



Public Health
England

National Congenital Anomaly and Rare Disease Registration Service (NCARDRS)

Data collection form – Antenatal

Please notify any suspected or confirmed anomaly identified antenatally – structural, chromosomal or biochemical.
DO NOT WAIT until final confirmation before sending this form.

Authorised under Section 251 of the NHS Act 2006 to collect information without patient consent (CAG 10-02(d)2015)

MOTHER'S DETAILS

(Sticky label, if available)

Surname:

Forename:

Hosp. no:

NHS no:

Address at booking:

Postcode: Date of birth:

Ethnic category: White Mixed Indian Pakistani
 Bangladeshi Other Asian* Black Caribbean Black African
 Other Black* Chinese Other* Not known

*If other, please state:

Occupation:

BOOKING DETAILS

Date of 1st booking appointment:

Booking hospital:

EDD:

Height: cm Weight: kg BMI:

Smoking status: Current Ex Non Never Not known

Weekly alcohol units at booking:

Substance use at booking: Yes No Not known

If yes, substance:

Prescription drugs (1st trimester) inc. dose:

Maternal illnesses:

Folic acid: Pre and post conception Post conception only
 Taken, timing unknown Not taken Not known

If taken, dose: Standard 400mcg High 5mg

Assisted conception: Yes No Not known

If yes, type: Ovulation induction IVF ICSI Not known

Number of previous live births:

Number of previous stillbirths (24+ weeks, incl. TOPs):

Number of previous losses (<24 weeks, incl. TOPs):

Number of previous neonatal deaths:

Previous congenital anomalies:

Father's age at booking: years

Family history of anomalies:

Maternal:

Paternal:

Consanguinity: No Yes, 1st cousin Yes, 2nd cousin
 Yes, other Yes, relation nk Not known

PREGNANCY DETAILS

Number of fetuses:

Twin type/chorionicity:

Please attach copies of any relevant scans/clinic letters/laboratory or post mortem reports.

Please send by secure electronic transfer to your regional NCARDRS office. Details of each regional NCARDRS office can be found at www.gov.uk/phe/ncardrs.

Click to lock all form fields and prevent future editing

ANEUPLOIDY SCREENING DETAILS

Date (specimen)	Test	Result
<input type="text"/>	<input type="radio"/> Combined	<input type="radio"/> Accepted
	<input type="radio"/> Quad	T21 risk:1 in <input type="text"/> T13/18 risk: 1 in <input type="text"/>
	<input type="radio"/> NIPT	<input type="radio"/> Declined <input type="radio"/> Not offered Reason <input type="text"/>
		<input type="radio"/> Positive <input type="radio"/> Negative <input type="radio"/> Inconclusive
		Risk: 1 in <input type="text"/>

DIAGNOSTIC TEST DETAILS

Date (procedure)	Sample	Result
<input type="text"/>	<input type="radio"/> CVS	<input type="radio"/> Normal <input type="radio"/> Abnormal <input type="radio"/> Declined
	<input type="radio"/> Amnio	<input type="radio"/> Offered <input type="radio"/> Not offered Reason <input type="text"/>
	<input type="radio"/> Fetal blood	<input type="radio"/> Other, specify: <input type="text"/>
Karyotype/microarray: <input type="text"/>		

ANTENATAL SCAN DETAILS

1st trimester (dating) scan:

Date: USS findings (*attach report*): Normal Abnormal Incomplete

NT measurement: mm

Fetal anomaly (18⁺⁰ – 20⁺⁶) scan:

1st attempt

Date: USS findings (*attach report*): Normal Abnormal Incomplete Not known

Not done, give details:

2nd attempt

Date: USS findings (*attach report*): Normal Abnormal Incomplete

Not done, give details:

Echo/MRI/Other:

Date: Findings (*attach report*):

REFERRAL DETAILS

Department/Hospital:

Consultant:

ADDITIONAL DETAILS

Use this box/back of the form to extend answers or include any extra information you think is relevant

NOTIFIER DETAILS

Name:

Hospital:

Department:

Email:

Tel: Date: