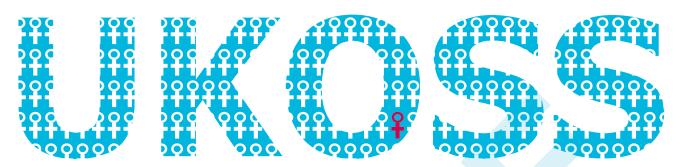
ID Number:



UK Obstetric Surveillance System

Pregnancy outcomes in mothers with a history of travel to a country with active Zika virus (ZIKV) transmission

Congenital Zika virus infection Study 02/16

Data Collection Form - CASE

Please report any woman delivering between 1st Mar 2016 and 28th Feb 2017.

Case Definition:

A history of travel to an area with active Zika virus transmission during pregnancy or in the 4 weeks before conception where a fetal abnormality has been detected, or miscarriage, stillbirth, neonatal death or termination of pregnancy occurred.



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: | | YYYY | |
| 1.2 | Ethnic group:1* (enter code, please se | e back cover for guidance | | |
| 1.3 | Was the woman in paid employment at booking?YesNo | | Yes No | |
| | If Yes, what is her occupation: | | | |
| | | | | |
| | If No, what is her partner's (if any) occu | ipation: | | |
| | | | | |
| 1.4 | Height at booking: | | cm | |
| 1.5 | Weight at booking: | | kg | |
| 1.6 | Smoking status: | never | gave up prior to pregnancy | |
| | | current | gave up during pregnancy | |
| 1.7 | Has this woman previously been rep (<i>i.e.</i> with history of travel to a country w during pregnancy but no adverse pregr | ith active Zika virus transn | • | |

| 1.8 | Did the mother travel to a country with active ZIKV transmission duringpregnancy or within 4 weeks before conception?Yes |
|-----|--|
| | If Yes, please indicate country/countries and subregion (if known) mother travelled to: |
| | 1. Country: Subregion: |
| | Date arrived: DD/MM/YY Date left: DD/MM/YY |
| | 2. Country: Subregion: |
| | Date arrived: DD/MM/YY Date left: DD/MM/YY |
| | 3. Country: Subregion: |
| | Date arrived: DD/MM/Y Date left: DD/MM/Y I |
| | arrival in the country with active ZIKV transmission? (Please tick all that apply) |
| | Fever Rash Itching |
| | Arthralgia/Arthritis Myalgia Conjunctivitis |
| | If Other, please specify: If Other |
| | Date of first symptom onset: |
| | Date last symptom resolved: |
| | |
| 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 the woman have any of the following in previous pregnancies? (Please tick all that apply) |
| | 3 or more miscarriages |
| | Preterm birth or mid trimester loss |
| | Neonatal death |
| | Congenital abnormalities (<i>E.g. autosomal recessive primary microcephaly (MCPH</i>)) |
| | Please specify: |
| | Other genetic disorders |
| | Please specify: |
| | Chromosomal disorders |
| | Please specify: |
| 2.3 | Did the woman have any other previous pregnancy problems? ^{2*} Yes No |
| | If Yes, please specify: |

