



UK Obstetric Surveillance System

Amniotic Fluid Embolism Study 01/15

Data Collection Form - CASE

Case Definition:

EITHER a clinical diagnosis of AFE (acute hypotension or cardiac arrest, acute hypoxia or coagulopathy in the absence of any other potential explanation for the symptoms and signs observed)

OR a pathological diagnosis (presence of fetal squames or hair in the lungs).

Case ID Number:



Royal College of
Obstetricians
and Gynaecologists

Bringing to life the best
in women's health care

Please return the completed form to:

ukoss@npeu.ox.ac.uk

UKOSS

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

Phone: 01865 617764 / 617774

Reporting Month: _____

Reporting Hospital: _____



NPEU

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 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 Obstetric History

- 2.1 Gravidity**
Number of previous completed pregnancies beyond 24 weeks
Number of previous pregnancies less than 24 weeks
If no previous pregnancies, please go to section 3.
- 2.2 Did the woman have any previous pregnancy problems?^{2*}** Yes No
If Yes, please specify _____

Section 3: Previous Medical History

- 3.1 Please indicate whether the woman had any of the following previous or pre-existing medical conditions:**
- | | | | | |
|---|-----|--------------------------|----|--------------------------|
| History of allergy | Yes | <input type="checkbox"/> | No | <input type="checkbox"/> |
| History of atopy (asthma, eczema, hayfever) | Yes | <input type="checkbox"/> | No | <input type="checkbox"/> |
| Essential hypertension | Yes | <input type="checkbox"/> | No | <input type="checkbox"/> |
| Diabetes mellitus | Yes | <input type="checkbox"/> | No | <input type="checkbox"/> |
- 3.2 Did the woman have any other pre-existing medical problems?^{3*}** Yes No
If Yes, please specify details _____

*For guidance please see back cover

Section 4a: This Pregnancy

4a.1 Final Estimated Date of Delivery (EDD)^{4*}

DD / MM / YY

4a.2 Was this pregnancy a multiple pregnancy?

Yes No

If Yes, specify number of fetuses

4a.3 Was placenta praevia diagnosed?

Yes No

If Yes, please specify the grade

4a.4 Did the woman have a placental abruption?

Yes No

4a.5 Did the woman develop any hypertensive disorder?

Yes No

If Yes, please specify

		Date of onset	Time of onset
Pregnancy induced hypertension	<input type="checkbox"/>	DD / MM / YY	
Pre-eclampsia (hypertension and proteinuria)	<input type="checkbox"/>	DD / MM / YY	
Eclampsia	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Other	<input type="checkbox"/>	DD / MM / YY	

If Other, please specify _____

4a.6 Did the woman have chorioamnionitis?

Yes No

4a.7 Did the woman have polyhydramnios?

Yes No

4a.8 Did the woman develop gestational diabetes?

Yes No

4a.9 Were there any **other** problems in this pregnancy?^{2*}

Yes No

If Yes, please specify _____

Section 4b: Diagnosis of amniotic fluid embolism

4b.1 Please indicate if any of the following features were present at or immediately preceding diagnosis

	Tick all that apply	Please rank the features in order of occurrence (1,2,3,etc)
Acute fetal compromise	<input type="checkbox"/>	<input type="checkbox"/>
Cardiac arrest	<input type="checkbox"/>	<input type="checkbox"/>
Cardiac rhythm problems	<input type="checkbox"/>	<input type="checkbox"/>
Coagulopathy	<input type="checkbox"/>	<input type="checkbox"/>
Hypotension	<input type="checkbox"/>	<input type="checkbox"/>
Maternal haemorrhage	<input type="checkbox"/>	<input type="checkbox"/>
Premonitory symptoms e.g. restlessness, agitation, numbness, tingling	<input type="checkbox"/>	<input type="checkbox"/>
Seizure	<input type="checkbox"/>	<input type="checkbox"/>
Shortness of breath	<input type="checkbox"/>	<input type="checkbox"/>

4b.2 Was an echocardiogram done following collapse?

Yes No

If Yes, did the woman have abnormal echocardiogram findings?

Yes No

If Yes, please indicate what the abnormal findings were? _____

Section 4c: Laboratory tests

Please specify the first results after diagnosis and the worst haematological parameters recorded at the time of the AFE or tick if not recorded?

	Diagnosis value	Tick if diagnosis value not recorded	Worst value	Tick if worst value not recorded
Hb g/dL	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>
Platelet count (x10 ⁹ /L)	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>
PT (sec)	<input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>
INR	<input type="text"/> . <input type="text"/>	<input type="checkbox"/>	<input type="text"/> . <input type="text"/>	<input type="checkbox"/>
APTT (sec)	<input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>
APTT (ratio) APTT	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>
Fibrinogen (g/dL)	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>
D-dimer (ng/ml)	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/>
Tryptase (µg/l)	<input type="text"/> <input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>	<input type="text"/> <input type="text"/> <input type="text"/> . <input type="text"/>	<input type="checkbox"/>

Section 4d: Maternal event

4d.1 Date and time of event

 / / : ^{24hr}

4d.2 Date and time diagnosis first considered

 / / : ^{24hr}

4d.3 Were membranes ruptured at time of event?

