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| Multiple Pregnancy and Birth | | | | | | |
|--|--|-----------|-------|---------|--------------|---------|
| Classification: | Guidelir | ne | | | | |
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| Authors Division: | Women | and Child | 's He | alth | | |
| Departments/Group this Document applies to: | Midwives, Obstetricians and Neonatal Team | | | | | |
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| Guideline to be followed by (target staff): Reference and guidance in the clinical management of multiple pregnancies for all women during the antenatal and intrapartum period. | | | | | | |

To be read in conjunction with the following documents:

MKUH Caesarean Section (MIDG/GL/36)

MKUH Fetal Monitoring (MIDG/GL/48)

MKUH Management of preterm rupture of membranes and preterm labour guideline

MKUH Screening guideline

Are there any eCARE implications? No

CQC Fundamental standards:

Regulation 9 – person centered care

Regulation 10 - dignity and respect

Regulation 11 – Need for consent

Regulation 12 – Safe care and treatment

Regulation 13 – Safeguarding service users from abuse and improper treatment

Regulation 14 – Meeting nutritional and hydration needs

Regulation 15 – Premises and equipment

Regulation 16 – Receiving and acting on complaints

Regulation 17 - Good governance

Regulation 18 - Staffing

Regulation 19 – Fit and proper

Disclaimer

Since every patient's history is different, and even the most exhaustive sources of information cannot cover every possible eventuality, you should be aware that all information is provided in this document on the basis that the healthcare

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professionals responsible for patient care will retain full and sole responsibility for decisions relating to patient care; the document is intended to supplement, not substitute for, the expertise and judgment of physicians, pharmacists or other healthcare professionals and should not be taken as an indication of suitability of a particular treatment for a particular individual.

The ultimate responsibility for the use of the guideline, dosage of drugs and correct following of instructions as well as the interpretation of the published material **lies solely with you** as the medical practitioner.

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Guideline Statement

To provide care which minimises the risk of complications and promotes the health and wellbeing of the woman and her babies in multiple childbearing.

To ensure that the correct procedures and effective communication has been adopted by staff who are dealing with multiple births.

Executive Summary

- Multiple pregnancies should have Consultant-led care.
- All multiple pregnancies should be started on aspirin from 12 weeks if they have one or more risk factors for hypertension.
- Upon identification at the dating scan, mono chorionic twin pregnancies should be referred
 to the Fetal Medicine Consultant where an individualised care plan can be agreed to include
 more frequent ultrasound monitoring for fetal well-being and earlier delivery than dichorionic
 twins.
- Multiple pregnancies with severe twin to twin transfusion should be referred to a Regional Fetal Medicine centre.
- Multiple pregnancies are associated with an increased incidence of obstetric complications therefore experienced midwives and Obstetric Consultant should be involved in labour management plans.

1.0 Roles and Responsibilities:

Doctors – decision making, discussion, planning and delivering care.

Midwives – decision making, antepartum, intrapartum and post-delivery care.

The role of Midwives and other healthcare specialists is integral to the management of multiple pregnancies. Additional support to women is available from Twin and Multiple Birth Association (TAMBA) and the Multiple Births Foundation.

2.0 Scope of document

Reference and guidance in the clinical management of multiple pregnancies for all women during the antenatal and intrapartum period.

3.0 Implementation and dissemination of document

This guideline is available on the Trust intranet and has followed the full guideline review process prior to publication.

4.0 Processes and procedures



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4.1 General Care (see also appendix 1)

- Multiple pregnancies should have a named consultant; they should have shared care with community midwives but with increased input from a consultant. Higher multiple pregnancies should have the same shared care but will have further individualised care plans.
- Women/birthing people with multiple pregnancies may require referral to a women's health physiotherapist, a perinatal mental health professional, an infant feeding specialist or a dietitian for further support in their pregnancy.
- Women/birthing people with multiple pregnancies require the same advice on diet, lifestyle and nutritional supplements as singleton pregnancies.
- There is a higher risk of iron deficiency anaemia and therefore an additional full blood count should be sent at 20-24 weeks as well as the routine at 28 weeks. Twin pregnancies should have their ferritin levels monitored and be kept above 30.
- Advise the woman/birthing person that multiple pregnancy is a risk factor for pre-eclampsia.
 If the Woman/birthing person has 1 additional risk factor or more, as listed below, they should commence 75 -150mg of aspirin daily from 12 weeks until the birth.
- Risk factors for pre-eclampsia:
 - first pregnancy
 - o age 40 years or older
 - o pregnancy interval of more than 10 years o BMI of 35 kg/m2 or more at first visit
 - o family history of pre-eclampsia
- Community midwives should refer to the GPs to prescribe aspirin to women with multiple pregnancies and the above risk factors, to commence between 12 -16 weeks.