| Section 3: Previous Medical History | | | | |
|-------------------------------------|--|--|--|--|
| 3.1 | 3.1 Please indicate whether any of the following were present: (<i>Please tick all that apply</i>) | | | |
| | Phenylketonuria Hypothyroidism Known alcohol dependence Known substance misuse | | | |
| | If known substance misuse, please specify drugs/substances used: | | | |
| 3.2 | Did the woman have any other pre-existing medical problem? ^{3*} Yes No | | | |
| | If Yes, please give details: | | | |
| 3.3 | Was the woman prescribed any regular medication in this pregnancy? Yes No | | | |
| | If Yes, please specify drug prescribed, date started and date stopped: | | | |
| | Drug Date Started Date Stopped | | | |
| | | | | |
| | | | | |
| | | | | |
| | (continue in Section 7 if required) | | | |
| | | | | |
| 0 | tion 4. This Dramonat | | | |
| 5 e | ction 4: This Pregnancy | | | |
| 4.1 | Final Estimated Date of Delivery (EDD):4* | | | |
| 4.2 | Was this a multiple pregnancy? Yes No | | | |
| | If Yes, please specify number of fetuses: | | | |
| 4.3 | Did the woman have any of the following in this pregnancy? (Please tick all that apply) | | | |
| | Small for gestational age fetus Hypertension | | | |
| | Gestational Diabetes Mellitus Abdominal injury Rhesus disease | | | |
| 4.4 | Were abnormalities detected on antenatal ultrasound? Yes No | | | |
| | If Yes, was Microcephaly (>2SD below mean for gestational age) detected? Yes No | | | |
| | If Yes, please provide details: | | | |
| | Date detected: | | | |
| | Head Circumference: | | | |
| | Occipitofrontal diameter: | | | |
| | Femur length (FL): | | | |
| | Biparietal diameter: | | | |
| | Abdominal circumference: | | | |
| | Were any of the following detected? | | | |
| | Joint Contractures: Yes No If Yes, Date detected: D / M / Y Craniofacial disproportion: Yes No If Yes, Date detected: D / M / Y | | | |
| | Craniofacial disproportion: Yes No If Yes, Date detected: D / M / Y Intracranial calcifications: Yes No If Yes, Date detected: D / M / Y Y | | | |
| | Any other abnormalities: Yes No If Yes, Date detected: DD/MM/YY | | | |
| | If Yes, please specify other abnormalities: | | | |

| 4.5 | Was an antenatal fetal brain MRI done? Yes No | |
|-----|--|---|
| | If Yes, were abnormalities detected on MRI? Yes No | |
| | If Yes, Were any of the following detected? | |
| | Cortical/subcortical atrophy: Yes No If Yes, Date detected: DD/MM//Y | Υ |
| | Ventricular dilation: Yes No If Yes, Date detected: DD/MM/ | Υ |
| | Lissencephaly: Yes No If Yes, Date detected: DD/MM// | Υ |
| | Pachygyria: Yes No If Yes, Date detected: DD/MM/ | Υ |
| | Any other abnormalities: Yes No If Yes, Date detected: DD/MM/ | Υ |
| | If Yes, please specify other abnormalities: | |
| 4.6 | Were maternal samples collected for ZIKV testing? Yes No | |
| | If Yes, please indicate which samples and the date and result below: | |
| | EDTA blood: Yes No Date: DD/MM/YY Pos Neg | |
| | Urine: Yes No Date: DD/MM/YY Pos Neg | |
| | Amniotic fluid for Zika virus: Yes No Date: DD/MM/YY Pos Neg | |
| | Clotted blood for local save serum: Yes No Date: DD/MM// | Υ |
| 4.7 | Were any other screening tests performed? Yes No | |
| | If Yes, please indicate tests, date and result below: | |
| | Toxoplasmosis: Yes No Date: DD/MM/ Pos Neg | |
| | Rubella: Yes No Date: D Model Neg | H |
| | Cytomegalovirus: Yes No Date: DD/MM/M Pos Neg | |
| | Herpes simplex: Yes No Date: D M Pos Neg | |
| | Syphilis: Yes No Date: D Mo Neg | |
| | HIV: Yes No Date: D / Yes Neg | |
| | Karyotype/array CGH: Yes No Date: Image: Constraint of the second seco | |
| 4.8 | Were there any other problems in this pregnancy? ^{2*} Yes No | |
| | | |
| | If Yes, please specify: | |
| | | |
| | | |
| | | |
| Se | ction 5: Delivery | |
| 5.1 | Did this woman have a miscarriage? Yes No | |
| | If Yes, please specify date: | Υ |
| | and cause (if known): | |
| 5.2 | Did this woman have a termination of pregnancy (including hysterotomy)? Yes No | |
| | | |
| | If Yes, please specify date: | |
| | | _ |
| | If Yes to 5.1 or 5.2, please go to sections 6a, 7 and 8 | |

