6b.7 Was the infant admitted to the neonatal unit? Yes No

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, for example:

3 or more miscarriages
Acute fatty liver
Amniotic fluid embolism
Ante-partum haemorrhage requiring transfusion
Baby with major congenital problem
Hyperemesis requiring admission
IUGR/small for gestational age
Neonatal death
Placenta praevia
Placental abruption
Placenta accreta/percreta/increta
Post-partum haemorrhage requiring transfusion
Preterm birth or mid-trimester loss
Severe infection (e.g. pyelonephritis)
Stillbirth (IUD)
Surgical procedure in pregnancy
Significant antepartum haemorrhage
Thrombotic event (DVT/Pulmonary embolus/Stroke)

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

Auto-immune disease
Cancer
Cardiac disease (congenital or acquired)
Epilepsy

Endocrine disorders, e.g. Hypo or hyperthyroidism,
Haematological disorders e.g. sickle cell disease,
diagnosed thrombophilia

Inflammatory disorders e.g. Inflammatory bowel
disease

Psychiatric disorders

Renal disease

Thrombotic event (pulmonary embolism)

Coagulopathy

Polycystic ovary disease

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, for example:

Adult respiratory distress syndrome
Cardiac arrest
Cerebrovascular accident/intercranial haemorrhage
Convulsions – not diagnosed as eclampsia
Disseminated intravascular coagulopathy (DIC)
Deranged clotting – not DIC
Multiple organ failure
Persistent vegetative state/anoxic/hypoxic brain injury
Pulmonary oedema
Septicaemia/septic shock
Thrombotic event

7. Infant complications, for example:

Chronic lung disease
Exchange transfusion
Intraventricular haemorrhage
Major congenital anomaly
Multiorgan failure
Necrotising enterocolitis
Neonatal encephalopathy/HIE/birth asphyxia
Respiratory distress syndrome/Ventilated/
Pneumothorax/Chest effusions/Haemothorax
Severe infection e.g. septicaemia, meningitis
Severe jaundice requiring phototherapy