Please refer to the Hypertensive Disorders in Pregnancy guideline:

Hypertensive Disorders of Pregnancy(including Pre-eclampsia and Eclampsia) .pdf (adobe.com)

4.2 Dating (First trimester) scan

- Offer women/birthing people with a multiple pregnancy a first trimester ultrasound scan to estimate the gestational age and determine chorionicity and amnionicity.
- Estimate gestational age should calculated from the largest baby
- Determine chorionicity and amnionicity at the time of detecting a twin or triplet pregnancy by ultrasound using:
 - the number of placental masses
 - o the presence of amniotic membrane(s) and membrane thickness
 - o the lambda or T-sign.
- Nomenclature for each twin assigned and documented clearly to ensure consistency throughout pregnancy. Twin in maternal right side is called twin 1.
- If woman/birthing person presents after 14+0 weeks, determine chorionicity and amnionicity at the earliest opportunity by ultrasound using all of the following:
 - o the number of placental masses
 - o the presence of amniotic membrane(s) and membrane thickness
 - o the lambda or T-sign
 - discordant fetal sex.



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- If it is not possible to determine chorionicity or amnionicity, refer the woman to Fetal Medicine consultant. If it is still not possible, plan the care for the pregnancy as a monochorionic pregnancy until proved otherwise.
- Nuchal translucency: This should be offered as the preferred method of chromosomal abnormality screening however this cannot be offered for higher order pregnancies.
- Upon identification of a multiple pregnancy please inform Antenatal Newborn (ANNB)
 Screening to attend the department to facilitate counselling to women before and after
 every screening test. A further discussion regarding informed consent for Trisomy
 screening in twin pregnancies is necessary prior to obtaining the blood sample. It is
 important the woman/birthing person understands the implications of receiving a separate
 result for each fetus.

Please refer to the Fetal Anomalies guideline: Fetal Anomalies Guideline .pdf (adobe.com)

4.3 Antenatal Care

- Offer screening for structural abnormalities in twin and higher order pregnancies as in routine antenatal care. This might be completed in the main ultrasound department.
 Consider a slightly later gestational age than in singleton pregnancies.
- Do not offer women with a twin or triplet pregnancy screening for fetal growth restriction or feto-fetal transfusion syndrome in the first trimester.
- Inform women/birthing person that 60% twin pregnancies will result in spontaneous delivery before 37 weeks. This increases to 75% in triplet pregnancies delivering before 35 weeks.
- Increased risk of admission into neonatal unit.

Dichorionic Diamniotic Twin Pregnancy

- Offer women/birthing people with an uncomplicated dichorionic diamniotic twin pregnancy antenatal clinic appointment with an Obstetrician along with a scan at 24, 28, 32 and 36 weeks.
- Women/birthing people will continue to have routine antenatal appointments with their community midwife.
- At each ultrasound scan from 24 weeks, offer women with a dichorionic twin pregnancy diagnostic monitoring for fetal weight discordance using 2 or more biometric parameters and amniotic fluid levels. To assess amniotic fluid levels, measure the deepest vertical pocket (DVP) on either side of the amniotic membrane.

Monochorionic Twin Pregnancy

• Offer women/birthing people with an uncomplicated monochorionic twin pregnancy antenatal clinic appointment with an Obstetrician along with a scan at 16, 18, 20, 22, 24, 28, 32 and 34. Some of these scans, especially the 16 weeks scan should be undertaken in the Fetal Medicine Clinic.



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- Women/birthing people will continue to have routine antenatal appointments with their community midwife.
- Refer women/birthing people for a fetal cardiac ultrasound scan in the fetal medicine tertiary centre at 20-21 weeks gestation.
- Offer women/ birthing people simultaneous monitoring for feto-fetal transfusion syndrome, fetal growth restriction and advanced-stage twin anaemia polycythaemia sequence (TAPS) at every ultrasound assessment to monitor effectively for all complications of monochorionicity. Explain that the relative likelihood of each complication changes with advancing gestation but that they can all occur at any gestational age.

