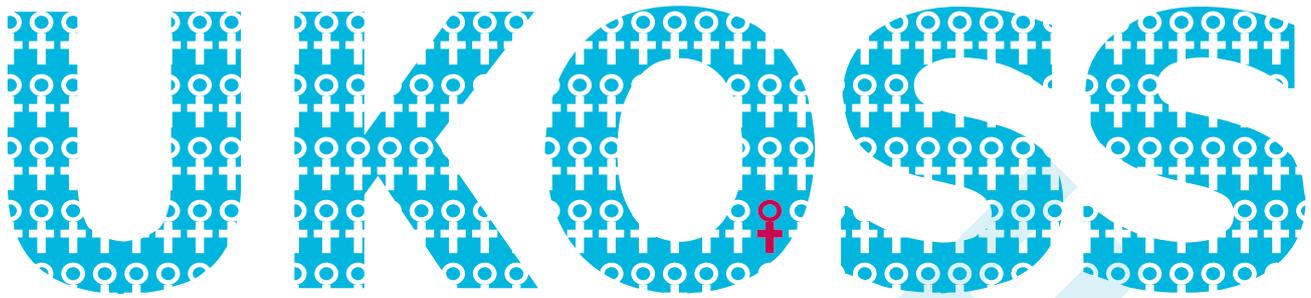


ID Number:



UK Obstetric Surveillance System

## Pregnancy in women with stage 5 Chronic Kidney Disease (chronic renal failure) Study 01/12

### Data Collection Form - CASE

Please report any woman delivering on or after 1<sup>st</sup> February 2012 and before 1<sup>st</sup> January 2014.

#### Case Definition:

Please report any pregnant woman with stage 5 Chronic Kidney Disease (chronic renal failure). This would usually include any pregnant woman in one of the following groups:

- A woman with an estimated glomerular filtration rate (eGFR) <15mls/min/1.73m<sup>2</sup> pre-pregnancy
- A woman receiving peritoneal or haemodialysis at conception
- A woman with a serum creatinine >300µmol/l pre-pregnancy
- A woman with a serum creatinine >250µmol/l on two or more occasions during pregnancy
- A woman commenced on peritoneal or haemodialysis to treat chronic kidney disease during this pregnancy



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 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 previous pregnancy problems (including kidney problems during previous pregnancies)?<sup>2\*</sup>** Yes  No   
If Yes, please specify: \_\_\_\_\_

## Section 3: Previous Medical History

**3.1** What was the underlying disease or condition causing chronic kidney disease?

\_\_\_\_\_

**3.2** Has this woman received a renal transplant?

Yes  No

If Yes, what was the date of the most recent transplant?

/   /

**3.3** Has this woman previously received dialysis?

Yes  No

If Yes, on what date was dialysis first commenced?

/   /

Was this during a previous pregnancy?

Yes  No

What was the indication for starting dialysis? \_\_\_\_\_

What were the following values when dialysis was commenced?

	Value	Unit	Date	Not Recorded
K+	_____	mmol/l	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/>	<input type="checkbox"/>
Urea	_____	mmol/l	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/>	<input type="checkbox"/>
Creatinine	_____	μmol/l	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/>	<input type="checkbox"/>
Bicarbonate	_____	mmol/l	<input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/> / <input type="text"/> <input type="text"/>	<input type="checkbox"/>

Was there proteinuria prior to pregnancy?

Yes  No

If Yes, what was the

Albumin/Creatinine Ratio (ACR) \_\_\_\_\_

OR Protein/Creatinine Ratio (PCR) \_\_\_\_\_

**3.4** What were the following values prior to pregnancy?

	Value	Unit	Not Recorded
Volume of urine per 24h	_____	ml	<input type="checkbox"/>
Most recent diastolic BP	_____	mmHg	<input type="checkbox"/>
Most recent serum creatinine	_____	μmol/l	<input type="checkbox"/>
eGFR	_____	mls/min/1.73m <sup>2</sup>	<input type="checkbox"/>

**3.5** Did the woman have any other pre-existing medical problems?<sup>3\*</sup>

Yes  No

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

## Section 4: This Pregnancy

### Section 4a: Antenatal care and management

4a.1 Final Estimated Date of Delivery (EDD)\*\*

/   /

4a.2 Was antenatal care undertaken in the usual hospital for this woman's area of residence?

Yes  No

If No, please indicate below reasons for care at a different hospital (*tick all that apply*)

Referred to a tertiary centre because of underlying medical condition  Patient preference

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

4a.3 Was this a multiple pregnancy?

Yes  No

If Yes, please specify number of fetuses: \_\_\_\_\_

4a.4 Did this woman conceive while taking ACE inhibitors (e.g. captopril)?

