

UK Obstetric Surveillance System

Cirrhosis in pregnancy Study 01/17

Data Collection Form - CASE

Case Definition:

Any woman giving birth in the UK between the 1st of June 2017 and 31st of May 2018 identified as having hepatic cirrhosis.

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.
- 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.



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: _



Sec	ction 1: Woman's details
1.1	Year of birth:
1.2	Ethnic group: ^{1*} (enter code, please see back cover for guidance)
1.3	
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:
1.6	Weight at booking:
Sec	ction 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 this woman's liver function deteriorate during a previous pregnancy? Yes No
2.3	Has this woman ever had any of the following: <i>Please tick all that apply</i>
	A pre-term delivery (<37 weeks) An infant admitted to the neonatal unit
	A stillbirth None of the above
2.4	Has this woman ever had any of the following in previous pregnancies:
	Please tick all that apply
	Pre-eclampsia HELLP Obstetric Cholestasis
	Acute Fatty Liver of Pregnancy Gestational diabetes None of the above
2.5	Did the woman have any other previous pregnancy problems? ^{2*} Yes No
	If Yes, please specify:
Sec	ction 3: Previous Medical History
3.1	When was cirrhosis diagnosed?
3.2	What was the diagnosis of cirrhosis based on:
	a) Liver biopsy: Yes No
	b) CT, MRI or Ultrasound scan: Yes No
3.3	Does the woman have portal hypertension: Yes No Not known
	If Yes, please give the date of diagnosis
3.4	Which underlying liver disease does this woman have (please tick one):

4	which underlying liver disease does this woman have (please lick one).
	Autoimmune Hepatitis 🦳 Primary Sclerosing Cholangitis 🗌 Primary Biliary Cirrhosis 🗌
	Hepatitis B Hepatitis C Portal Vein Thrombosis Other Not known

If Other, please specify: _

3.5	3.5 Did the woman have oesophageal varices before this pregnancy:					
		jj		Yes	No 🗌 Not	known
	If Yes, what was the si	ze of the varices	at the last pre-p	regnancy endosco	ppy (please tick	(one):
				ledium 📄 III - L		known
3.6	Has this woman ever	had treatment f	or a variceal bl	eed: Yes	No 🗌 Not	known
	If Yes, did she have:		Band	Ligation: Yes	No Not	known
			Injection Sclere	otherapy: Yes	No Not	known
3.7	Has this woman had	a liver transplan	it:		Yes	No 🗌
	If Yes, please give the	year of transplan	t:		[YYYY
3.8	Does the woman have	e any history of	gallstones:	Yes	No 🗌 Not	known
3.9	Has this woman ever	had drug induc	ed cholestasis:	Yes	No 🗌 Not	known
	If Yes, please give the	name of the drug	responsible: _			
3.10	Does this woman hav	ve any other prev	vious / pre-exis	ting medical con	ditions: Yes	No
	If Yes, please specify:					
3.11	Did this woman have with a specialist know				No 🗌 Not	known
	•	•				
Sec	tion 4: This Pregr	nancy				
4.1	Final Estimated Date	of Delivery (EDI	D): ^{4*}		D D / M	ΜΥΥ
4.2	Was this a multiple p	regnancy?			Yes	No
	If Yes, please specify r	number of fetuses	3:			
4.3	Was conception?			Sponta	aneous 🗌 A	ssisted
4.4	What was the date of	the first (booking	ng) appointmen	t in pregnancy?	D D / M	ΜΥΥ
4.5	Regarding this woma	n's medication,	were any of the	e following: Pleas	se tick all that a	apply
			Stopped	Stopped	Started	
		Taken at conception	prior to	following	during	Never taken
		• •	conception	conception	pregnancy	
	Ursodeoxycholic acid					
	Steroids					
	Vitamin K					
	Immunosuppression					
	(specify which)					

4.6	Were any other drugs not listed above taken during pregnancy for
	underlying liver disease?

Yes 📃 No 🛛

If Yes, please list drugs taken _

4.7 Did this woman have any of the following symptoms during pregnancy:

Did this woman have any of th	le following symptoms	s during pregnan	icy.
	Tick all that apply		se provide the date first ed in pregnancy
Pruritus		D) / M M / Y Y
Jaundice		D) / M M / Y Y
Ascites		D) / M M / Y Y
Gastrointestinal bleeding		D	
Encephalopathy		D) / M M / Y Y
None of the above			
8 Were any of the following bloc Please tick all that apply	od tests abnormal at th	ne time of conce	ption?
Raised ALT?	Lo	w Albumin?	
Raised Bilirubin?	Lo	w Sodium?	
Raised Bile Acids?	Lo	w Haemaglobin?	
Raised Creatinine?	Lo	w Platelets?	
Raised Prolonged prothrombin	time? No	one of the above	
9 Please give details of the wor	st levels of the followir	ng blood tests du	uring pregnancy
w	orst level Or	tick if not known	Date
ALT			
Bilirubin			D D / M M / Y Y
Bile Acids			
Albumin			D D / M M / Y Y
Creatinine			
Sodium			DD/MM/YY
Haemoglobin			
Platelets			DD/MM/YY
Prothrombin time			
10 Please give details of the pre-	delivery levels of the f	ollowing blood to	ests
Pre-delive	ery level Or t	tick if not known	Date
ALT			
Bilirubin			D D / M M / Y Y
Bile Acids			
Albumin			D D / M M / Y Y
Creatinine			
Sodium			DD/MM/YY
Haemoglobin			
Platelets			DD/MM/YY
Prothrombin time			