Higher Order Pregnancy

• Higher order multiple pregnancies are rare and for these individualised management plans should be developed by the Fetal Medicine Consultant. A referral should be made to a tertiary centre if appropriate.

4.4 Fetal Complications

Twin to Twin Transfusion Syndrome (TTTS)

- Women/birthing people with monochorionic twins should be asked to report a sudden increase in abdominal size or breathlessness as these may be manifestations of TTTS.
- Ultrasound evidence of TTTS includes:
 - o difference between deep vertical pool (DVP) depth of 4 cm or more
 - o One baby has a DVP depth of less than 2 cm, **and** another baby has a DVP depth of:
 - > over 8 cm before 20+0 weeks of pregnancy or
 - over 10 cm from 20+0 weeks
 - One baby has normal DVP depth, and another baby has a DVP depth of:
 - Less than 2 cm or
 - More than 8 cm
- If twin to twin transfusion is present at any ultrasound scan, referral should be made to a Fetal Medicine Consultant Obstetrician.
- In the presence of moderate to severe twin to twin transfusion referral should be made to a
 Regional Fetal Medicine Centre. Ongoing care may be managed by the regional centre or
 may be returned to the local Fetal Medicine Consultant Obstetrician for ongoing
 management. These cases may be discussed at the monthly fetal medicine forum.
- Baby alert forms will be completed.

Selective Fetal Growth Restriction

- Do not use abdominal palpation or symphysis—fundal height measurements.
- Use ultrasound scan from 24 weeks using two or more biometric parameters to estimate fetal weight (EFW).



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- Calculate and document EFW discordance in mono chorionic and dichorionic twins using the formula below: (EFW larger fetus – EFW smaller fetus) ÷ EFW larger fetus
- Increase ultrasound frequency to at least weekly, and include doppler assessment of the umbilical artery flow for each baby, if:
 - o there is an EFW discordance of 20% or more and/or
 - o the EFW of any of the babies is below the 10th centile for gestational age
- Refer woman to fetal medicine clinic if there is an EFW discordance of 25% or more and
 the EFW of any of the babies is below the 10th centile for gestational age because this is a
 clinically important indicator of selective fetal growth restriction.

Preterm Labour

- Women with multiple pregnancy are at higher risk of spontaneous preterm birth than women with a singleton pregnancy (up to 60% of twin pregnancies).
- This risk is further increased if they have other risk factors, such as a spontaneous preterm birth in a previous pregnancy.
- Do not use fetal fibronectin or cervical length alone to diagnose preterm labour.
- Do not offer the following interventions (alone or in combination) routinely to prevent spontaneous preterm birth in women with a multiple pregnancy:
 - Arabin pessary
 - o cervical cerclage
 - oral tocolytics
 - Progesterone intramuscular therapy

See Appendix 2: Indications for referral to a tertiary level fetal medicine centre.

4.5 Planning Birth

Discussion

Discuss the following with the woman/birthing person from 24 weeks by 28 weeks:

- place of birth and the possible need to transfer in case of preterm birth timing and possible modes of birth
- analgesia during labour (or for caesarean birth) intrapartum fetal heart monitoring
- o management of the third stage of labour.
- Increase risk of admission to the neonatal unit.

Timing of birth

Uncomplicated DCDA pregnancy:

- Explain to women/birthing person that elective birth from 37 weeks 0 days does not appear
 to be associated with an increased risk of serious adverse outcomes, and that continuing
 uncomplicated twin pregnancies beyond 38 weeks 0 days increases the risk of fetal death.
- Offer delivery after 37 weeks 0 days if uncomplicated pregnancy.

Uncomplicated MCDA pregnancy:

 Explain to women/birthing person that elective birth from 36 weeks 0 days does not appear to be associated with an increased risk of serious adverse outcomes, and that continuing



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uncomplicated twin pregnancies beyond 38 weeks 0 days increases the risk of fetal death.

Offer delivery after 36 weeks 0 days following consideration for corticosteroids.

Uncomplicated MCMA pregnancy:

- Explain to women/birthing person that planned birth between 32+0 and 33+6 weeks with corticosteroids does not appear to be associated with an increased risk of serious neonatal adverse outcomes, and that continuing the pregnancy beyond 33+6 weeks increases the risk of fetal death.
- Babies will usually need to be admitted to the neonatal unit and have an increased risk of respiratory problems.
- Higher order pregnancy or in complicated twin pregnancy the timing of birth will be decided and discussed with each woman/birthing person individually by Fetal Medicine Consultant Obstetrician.

