

Severe Obesity

Study 01/16

Sample Data Collection Form - CONTROL

Instructions for identifying controls:

- 1. Identify the date and time of admission for labour care for the CASE you have reported.
- 2. From the unit admission book/electronic records identify the two women admitted for labour care in the unit immediately BEFORE the CASE. These women will act as CONTROLS.
- 3. Record identifying details for controls alongside the relevant case in your paper case log.
- 4. Retrieve the hospital case notes for the controls and enter data on controls via the website.

This is a Sample Data Collection Form for information only. **Please do not use this form to provide us with information on your controls.** Data should be entered using our OpenClinica system at https://openclinica.npeu.ox.ac.uk/OpenClinica

Section 1 Woman's details

Black African

Section 1. Woman's actums		
1.1 Body Mass Index (BMI) at time of booking		
Should be ≤35 kg/m² for controls.		
kg/m²		
1.2 Age at delivery (years)		
years		
1.3 Ethnic group		
Please tick one		
White British		
White Irish		
Any other White background		
Mixed White & Black Caribbean		
Mixed White & Black African		
Mixed White & Asian		
Any other mixed background		
Indian		
Pakistani		
Bangladeshi		
Any other Asian background		
Black Caribbean		

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Any other Black Chinese Any other ethnic Not recorded			
Yes, please give	man in currently in paid en woman's occupation		_
0.000 [Note: This is de	In Low-income Families Merived from the woman's postcode tion form and enter the woman's postcoto the form]	You will r	
1.6 Height at	booking (cm) cm [Or not recorded]	~ <	0,
1.7 Minimum	recorded weight (kg) & d		
1.8 Maximum	recorded weight (kg) & d		
1.9 Smoking s Never smoked Gave up prior to Gave up during p Current smoker Not recorded			
Section 2. Pres	gnancy/antenatal histor	y	
2.1 Has this w Yes [If Yes, go to No [If no, go to 2		regnancies?	
	umber of completed preg regnancy	nancies ≥24 weeks, p	rior to this
2.1.2 N	umber of pregnancy losse	es<24 weeks	

2.1.3 Was this woman known to have had complications in a previous pregnancy?

	pregnancy:
	For example, unexplained stillbirth/neonatal death; pre-eclampsia requiring preterm
	birth; primary PPH requiring treatment or transfusion; retained placenta requiring
	manual removal; caesarean section; shoulder dystocia.
	Yes, please specify
	No
=	Was this pregnancy conceived through assisted conception? Yes No
2.3	Final Estimated Date of Delivery (EDD)
Note:	: Use the best estimate (ultrasound or date of last menstrual period) based on a 40 week gestation (Date)
2.4	Immediately prior to the onset of labour was this woman known to have any medical conditions?
For e	xample: confirmed cardiac disease; essential hypertension; asthma (please specify if requiring increase
in tre	atment or hospital treatment); thromboembolic disorders; atypical antibodies; Group B Streptococcus
hype	rthyroidism; epilepsy.
\sqsubseteq	Yes, please specify
	No
2.5	During antenatal care were any current pregnancy problems identified?
For e	xample: pre-eclampsia or pregnancy induced hypertension; small for gestational age.
Ш	Yes, please specify
	No
2.6	Did this woman have an oral glucose tolerance test during pregnancy?
	Yes [If Yes, go to 2.6.1]
	No [If no, go to Section 3]
	2.6.1 Did this test indicate gestational diabetes? Yes No

Section 3. Labour and birth care

3.1	Date and time of start of labour care in the AMU
	(Date) (Time, 24hr clock)
3.2 up to	Stage of labour at start of labour care Latent stage: painful contractions & some cervical change, including cervical effacement and dilatation of 4 cm;
mate	Active 1st stage: regular painful contractions & progressive cervical dilatation from 4 cm Passive 2nd stage: full dilatation before or in absence of involuntary expulsive contractions Active 2nd stage: baby visible or expulsive contractions with findings/signs of full dilatation or active ernal effort with full dilatation in absence of expulsive contractions
3.3 Pleas	On initial assessment at the start of labour care were any of the following identified? se tick at least one box:
	Maternal tachycardia (Pulse >120 beats/minute on 2 occasions, 30 minutes apart) Hypertension (Single blood pressure reading - diastolic ≥110 mmHg or systolic ≥160 mmHg OR diastolic ≥90 mmHg or systolic ≥140mmHg on 2 readings 30 minutes apart)
	Proteinuria (2+ of protein or more AND single reading of either diastolic blood pressure ≥90 mmHg or systolic ≥140mmHg) Maternal pyrexia (Temperature of ≥38°C on a single reading, or ≥37.5°C on 2 readings 1 hour apart) Vaginal blood loss (Other than a show) Prolonged rupture of membranes (>24 hours before onset of established labour) If Yes, please specify duration Significant meconium (Dark green or black amniotic fluid that is thick or tenacious, or any meconium-stained amniotic fluid containing lumps of meconium) Reported pain differing from pain normally associated with contractions Abnormal presentation, including cord presentation Transverse or oblique lie High or free-floating head (4/5–5/5 palpable or free-floating head in a nulliparous woman) Suspected fetal growth restriction or macrosomia Suspected anhydramnios or polyhydramnios Fetal heart rate abnormality (<100 or >160 beats/minute) Deceleration in fetal heart rate Reduced fetal movements in the last 24 hours None of the above
3.4	Did this woman use immersion in water for pain relief at any time during labour? Yes
	No

3.5 Was this woman transferred to the care of an obstetrician at any time during labour care or immediately after the birth?

