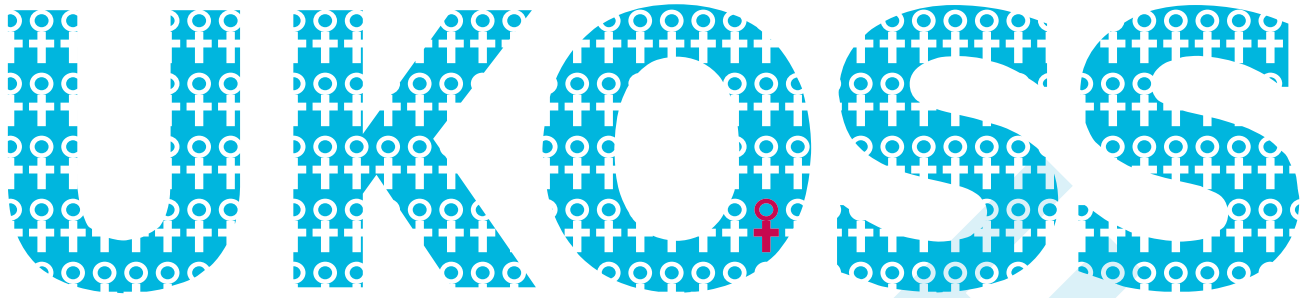


ID Number:



UK Obstetric Surveillance System

Maternal Pulmonary Aspiration in Pregnancy Study 04/13

Data Collection Form - CASE

Please report any woman delivering on or after 01/09/2013 and
before 01/09/15

Case Definition:

Please report any woman with a final diagnosis of pulmonary aspiration during pregnancy or delivery up to postpartum discharge from hospital.

Maternal pulmonary aspiration includes women with the following features

- Women who have had an unprotected airway while unconscious, semi-conscious or paralysed

AND

- A clinical history consistent with regurgitation of stomach contents and pulmonary aspiration (e.g vomiting after induction of anaesthesia or gastric contents seen in the oropharynx)

AND

- Symptoms / signs of respiratory compromise requiring supplementary oxygen and antibiotics or level 2 or level 3 (HDU or ITU) respiratory support, in the absence of any other clear cause

Classical radiological findings may or may not be present



Royal College of
Obstetricians
and Gynaecologists

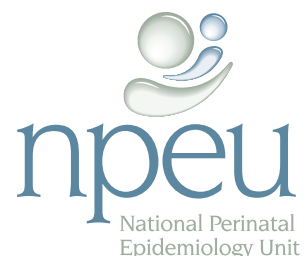
Bringing to life the best
in women's health care

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^{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 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 problems in her previous pregnancies?^{2*}** Yes No
If Yes, please specify _____

Section 3: Previous Medical History

- 3.1 Did the woman have any of the following?** Yes No
If Yes, please tick all that apply
Hiatus hernia Known history of aspiration
Known swallowing problems Gastro-oesophageal reflux disease (GORD) Epilepsy
- 3.2 Does the woman have any respiratory illness, recurrent chest infections or other chest complaints?** Yes No
If Yes, please specify _____
- 3.3 Does the woman have any other pre-existing medical problems^{3*}** Yes No
If Yes, please specify _____

*For guidance please see back cover

Section 4: This Pregnancy

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

/ /

4.2 Was this pregnancy a multiple pregnancy?

Yes No

If Yes, specify number of fetuses

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

Yes No

If Yes, please specify _____

4.4 What are the usual hospital guidelines for oral intake during active labour? (please tick one for high risk and one for low risk women)

	High risk	Low risk
Restricted intake (clear fluids only)	<input type="checkbox"/>	<input type="checkbox"/>
Light diet	<input type="checkbox"/>	<input type="checkbox"/>
Other (please specify) _____	<input type="checkbox"/>	<input type="checkbox"/>

4.5 Does the hospital have a fasting policy prior to elective surgery?

Yes No

If Yes, for how many hours are women recommended to fast prior to surgery?

Section 5: Diagnosis and Delivery

Section 5a: Diagnosis of Aspiration

5a.1 What was the date and time the woman was admitted prior to the aspiration event?

/ / :

5a.2 What was the date and time that aspiration was diagnosed?

/ / :

5a.3 Where was the woman when the aspiration occurred? (tick one only)

- Home / community Delivery room Labour ward anaesthetic room
 Labour ward theatre Labour ward recovery area Labour ward high dependency area
 Elsewhere in the hospital (please specify) _____

5a.4 Which of the following features were noted prior to or immediately following the presumed aspiration event: (tick all that apply)

- Vomiting
 Gastric contents seen in oropharynx
 Gastric contents seen on pillow
 Low oxygen saturation
 Unprotected airway at induction/intubation for general anaesthesia
 Unprotected airway at extubation following general anaesthesia
 Unprotected airway while semi-conscious

If ticked, please specify circumstances (e.g. in recovery after anaesthesia, following epileptic or eclamptic fit): _____

5a.5 Was a chest X-ray done?

Yes No

If Yes, was consolidation noted?

Yes No

Were any other abnormalities noted?