| | Is this woman still undelivered? If Yes, will she be receiving the rest of her antenatal ca If No, please indicate name of hospital providing fut | | Yes No |
|----------------------------|---|---|------------------|
| | Will she be delivered at your hospital? If No, please indicate name of delivery hospital, the | n go to Section 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: ^{5*} | | |
| | Indication for caesarean section: | | |
| | Method of anaesthesia: | Regional Gen | eral anaesthetic |
| Section 6: Outcomes | | | |
| Sec | ction 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: | | |
| | OD Tiels if we are an even a factor of a second to an other a large still be | | |
| 62.2 | OR Tick if woman was transferred to another hospital: | | |
| 6a.2 | Did any other major maternal morbidity occur?6* | | Yes No |
| | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: | | |
| 6a.2 6a.3 | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? | | Yes No Yes No |
| | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: | death certificate? | |
| | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death | death certificate? | |
| 6a.3 | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the | death certificate? | |
| 6a.3 | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the (Please state if not known) | se photocopy the infant s | Yes No |
| 6a.3 | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the (Please state if not known) ction 6b: Infant 1 If more than one infant, for each additional infant, pleas (before filling it in) and attach extra sheet(s) or downl | se photocopy the infant s | Yes No |
| 6a.3 Sec NB: | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the (<i>Please state if not known</i>) ction 6b: Infant 1 If more than one infant, for each additional infant, pleas (before filling it in) and attach extra sheet(s) or downl www.npeu.ox.ac.uk/ukoss Date and time of delivery: | se photocopy the infant s | Yes No |
| 6a.3 Sec NB: 6b.1 | Did any other major maternal morbidity occur? ^{6*} If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the (<i>Please state if not known</i>) tion 6b: Infant 1 If more than one infant, for each additional infant, pleas (before filling it in) and attach extra sheet(s) or downly www.npeu.ox.ac.uk/ukoss Date and time of delivery: | se photocopy the infant s load additional forms from | Yes No |
| 6a.3 Sec NB: 6b.1 | Did any other major maternal morbidity occur?6* If Yes, please specify: Did the woman die? If Yes, please specify date and time of death What was the primary cause of death as stated on the (Please state if not known) ction 6b: Infant 1 If more than one infant, for each additional infant, pleas (before filling it in) and attach extra sheet(s) or downl www.npeu.ox.ac.uk/ukoss Date and time of delivery: Mode of delivery: Spontaneous vaginal Pre-labour caesarean section | se photocopy the infant s load additional forms from | Yes No |

| 6b.5 Was the infant stillborn? | | Yes | No |
|--|----------------------------|----------|-------|
| If Yes, when did the fetus die? | Antepartum OR | Intrapar | tum 🗌 |
| Cause of still birth (if known): | | | |
| Please go to section 7 | | | |
| 6b.6 5 min Apgar | | | |
| 6b.7 Was the infant admitted to the neonatal unit? | | Yes | No |
| 6b.8 Was Microcephaly (>2SD below mean for gestational a confirmed? | ge and sex) | Yes 🗌 | No 🗌 |
| 6b.9 Did the infant have any other confirmed anomalies? | | Yes | No |
| If Yes, please specify: | | | |
| 6b.10 Did any other major infant complications occur? ^{7*} | | Yes | No |
| If Yes, please specify: | | | |
| 6b.11 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 deat | DD/MM/Y th certificate? | Yhh | 24hr |
| (Please state if not known) | | | |
| Was microcephaly confirmed at postmortem? | | Yes | No 🗌 |
| Were there any other abnormalities confirmed postmortem? | ? | Yes | No 🗌 |
| If Yes, please specify: | | | |

Section 7:

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

| Sec | tion 8: | |
|-------|---|--------------------------|
| 8.1 | Name of person completing the form: | |
| 8.2 | Designation: | |
| 8.3 | Today's date: | |
| You r | nay find it useful in the case of queries to ke | eep 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