Yes  No

4a.5 Did this woman receive Erythrocyte Stimulating Agents (ESA)/ Erythropoetin (EPO)?

Yes  No

If Yes, please give agent and maximum dose: (e.g. aranesp, micera)

Agent

Dose

Frequency

\_\_\_\_\_

4a.6 Did the woman receive intravenous iron during pregnancy?

Yes  No

If Yes, please give dose and number of doses:

Dose

Number

\_\_\_\_\_

4a.7 Did this woman receive any of the following during pregnancy?

Yes  No

If Yes, please tick all that apply Aspirin  LMWH (in addition to anticoagulation on dialysis)

Unfractionated heparin  Vitamin D (cholecalciferol or adcal D3)

### Section 4b: Dialysis therapy

4b.1 Has this woman received dialysis in this pregnancy?

Yes  No

If Yes, please indicate whether any of the following dialysis therapies were used in this pregnancy (*Please tick all that apply*)

If more than one dialysis treatment was given please give dates of changes

	Used	Maximum hours per week during pregnancy	Date of change
Peritoneal Dialysis	<input type="checkbox"/>	<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"/>
Haemodialysis	<input type="checkbox"/>	<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"/>
Nocturnal dialysis	<input type="checkbox"/>	<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"/>
Haemodiafiltration (Dialysis using individually prescribed replacement fluid)	<input type="checkbox"/>	<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"/>

Were there any admissions for dialysis related events?

Yes  No

If Yes, please tick all that apply:

Hypotension  Line infection  Line not working  Fistula or graft clotting

Please give details of access for dialysis (*Please tick all that apply*)

	Used	Date started	Still used	Date ended
Line 1	<input type="checkbox"/>	DD / MM / YY	<input type="checkbox"/>	DD / MM / YY
Line 2	<input type="checkbox"/>	DD / MM / YY	<input type="checkbox"/>	DD / MM / YY
Line 3	<input type="checkbox"/>	DD / MM / YY	<input type="checkbox"/>	DD / MM / YY
Fistula	<input type="checkbox"/>	DD / MM / YY	<input type="checkbox"/>	DD / MM / YY
Graft (type of fistula)	<input type="checkbox"/>	DD / MM / YY	<input type="checkbox"/>	DD / MM / YY

4b.2 Please indicate the number of antihypertensive drugs used:

number

Prior to pregnancy

First trimester (up to 14 weeks)

Second trimester (14-28 weeks)

Third trimester (after 28 weeks)

## Section 4c: Laboratory results and complications

4c.1 Please record the levels of the following:

	Highest serum urea (mmol/l)	Lowest serum urea (mmol/l)	Lowest haemoglobin (g/dl)
First trimester (up to 14 weeks)	_____	_____	_____
Second trimester (14-28 weeks)	_____	_____	_____
Third trimester (after 28 weeks)	_____	_____	_____

4c.2 Did the woman have a serum creatinine >250µmol/l on two or more occasions in this pregnancy?

Yes  No

4c.3 Did the woman have polyhydramnios (Amniotic Fluid Index >20cm) diagnosed at any point in pregnancy?

Yes  No

4c.4 Was pre-eclampsia (or superimposed pre-eclampsia) diagnosed in this pregnancy?<sup>5\*</sup>

Yes  No

If Yes – what was:

Highest systolic blood pressure (mmHg)?

Highest diastolic blood pressure (mmHg)?

4c.5 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?**

Yes  No

If Yes, please specify date:

/   /

If Yes to 5.1 or 5.2, please now complete 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 the 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>6\*</sup>

Indication for caesarean section:

\_\_\_\_\_

Method of anaesthesia:

Regional

General anaesthetic

## 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>7\*</sup>**

Yes  No

If Yes, please specify:

\_\_\_\_\_

**6a.3 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.)

\_\_\_\_\_

\*For guidance please see back cover

## 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:**

/   /   :

**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

**6b.8 Did any other major infant complications occur?<sup>8\*</sup>**

Yes       No

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

**6b.9 Did this infant 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) \_\_\_\_\_

## Section 7:

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

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

## Section 8:

**8.1 Name of person completing the form:** \_\_\_\_\_

**8.2 Designation:** \_\_\_\_\_

**8.3 Today's date:**   /   /

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

## 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. Definition of pre-eclampsia:

Systolic BP  $\geq 140$  and/or diastolic BP  $\geq 90$  and proteinuria  $\geq 300\text{mg}/24\text{hrs}$  (30mg/mmol Protein creatinine ratio). If hypertension already present - the new onset of proteinuria; if proteinuria already present - the new onset of hypertension; if both hypertension and proteinuria present - the development of one additional clinical or biochemical feature of pre-eclampsia e.g abnormal LFTs)

### 6. 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

### 7. 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  
Thrombotic event  
Septicaemia  
Required ventilation

### 8. 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