Please refer to: Management of Preterm Labour Guideline .pdf (adobe.com)

For women/birthing person who decline planned birth at the expected timing as above, offer
weekly appointments with the specialist obstetrician. At each appointment, offer an
ultrasound scan and perform assessments of amniotic fluid level and doppler of the umbilical
artery flow for each baby in addition to fortnightly fetal growth scans.

Mode of Birth

- DCDA and MCDA pregnancy:
 - Explain to women/birthing person with uncomplicated pregnancy that planned vaginal birth and planned caesarean section are both safe choices for them and their babies if all of the following apply:
 - · the pregnancy remains uncomplicated and has progressed beyond 32 weeks
 - there are no obstetric contraindications to labour
 - the first baby is in a cephalic (head-first) presentation
 - there is no significant size discordance between the twins.
 - Planned Induction of labour could be started on Ward 9.
 - Discuss caesarean section to women/birthing person if the first twin is not cephalic at the time of planned birth.
 - Discuss an individualised assessment of mode of birth to women/birthing people in suspected, diagnosed or established preterm labour before 26 weeks. Consider the risks of caesarean section and discuss these with the Woman/birthing person.
 - Explain to women with an uncomplicated twin pregnancy that for women giving birth after 32
 weeks more than a third of women who plan a vaginal birth go on to have a caesarean
 section.
 - Explain to women who plan a caesarean section that a few will have a vaginal birth before the caesarean section can be carried out.
 - Explain to women/birthing person who plan for vaginal birth that a small number of women will need an emergency caesarean section to deliver the second twin after vaginal birth of the first twin.

MCMA pregnancy:

- Delivery by caesarean section should be discussed. Delivery should be planned for between 32+0 and 33+6 weeks as there is a higher risk of cord entanglement.
- Higher order pregnancy or in complicated twin pregnancy:



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 Delivery by caesarean section should be discussed. A management plan will have been formalised with the woman/birthing person antenatally.

4.6 Care in Labour Analgesia

- Discuss options for analgesia and anaesthesia with the women/birthing person antenatally, prior to 28 weeks, whether they are planning a vaginal birth or caesarean section.
- Discuss an epidural with women/birthing people who choose to have a vaginal birth. Explain that this is likely to:
- enable a quicker birth by emergency caesarean section if needed.
- Discuss regional anaesthesia to women with a twin or triplet pregnancy who are having a caesarean section.

Intrapartum Care

Once admitted in labour

- When a woman/birthing person with a multiple pregnancy is in labour the obstetric consultant on call and Specialist Registrar will be informed. The most senior obstetric team member will review the plan of care and birth preferences with the Woman/birthing person
- Labour care will be provided by a midwife
- The obstetric Anaesthetist and Neonatal team should be informed the Woman/birthing person is in labour
- If there is no clear plan of labour and documentation of the discussed mode of delivery in the maternal records, then the Obstetric Registrar will discuss the Womans/birth persons wishes and discuss the plan with the Consultant on-call, this will then be document in the maternal records.
- A large bore intravenous cannula (grey 16g venflon) should be inserted. Take full blood count and group & save serum.
- Commence Omeprazole orally 12 hourly
- Prepare the room for delivery and prepare resuscitation equipment for the babies
- For women between 23+0 and 25+6 weeks of pregnancy who are in established labour, involve a senior obstetrician in discussions with the woman/birthing person about how to monitor the fetal heart rates
- Perform a portable ultrasound scan when established labour starts, to confirm which twin is which, the presentation of each twin, and to locate the fetal hearts
- Continuous electronic fetal monitoring is indicated for all women/birthing people above 26
 weeks gestation, with a twin pregnancy in labour. Please refer to Fetal Monitoring
 Guideline.

Fetal Monitoring Guideline .pdf (adobe.com)

- Consider FSE monitoring for the presenting twin if abdominal monitoring difficult or a continuous CTG is not being obtained.
- If the registrar or lead midwife has any concerns about CTG monitoring including concerns that only one twin is being monitored, this must be escalated immediately to the Consultant on-Call.

On diagnosis of 2nd stage of Labour

- Inform Labour Ward shift lead.
- Inform the obstetric consultant on call and specialist registrar. The obstetric consultant on call





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should be present on labour ward during the 2nd stage of labour.