Yes No

If Yes, please specify _____

5a.6 Was any other cause for the symptoms and signs identified? Yes No

If Yes, please specify _____

5a.7 Did the woman have any oral intake in the 6 hours prior to the aspiration event? Yes No Not known

If Yes, please record what was eaten/drunk (*tick one only*)

Food and fluids Fluids only Clear fluids only

Was the date and time of the last oral intake known? Yes No

If Yes, please give date and time of last known oral intake / / : : 24hr

5a.8 Was antacid prophylaxis against aspiration prescribed in the 6 hours prior to aspiration? (e.g. metoclopramide, H2 antagonist, proton pump inhibitor, sodium citrate) Yes No

If Yes, please state:

Drug(s) name	Dose	Units	Date and time of administration
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>

5a.9 Did the woman receive any opioids in the 6 hours prior to aspiration? (e.g. morphine, pethidine) Yes No

If Yes, please state:

Drug(s) name	Dose	Units	Date and time of administration
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>
_____	_____	_____	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> : <input type="text"/> <input type="text"/> <small>24hr</small>

Section 5b: Aspiration Associated with General Anaesthesia

5b.1 Was the aspiration event associated with general anaesthesia? Yes No

If No, please go to section 5c.

If Yes, please continue

5b.2 Did a consultant anaesthetist administer the anaesthetic? Yes No

If No, what was the grade of the anaesthetist administering the anaesthetic? _____

Was a consultant anaesthetist present? Yes No

5b.3 Was an airway device in use at the time of aspiration? Yes No

If Yes, please specify (*tick one only*)

Face mask Oropharyngeal airway Classic laryngeal mask airway

Other supraglottic airway device (*please specify with trade name*) _____

Endotracheal tube with cuff inflated Endotracheal tube with cuff NOT inflated

Other (*please specify*) _____

5b.4 Was the anaesthetic associated with the aspiration given for caesarean section? Yes No

If No, for what procedure was anaesthesia given? _____

5b.5 Were any of the following mechanisms used to reduce the aspiration risk before, during or after anaesthesia Yes No

If Yes, tick all that apply

Nasogastric or orogastric tube insertion and stomach drainage

Rapid sequence induction

Application of cricoid pressure during induction

Release of cricoid pressure to assist airway insertion

Extubation in the left lateral position

Extubation when awake

Section 5c: Management of Aspiration

5c.1 Following aspiration, were any of the following performed? Yes No

If Yes, tick all that apply

Intubation Bronchoscopy Cricoid pressure Oropharyngeal suction

Patient placed in left-lateral, head-down position Supplemental oxygen

5c.2 Were antibiotics given? Yes No

If Yes, please state:

Drug(s) name	Dose	Units	Route	Duration (days)
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____

5c.3 Were any steroid or bronchodilator drugs given? Yes No

If Yes, please state:

Drug(s) name	Dose	Units	Route
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____
_____	_____	_____	_____

Section 5d: Delivery

5d.1 Did this woman have a miscarriage? Yes No

If Yes, please specify date

/ /

5d.2 Did this woman have a termination of pregnancy? Yes No

If Yes, please specify date

/ /

5d.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 the name of the hospital providing future care

Will she be delivered at your hospital?

Yes No

If No, please indicate the name of delivery hospital then, complete ONLY sections 6a, 7 and 8

5d.4 Was delivery induced?

Yes No

If Yes, please state indication? _____

5d.5 Did the woman labour?

Yes No

5d.6 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 HDU (critical care level 2)?

Yes No

If Yes, please specify:

Duration of stay _____

days

Or Tick if woman is still in HDU (critical care level 2)

Or Tick if woman was transferred to another hospital

6a.2 Was the woman admitted to ITU (critical care level 3)?

Yes No

If Yes, please specify:

Duration of stay _____

days

Or Tick if woman is still in ITU (critical care level 3)

Or Tick if woman was transferred to another hospital

6a.3 Did the woman require mechanical ventilation after the aspiration?

Yes No

6a.4 Did the woman require extracorporeal membrane oxygenation (ECMO)?

Yes No

6a.5 Did the woman have a cardiorespiratory arrest?

Yes No

If Yes, please give date and time of arrest

/ / : :

24hr

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

Yes No

If Yes, please specify _____

6a.7 Was the woman discharged following the admission in which she had the aspiration event?

Yes No

If Yes, please give date of discharge

/ /

6a.8 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 cause of death?

Yes No Not known

*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

D	D	/	M	M	/	Y	Y	h	h	:	m	m
---	---	---	---	---	---	---	---	---	---	---	---	---

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 Intra-partum

If Yes, go to section 7

6b.6 Did the infant require resuscitation at birth?

Yes No

6b.7 5 min Apgar

--	--

6b.8 Was the infant admitted to the neonatal unit?

Yes No

6b.9 Did any major infant complications occur?*

Yes No Unknown

If Yes, please specify _____

6b.10 Did this infant die?

Yes No

If Yes, please specify date and time of death

D	D	/	M	M	/	Y	Y	h	h	:	m	m
---	---	---	---	---	---	---	---	---	---	---	---	---

24hr

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

D	D	/	M	M	/	Y	Y
---	---	---	---	---	---	---	---

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:

Hospital admissions during the pregnancy, either related to the pregnancy or not related

Thrombotic event (PE/ DVT)

Amniotic fluid embolism

Pre-eclampsia

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

Pulmonary aspiration

HELLP

Anaphylaxis

Obstetric cholestasis

Placenta accreta/ increta/ percreta

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

Cardiac disease (congenital or acquired)

Renal disease

Endocrine disorders e.g. hypo or hyperthyroidism, diabetes

Psychiatric disorders

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

Inflammatory disorders e.g. inflammatory bowel disease

Autoimmune diseases

Cancer

HIV

Immunosuppressive disorders

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:

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