Yes No

If Yes, please state date and time of rupture

 / / : ^{24hr}

Was rupture

Artificial Spontaneous

4d.4 Was there meconium staining of liquor?

Fresh Old No

4d.5 Was there fetal distress before maternal collapse?

Yes No

If Yes, please specify _____

4d.6 Was woman:

Not in labour First stage Second stage Post-delivery

4d.7 What was the contraction frequency at time of event? (number in 10 mins)

4d.8 Did the woman have any anaesthetic/analgesia at time of collapse?

Spinal Epidural Combined spinal epidural GA None

4d.9 Were any of the following clinical staff present at the time of collapse?

If No, please indicate date and time first saw woman after collapse

OR Tick if did not see woman

Consultant obstetrician

Yes No

 / / : ^{24hr}

Consultant anaesthetist

Yes No

 / / : ^{24hr}

Senior midwife
(band 7 or above)

Yes No

 / / : ^{24hr}

Section 4e: Management

4e.1 Please indicate what treatments were undertaken, when they were first used and total units/dose given where applicable

	Tick all that apply	Date first given	Time first given	Total dose	Units
Syntocinon infusion	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Ergometrine	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Prostaglandin F2α	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Misoprostol	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Hemabate	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Whole blood or packed red cells	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Cryoprecipitate	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Fresh frozen plasma	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Platelets	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Cell salvage	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Fibrinogen	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Factor VIIa	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Heparin	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Tranexamic acid	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____
Other	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>	_____	_____

If Other, please specify _____

	Tick all that apply	Date	Time
Intrauterine balloons	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Intrauterine packing	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
B-lynch or other brace suture	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Vessel embolisation	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Vessel ligation	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Intra-arterial balloons	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Hysterectomy	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Intra-abdominal packing	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Exchange transfusion	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Plasma exchange	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>
Apheresis	<input type="checkbox"/>	DD / MM / YY	hh : mm <small>24hr</small>

*For guidance please see back cover

Section 5: Delivery

5.1 Was delivery induced?

Yes No

If Yes, what was the reason for induction? _____

If Yes, was vaginal prostaglandin used? Yes No If Yes, please record the preparation and total dose of prostaglandin given (mg)

5.2 Did the woman labour?

Yes No If Yes, what date and time was labour diagnosed? / / : : Was syntocinon used during labour? Yes No Duration of syntocinon during labour : Did hyperstimulation occur? (contractions more than 5 in 10 minutes) Yes No If Yes, for how long did hyperstimulation occur? hrs mins

5.3 Was delivery by caesarean section?

Yes No If Yes, please state whether Elective OR Emergency Grade of urgency^{5*}

Indication for caesarean section _____

Method of anaesthesia: Regional General anaesthetic

5.4 Did the woman have manual removal of her placenta?

Yes No

Section 6: Outcomes

Section 6a: Woman

6a.1 Was the woman admitted to ITU/HDU?

Yes No If Yes, please indicate date and time of admission: / / : : Duration of stay daysOr Tick if woman is still in ITU/HDU Or Tick if woman was transferred to another hospital

6a.2 Did the woman have permanent neurological injury (e.g. hypoxic brain injury, persistent vegetative state)?

Yes No

If Yes, please give details _____

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

Yes No

If Yes, please specify _____

6a.4 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, were fetal squames or hair found in the lungs? _____

*For guidance please see back cover

Section 6b: Infant

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 Sex of infant

Male Female Indeterminate

6b.5 Was the infant stillborn?

Yes No

If Yes, was this

Ante-partum **OR** Intra-partum

If Yes, go to section 7

6b.6 5 min Apgar

6b.7 Was the infant admitted to the neonatal unit?

Yes No

If Yes, please state the duration of stay

days

Or Tick if the infant is still in the neonatal unit

Or Tick if the infant was transferred to another hospital

6b.8 Did any other major infant complications occur?*

Yes No

If Yes, please specify details _____

6b.9 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.

*For guidance please see back cover

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:

3 or more miscarriages
Amniocentesis
Baby with a major congenital abnormality
Gestational diabetes
Haemorrhage
Hyperemesis requiring admission
Infant requiring intensive care
Neonatal death
Placenta praevia
Placental abruption
Post-partum haemorrhage requiring transfusion
Pre-eclampsia (hypertension and proteinuria)
Premature rupture of membranes
Preterm birth or mid trimester loss
Puerperal psychosis
Thrombotic event
Severe infection e.g. pyelonephritis
Stillbirth
Surgical procedure in pregnancy

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

Cardiac disease (congenital or acquired)
Diabetes
Epilepsy
Endocrine disorders e.g. hypo or hyperthyroidism
Essential hypertension
Haematological disorders e.g. sickle cell disease, diagnosed thrombophilia
Inflammatory disorders e.g. inflammatory bowel disease
Psychiatric disorders
Renal 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 morbidity, including:

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

7. Fetal/infant complications, including:

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