- The obstetric registrar, obstetric SHO, neonatologist, anaesthetist and ODP must be present on labour ward when presenting twin is imminent and stay until the 3rd stage of labour.
- The woman/birthing person and partner must be fully informed at all times.
- Consider birth in theatre if difficulties with birth are anticipated.
- The ultrasound scanner should be available outside the birthing room.
- Prepare Oxytocin 10 units in 500ml 0.9% Saline for possible use to augment contractions after first baby is delivered if required.

Following birth of first baby

- One cord clamp is applied to mark the first cord
- Palpate abdomen to assess lie of second baby.
- If the lie of second baby is not longitudinal then stabilise the lie to longitudinal lie, if necessary, an external version could be performed by the obstetric registrar or consultant. Ultrasound should be used to confirm the lie and presentation.
- Continue to monitor the second baby using cardiotocography.
- If no contractions are apparent within 5-10 minutes, the woman/birthing person should be reviewed by the obstetric registrar or consultant and to discuss commencing oxytocin infusion, 10 units in 500mls as per augmentation regime.
- Artificial rupture of membranes should NOT be performed until the presenting part is engaged to reduce the risk of cord prolapse. If it is not, to stabilise to a longitudinal lie by performing external version.
- External version to achieve longitudinal lie should be performed if needed. If unsuccessful, internal podalic version or Caesarean section needs to be considered depending on clinical judgement
- For second twin second stage of labour should be managed as per the intrapartum care guideline

Intrapartum Care.pdf (adobe.com)

- If delivery of the second twin is not anticipated within 30mins of the first, the Consultant on Call is to be informed.
- The cardiotocograph should be interpreted based on the clinical picture and the delivery managed accordingly.
- Two cord clamps are applied to mark the cord of the second twin.

The 3rd stage of labour

- By 28 weeks of pregnancy, discuss with women with a twin or triplet pregnancy the potential need for blood transfusion, including the need for intravenous access
- After the birth of both babies, consider double clamping the cord to allow umbilical cord blood gases to be sampled. Ensure that the samples are correctly labelled for each baby
- Postpartum haemorrhage is more common after multiple delivery therefore active management of the third stage is advised.
- IM syntometrine (or IV Oxytocin if the woman is hypertensive) is administered following birth of the 2nd twin.
- 40 units of Oxytocin in 500mls of Normal Saline should be commenced at a rate of 125mls/hour (10iu/hr) after delivery of the placenta.



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5.0 Statement of evidence/references

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6.0 Governance

6.1 Document review history

| Version number | Review date | Reviewed by | Changes made |
|----------------|----------------|---------------------------------------|---|
| 5 | January 2018 | Ghaly Hanna | Reviewed and updated. |
| 5.1 | October 2018 | Sharon Page | Page 5 continuation of 4.1 add a bullet point after Nuchal Translucency: This should be offered as the preferred method |
| 6 | January 2021 | Ghaly Hanna /Emad Nasr/Katie Selby | Complete review and GAP |
| 6.1 | September 2023 | Lila Ravel | Complete review and GAP with SBL 3 |

6.2 Consultation History

| Stakeholders Name/Board | Area of Expertise | Date Sent | Date Received | Comments | Endorsed Yes/No |
|----------------------------|-----------------------|------------|------------------|-------------------|-----------------|
| Janice Styles | Consultant Midwife | 25/01/2022 | | Change to wording | yes |
| Anita Males | Screening Lead | 25/01/2022 | | Read guideline | |
| Maternity CIG | Maternity | 25/02/2022 | | | |



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| Maternity guideline review group | Maternity | 25/02/2022 | | |
|----------------------------------|--|------------|---|-----|
| Kathryn O'Gorman | Fetal Medicine and Preterm birth midwife | 02/10/2023 | Add referral forms to appendix – Minor changes to wording | yes |
| Maternity guideline review group | Maternity | 03/10/2023 | | |
| | | | | |