Note: Includes transfers for epidural/pain relief. Does NOT include postnatal transfer where baby was transferred to specialist care, but mother's care was not transferred
Yes [If Yes, go to 3.5.1] No [If no, go to 3.6]
3.5.1 Date and time of decision to transfer(Date)(Time, 24hr clock)
3.5.2 Primary reason for transfer Hypertension Significant meconium Confirmed delay in first stage of labour Confirmed delay in second stage of labour Epidural/other pain relief request Fetal heart rate abnormalities in first stage Fetal heart rate abnormalities in second stage Retained placenta Repair of perineal trauma Other, please specify
3.5.3 Was labour augmented with syntocinon? Yes (Date) (Time, 24hr clock) No
3.5.4 Did this woman have an epidural or spinal? Yes (Date) (Time, 24hr clock) No
3.5.5 Did this woman have a general anaesthetic? Yes No
3.6 Was this a multiple birth? Yes, number of babies No
3.6.1 Date and time of delivery (Date) (Time, 24hr clock)
3.7 What was the place of birth? AMU, under midwifery care AMU, under care of obstetrician Obstetric unit, under midwifery care Obstetric unit, under care of obstetrician

3.8	Did this woman give birth in water? Yes
	No
3.9	What was the mode of birth? Spontaneous vertex birth (go to 3.10) Vaginal breech (go to 3.10) Ventouse (go to 3.10) Forceps (go to 3.10) Caesarean section (got to 3.9.1)
	3.9.1 Grade of urgency of Caesarean section Category 1: Immediate threat to life of woman or fetus Category 2: Maternal or fetal compromise, not immediately life-threatening Category 3: Needing early delivery, no maternal or fetal compromise Category 4: At a time to suit the woman and maternity team
	3.9.2 Primary reason for Caesarean section Abnormal presentation Fetal compromise Maternal compromise Slow progress Other, please specify Not known
3.1	O Was shoulder dystocia documented? Yes, please describe management technique used
3.1	Did this woman receive a prophylactic oxytocic (syntocinon) in the 3 rd stage?
	Yes No
Sec	ction 4. Maternal outcomes
4.1	Did this woman have any perineal trauma involving the anal sphincter $(3^{rd}/4^{th})$ degree tear)?
	Yes No
4.2	Did this woman receive postnatal low molecular weight heparin (LMWH) thromboprophylaxis?
	Yes. Please say for how long days or weeks No

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4.3	Within the first 48 hours after giving birth was this woman admitted to a higher level of care?
\vdash	No (go to 4.4)
Ц	Yes, High dependency unit or area (go to 4.3.1)
Ш	Yes, Intensive care unit (go to 4.3.1)
	Yes, other (please specify) (go to 4.3.1)
	4.3.1 What was the main reason for admission to HDU/ICU:
	4.3.2 What was the total duration of stay in HDU/ICU: hours or days
	My athir and a second of the form DDM (AFOO) of the second
4.4	Was this woman recorded as having a PPH of 1500ml or more?
Ц	Yes
	No
4.5	Did this woman receive a blood transfusion within 48 hours of giving
	birth?
	Yes (go to 4.5.1)
	No (go to 4.6)
	4.5.1 When was the first blood transfusion given? Intrapartum End of third stage – 23 hours after birth 24-48 hours after birth
	4.5.2 How many units of whole blood or packed cells did this woman
	receive?
	units
	units
	4.5.3 Was a cell saver used?
	Yes. Volume of patient's blood transfused ml
	No No
	4.5.4 What was the main reason for blood transfusion?
	Uterine atony
	Genital tract trauma
	Retained products/morbidly adherent placenta
	Other, please specify
	Other, please specify
4.6	Was there any other maternal morbidity?
	Yes, please specify
	No ,
_	
4.7	Did this woman die?
	Yes (go to 4.7.1)
	No (go to 4.8)

	4.7.1	Date and time of ma		
		What was the under	rlying cause of maternal de	eath?
4.8		Not yet known as the date of materi ate)	nal discharge?	
		aby outcomes ection if more than one ba	aby	
5.1	What wa	as the birthweight?		(O)
5.2	Sex of bandle Male Female Indeterminate			
5.3	Yes (go to 5.3. No (go to 5.4)			
	E	When did the baby Before the start of care in late After the start of care in late	abour	
5.4	What wa	as the Apgar score a	t 5 minutes?	
5.5	Was the Yes No	baby breastfed at le	east once?	
5.6		thin 48 hours of birtl	neonatal unit or any other h?	paediatric high leve
	5.6.1		tion of stay in the neonatal	unit?

	5.6.2 What was the main reason for admission to the	neonatal unit?
	Hypoxic-ischaemic encephalopathy	
	Hypoglycaemia	
	Birth trauma	
	Feeding problems	
	Other, please specify	
5.7	Was there any other neonatal morbidity?	
	Yes, please specify	
	No	
F 0		
5.8		
닏	Yes (go to 5.8.1)	
Ш	No (go to 5.9)	
	5.8.1 Date and time of neonatal death	
	(Date) (Time, 24hr clock)	
	5.8.2 Primary cause of neonatal death	
	Congenital anomaly	
	Antepartum infection	
	Immaturity related conditions	
	Intrapartum asphyxia, anoxia or trauma	
	Infection	
	Other, please specify	
	Not yet known	
5.9	Date of neonatal discharge	
	(Date)	

Section 6. Any other information

6.1 Please enter any other information you feel may be important