4.11	Did this woman have any of the following pregnancy problems: Please tick all that apply			
	Pre-eclampsia Pregnancy induced hypertension HELLP			
	Cholestasis AFLP Gestational diabetes None of the above			
4.12	Did the woman have a post-partum haemorrhage:YesNo			
	If Yes, please give estimated blood loss ml			
4.13	Did the woman have a diagnostic endoscopy during pregnancy:			
	Yes No Not known			
	If Yes, did she have varices Yes No Not known			
	If Yes, what was the size of the varices (please tick one):			
	0 - None I - Small II - Medium III - Large Not known			
	Was any treatment given during the procedure? (please tick one)			
	None Sclerotherapy Banding Not known			
4.14	Did the woman have a variceal bleed during pregnancy: Yes No			
	If Yes, please give date of bleed and treatment (indicate banding/sclerotherapy/octreotide/none)			
	Date of bleed Treatment			
4.15	Did this woman have a liver ultrasound scan during pregnancy: Yes No			
4.15	Did this woman have a liver ultrasound scan during pregnancy: Yes No If Yes, what was the size of the spleen Image: margin content of the spleen Image: margin content of the spleen			
4.15 4.16				
	If Yes, what was the size of the spleen			
	If Yes, what was the size of the spleen			
	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes No If Yes, what was the grade of encephalopathy: None Mild Moderate Severe Coma Not known What treatment did she receive?			
	If Yes, what was the size of the spleen			
4.16 4.17	If Yes, what was the size of the spleen			
4.16 4.17	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No			
4.16 4.17	If Yes, what was the size of the spleen			
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4.16 4.17 4.18	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No If Yes, please specify:			
4.16 4.17 4.18 Sec	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No If Yes, please specify:			
4.16 4.17 4.18	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No If Yes, please specify: Toto 5: Delivery Did this woman have a miscarriage? Yes No			
4.16 4.17 4.18 Sec 5.1	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No If Yes, please specify:			
4.16 4.17 4.18 Sec	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Yes No If Yes, please specify: Yes No If Yes, please specify date: Did this woman have a miscarriage? Yes No Yes No Yes No Yes No Yes No Yes No <p< th=""></p<>			
4.16 4.17 4.18 Sec 5.1	If Yes, what was the size of the spleen Did the woman have an episode of encephalopathy during pregnancy: Yes None Mild Moderate Severe Coma Not known What treatment did she receive? Did this woman have a liver transplant during the pregnancy: Yes No If Yes, please give date Were there any other problems in this pregnancy? Yes No If Yes, please specify:			

5.3	Is this woman still undelivered? If Yes, will she be receiving the rest of her antenatal care from your hospital? If No, please indicate name of hospital providing future care:	Yes No Yes No
	Will she be delivered at your hospital? If No, please indicate name of delivery hospital, then <i>go to Section</i> 7	Yes No
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: ^{7*}	
	Indication for caesarean section:	
	Method of anaesthesia: Regional Gene	eral anaesthetic
Sec	ction 6: Outcomes	
Sec	ction 6a: Woman	
6a.1		Yes No
	If Yes, duration of stay: OR Tick if woman is still in ITU (critical care level 3):	days
6- 0	OR Tick if woman was transferred to another hospital:	
6a.2	Did any other major maternal morbidity occur? ^{6*}	Yes No
	If Yes, please specify:	
6a.3	Did the woman die?	Yes No
	If Yes, please specify date and time of death	YY hh mm 24hr
	(Please state if not known)	
	ction 6b: Infant 1	
NB:	If more than one infant, for each additional infant, please photocopy the infant (before filling it in) and attach extra sheet(s) or download additional forms from npeu.ox.ac.uk/ukoss	
6b.1	Date and time of delivery:	Y Y h h m m
6b.2	Mode of delivery: Spontaneous vaginal Ventouse or Forceps	Vaginal Breech
	Pre-labour caesarean section Caesarean section after	•
6b.3		
	-	
6b.4		
6b.5	Was the infant stillborn?	Yes No
	If Yes, please go to section 7	

6b.6 5 min Apgar				
6b.7 Was meconium staining of the liquor noted at any point prior to delivery? Yes No				
6b.8 Was the umbilical arterial or venous pH measured?YesNo				
If Yes, what was the umbilical arterial pH? or tick if not measured or tick if not measured or tick if not measured 				
6b.9 Was the infant admitted to the neonatal unit? Yes No				
If Yes, what was the indication?				
6b.10 Did any major infant complications occur? ^{7*} Yes No				
If Yes, please specify				
6b.11 Did this infant have a congenital abnormality: Yes No				
If Yes, please give details				
6b.12 Did this infant die? Yes No				
If Yes, please specify date of death	Y			
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				

DD/MM/Y

Υ

Section 8:

8.1	Name of person	completing the form:
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- 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. 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