6.3 Audit and monitoring

| Audit/Monitoring Criteria | Tool | Audit Lead | Frequency of Audit | Responsible Committee/Board |
|---|-------|--|--------------------|--------------------------------|
| a. Women with a multiple pregnancy have the chorionicity and amnionicity of their pregnancy determined using ultrasound and recorded between 11 weeks 0 days and 13 weeks 6 days. | Audit | Midwives, doctors, sonographers designated by the audit leads | Quarterly | Audit group meeting |
| b. Evaluation of the accuracy of determining chorionicity and amnionicity | | Doctors and sonographers designated by the audit leads | | |
| c. Women with a multiple pregnancy have their fetuses labelled using ultrasound and recorded between 11 weeks 0 days and 13 weeks 6 days. | | | | |
| d. Women with a multiple pregnancy are cared for by a multidisciplinary core team | | | | |
| e. Women with a multiple pregnancy follow the antenatal care pathway defined in appendix 1 | | | | |
| f. Women with a multiple pregnancy are monitored for fetal complications according to the chorionicity and amnionicity of their pregnancy g. Women with a higher-risk | | | | |



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|-----|---------|---|----------|--|--|
| | | or complicated multiple | | | |
| | | pregnancy have a | | | |
| | | consultant from a tertiary | | | |
| | | level fetal medicine | | | |
| | | centre involved in their | | | |
| | | care. | | | |
| | h. | Women with a multiple | | | |
| | | pregnancy have a | | | |
| | | discussion by 24 weeks | | | |
| | | with one or more | | | |
| | | members of the | | | |
| | | multidisciplinary core | | | |
| | | team about the risks, | | | |
| | | signs and symptoms of | | | |
| | | preterm labour and | | | |
| | | possible outcomes of | | | |
| | | preterm birth | | | |
| | i. | Women with a multiple | | | |
| | | pregnancy have a | | | |
| | | discussion by 32 weeks | | | |
| | | with one or more | | | |
| | | members of the | | | |
| | | multidisciplinary core | | | |
| | | team about the timing of | | | |
| | | birth and possible modes | | | |
| | | of delivery so that a birth | | | |
| | | · | 1 | | |

6.4 Equality Impact Assessment

plan can be agreed.

As part of its development, this Guideline and its impact on equality has been reviewed. The purpose of the assessment is to minimise and if possible remove any disproportionate impact on the grounds of race, gender, disability, age, sexual orientation, religion or belief, pregnancy and maternity, gender reassignment or marriage and civil partnership. No detriment was identified. Equality Impact assessments will show any future actions required to overcome any identified barriers or discriminatory practice.

| Equality Impact Assessment | | | | | | |
|--|--------------------|-----------|---------------------|------------|--|--|
| Division | Women and Children | | Department | Maternity | | |
| Person completing the EqIA | Erica Puri | | Contact No. | Ex 87153 | | |
| Others involved: | yes | | Date of assessment: | 15/03/2022 | | |
| Existing policy/service | yes | | New policy/service | no | | |
| | | | | | | |
| Will patients, carers, the affected by the policy/ | | Yes | | | | |
| If staff, how many/which groups will be effected? | | All Staff | | | | |



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| Protected characteristic | Any impact? | Comments |
|---|------------------|---|
| Age | NO | Positive impact as the policy aims to recognise diversity, |
| Disability | NO | promote inclusion and fair treatment for patients and staff |
| Gender reassignment | NO | |
| Marriage and civil partnership | NO | |
| Pregnancy and maternity | NO | |
| Race | NO | |
| Religion or belief | NO | |
| Sex | NO | |
| Sexual orientation | NO | |
| | | |
| What consultation met carried out? | thod(s) have you | face-to-face teams and meetings |
| How are the changes/a policies/services com | | Email, face to face and teams |
| Review date of EqIA | | 26/02/2025 |



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Checklist for Guideline and guidelines documentation

Fonts should be Arial 14 for headers 12 for main body
Clear Title and replace with document title Font Arial 22

By submitting a document for review/approval you are confirming that the document has been checked against the checklist below to ensure it meets the Trust standards for producing Trust Documentation (for support please contact your Governance Facilitator/Patient Safety Lead.

| | | | - | | | | |
|---|-------------------------|-----------------------------|--------------|--|--|--|--|
| Authors Division: | | | | | | | |
| Department/Groups this | document applies to |): | | | | | |
| Date of approval: | | | | | | | |
| Review date: | | | | | | | |
| Approval Group/approved by (according to policy requirements): | | | | | | | |
| Last review date: | | | | | | | |
| Unique Identifier: if known (new documents will be assigned at publication) | | | | | | | |
| Status: Approved | | | | | | | |
| Version numbers are the | e same throughout d | ocument | | | | | |
| Scope: Who will use this | s document? | | | | | | |
| To be read in conjunction | n with the following of | documents: | | | | | |
| Are there any eCARE in | nplications? | | | | | | |
| | al standards referenc | ed:Trust intranet page with | | | | | |
| <u>fundamental standards</u> | | | | | | | |
| Footers completed to match main page : (on all pages) | | | | | | | |
| References are updated (contact the library (Jayne Plant 3077) for help if | | | | | | | |
| required) | | | | | | | |
| Consultation history incl | - | • | | | | | |
| , | | ocument contains medication | on | | | | |
| Audit and monitoring cri | • | id clear (where possible | | | | | |
| reference the relevant se | | | | | | | |
| Include full & correct consultation history | | | | | | | |
| Dissemination should be clear | | | | | | | |
| Check relevant hyperlinks work | | | | | | | |
| Completed by name: | Position: Division Date | | | | | | |
| | | | (DD/MM/YYY) | | | | |
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Appendix 1: Routine Antenatal Care Outline

| Visit Number | Pregnancy Weeks | What to expect at your appointment | | | | | |
|-----------------|--------------------------|--|-------------------|--|--|--|--|
| 1 | When pregnancy confirmed | You will see your Midwife or GP and a dating scan will be arranged, which you will have around your twelfth week of pregnancy. | | | | | |
| 2 | Booking at 8-12 weeks | Your medical and obstetric history will be taken and recorded. You will be given information on antenatal screening tests and blood may be taken. Information will be given on diet and lifestyle, options on place of birth and any additional care needed. Your will receive your pregnancy record which you should carry with you at all times. Various information leaflets will be provided for you to read at your leisure. | | | | | |
| 3 | 10-13 weeks | Nuchal Translucency & dating sca Consultant appointment arranged | within one or tw | o weeks. | | | |
| 4 | 13-14 weeks | Consultant appointment and discu pregnancy. Discuss screening opt | ions. | | | | |
| | Monoch | orionic Diamniotic Twins | Di | chorionic Twins | | | |
| Hospital | 16 weeks | scan for fetal well-being, antenatal review | 16 weeks- CMW | Routine antenatal review | | | |
| Hospital | 18 weeks | Scan for fetal well-being, antenatal review | | | | | |
| Hospital | 20 weeks | Scan for fetal anomaly cardiac scan including outflow tracts and fetal well-being. Full blood count to be taken by MCA in ANC. | 20 weeks | 20-week scan for fetal anomaly cardiac scan including outflow tracts and fetal well-being. Full blood count. | | | |
| Hospital | 22 weeks | Scan for fetal well-being, antenatal review | | | | | |
| Hospital | 24 weeks | Scan for fetal well-being and review by Consultant. | 24 weeks | Scan for fetal well-being and antenatal review | | | |
| Hospital | 26 weeks | Scan for fetal well-being and review by Consultant. | 28 weeks | 28 weeks scan for fetal wellbeing and review by consultant. | | | |
| Hospital | 28 weeks | Scan for fetal well-being and review by Consultant. | 31 weeks - CMW | Routine antenatal review | | | |
| Hospital | 30 weeks | Scan for fetal well-being and review by Consultant. 32 weeks | | Scan for fetal well-being and review by Consultant. Discussion on mode, place, and timing of delivery | | | |
| Hospital | 32 weeks | Scan for fetal well-being and review by Consultant. Discussion on mode, place, and timing of delivery Scan for fetal well-being and 34 weeks-CMW CMW | | | | | |
| Hospital | 34 weeks | Scan for fetal well-being and review by Consultant. Discussion on mode, place, and timing of delivery t 2 appointments should be with a second secon | 36 weeks | Scan for fetal well-being and review by Consultant. Discussion on mode, place, and timing of delivery | | | |



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Appendix 2: Indications for referral to a tertiary level fetal medicine centre

Seek a consultant opinion from a tertiary level fetal medicine centre for:

Pregnancies with a shared amnion:

- monochorionic monoamniotic twins o dichorionic diamniotic triplets
- monochorionic diamniotic triplets
- monochorionic monoamniotic triplets

Pregnancies complicated by any of the following:

- fetal weight discordance (of 25% or more) and an EFW of any of the babies below the 10th centile for gestational age
- fetal anomaly (structural or chromosomal)
- discordant fetal death
- feto-fetal transfusion syndrome
- twin reverse arterial perfusion sequence (TRAP)
- conjoined twins or triplets
- suspected Twin Anaemia Polycythaemia sequence (TAPS)





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Appendix 3: Referral form to tertiary fetal medicine unit

Oxford University Hospitals WHS

NHS Foundation Trust

OXFORD FETAL MEDICINE UNIT

LEVEL 6, WOMEN'S CENTRE, JOHN RADCLIFFE HOSPITAL, HEADINGTON, OXFORD, OX3 9DU Tel: 01865 221716 (Mon-Fri; 08.30-17.30); Email: fetalmedicine.pnd@nhs.net

| Referring Unit | | | GI | Surgery | | |
|--|--------|-------------------------------------|---|---------------------------|---|--|
| Consultant | | | Date of referral | | / /20 | |
| | | | | | | |
| | | | | CHECKLIST (O | bligate attachments) | |
| PATIENT DETAILS | | | | ☐ Laboratory blood group* | | |
| Name | | | *If Rh Neg and fetal blood group available, please attach | | | |
| | | | | ☐ Latest USS | • | |
| NHS Number | | | | ☐ Booking bl | | |
| DOB | | | | ☐ Combined | /Quadruple screening | |
| | | | | (Diamentish all to | hat and to Barthack l | |
| Address | | | | ☐ Safeguardi | hat apply & attach) | |
| | | | | L Saleguardi | ing concerns | |
| | | | | □ Virology/T | ORCH results | |
| | | | | □ NIPT | ONCH Tesuits | |
| | | | | □ QF-PCR | | |
| TELEPHONE (1) | | | | ☐ Microarray | | |
| | | I | | | □ WES result | |
| TELEPHONE (2) | | | | | s, please send at earliest <u>opportunity</u> | |
| | | | | | , presse serie at contest expenses | |
| PATIENT EMAIL | | | | | | |
| (essential) | | | | | | |
| CLINICAL DETAILS: | oleas | e tick all that <u>apply)</u> D FMU | REFE | RRAL DEET | AL CARDIAC REFERRAL | |
| EDD | \top | | | | | |
| Current Gestation | | ı | Local Anomaly Scan Da (essential) | | ite | |
| CURRENT REFERRAL REASON | Ŀ | L | | (Casericial) | I | |
| | | | | | | |
| • | | | | | | |
| l . | | | | | | |
| • | | | | | | |
| | | | | | | |
| | | | | | | |
| • | | | | | | |
| Additional comments (ag | sor | ial issues/contact specificatio | ne/nra | vious referrals/f: | ailed tests/hospital issues etc) | |
| Additional comments (55 | 300 | iai issues/contact specificatio | nis, pie | rious referralsy in | aneu testa/nospitarissues etc/ | |
| | | | | | | |
| | | | | | | |
| Cardiac Referrals Only: | | | | | | |
| | | evious child or baby's father): | | | | |
| Was the problem present Have they had cardiac s | | | | | | |
| | | | | | | |
| | | | | | | |

Urgent referrals eg TTTS need to be telephoned through to midwife please Routine referrals will not necessarily be actioned same day



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Appendix 4: Referral form to Anaesthetic services

| | Milton Keynes Hospital |
|---------------------------------------|--|
| Milton Keynes Hospital | |
| NOTIFICATION OF EXPE | CTANT MOTHER WITH PROBLEMS POSSIBLY RELEVANT TO ANAESTHESIA |
| Re: | Or: Name: |
| | Address: |
| | |
| Patients contact Tel No: Home Tel: | |
| Mobile: | |
| PATIENT'S | CONSULTANT: |
| | |
| EDD: | |
| The expectant mother is nov | w Weeks pregnant and this is herpregnancy |
| | |
| anaesthesia or regional bloc | m(s) which may become particularly relevant if she needs k: |
| | |
| | |
| | |
| | *************************************** |
| required, more detailed info | ormation is available in her case notes or from me directly |
| | 940 14.0 147.13.0 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 147.1 14 |
| ours sincerely | |
| ours sincerely | |
| 10 300 T 3 C 10 00 C 10 C 10 C | Job Title: Ext |
| 10 300 T 3 C 10 00 C 10 C 10 C | Job Title: Ext |
| Name (in capitals): | AN SECTION – PATIENT CHECKLIST |
| Name (in capitals): | |
| Name (in capitals): | |
| Name (in capitals): | |