

All Wales Antenatal Care Criteria

Women booked by community midwives by 10 + 6 weeks. Additional care needs and recommended antenatal care pathways Where women are under OLC the initial appointment should be with the named consultant, where possible, to aid efficient care planning		
Pathway A	Pathway B	Pathway C
Additional care needs requiring - Obstetric led antenatal care (OLC)	Where some level of additional monitoring is required and/or initial Obstetric antenatal care planning.	Midwifery-led antenatal care (MLC)
<u>Cardiovascular and medical</u> <ul style="list-style-type: none">Confirmed cardiac diseaseSubstance MisuseEpilepsyPrevious epilepsyHypertensionMalignant DiseaseRenal disease.	<u>Cardiovascular and medical</u>	<u>Cardiovascular and medical</u>
<u>Respiratory conditions</u> <ul style="list-style-type: none">Asthma requiring oral steroid treatmentAsthma requiring in-hospital treatment or an increase in treatment during pregnancyCystic Fibrosis	<u>Respiratory conditions</u>	<u>Respiratory conditions</u> <ul style="list-style-type: none">Well controlled Asthma
<u>Haematological</u> <ul style="list-style-type: none">Blood clotting disordersAutoimmune disorders e.g. Systemic Lupus, Antiphospholipid syndromeHaematological – History of sickle-cell, beta thalassemia MajorHistory of thromboembolic disordersHb less than 110 g/L at booking or < 105 g/L at 28/40 with no response to oral iron therapy after 4 weeks.Hb < 90 g/L n isolation.Immune thrombocytopenia purpura or platelet count below <150 µ/L at booking or during the course of pregnancy.Von Willebrand's diseaseBleeding disorder in the women or her unborn babyAtypical antibodiesJehovah's witnessWomen scoring 4 on VTE assessmentAntenatal thromboprophylaxis administration.	<u>Haematological</u> <ul style="list-style-type: none">Women scoring 3 on VTE assessment should be offered an obstetric appointment to discuss thrombo-prophylaxis from 28/40.Women scoring 2 on VTE assessment should have discussion around postnatal thrombo prophylaxis this should be prescribed antenatally where accepted.	<u>Haematological</u> <ul style="list-style-type: none">Hb of <110g/L at booking and <105g/L at 28 weeks require iron therapy and 4 weekly FBC in the community. If not responsive to iron therapy or if Hb<90 g/l at any time or any concerns refer to OLC.
<u>Endocrine</u> <ul style="list-style-type: none">Type 1 and 2 diabeticsGestational diabeticHyperthyroidism (may present as hypo)HypothyroidismWomen on oral steroids	<u>Endocrine</u>	<u>Endocrine</u>
<u>Auto-immune</u> <ul style="list-style-type: none">Systemic lupus erythematosus, SclerodermaConnective tissue disorders	<u>Auto-immune</u>	<u>Auto-immune</u>
<u>Infective</u> <ul style="list-style-type: none">Hepatitis B or CCarriers of, or infected HIVToxoplasmosis currently being treatedActive infection or chicken pox/rubellaPrimary episode of genital herpes or recurrent active lesions after 36/40Tuberculous under treatmentCOVID positive during pregnancy requiring hospital admission.	<u>Infective</u> <ul style="list-style-type: none">Current COVID 19 infectionPrevious baby affected by GBS or diagnosed with GBS this pregnancy.	<u>Infective</u> <ul style="list-style-type: none">Recurrent episodes of genital herpes, women to be treated by GP with Acyclovir from 36/40Group B streptococcus in current pregnancy, who decline IPAB (offer NN observation on obstetric unit as per SRC).GBS in last pregnancy; Offer Vaginal/rectal swab 35-37/40. If negative IPAB not required can be MLC for birth.
<u>Neurological</u> <ul style="list-style-type: none">EpilepsyMyasthenia gravisMultiple sclerosisPrevious cerebrovascular accident	<u>Neurological</u>	<u>Neurological</u>
<u>Gastro-intestinal/Renal</u> <ul style="list-style-type: none">Liver disease (not obstetric-cholestasis)Abnormal renal functions/known renal diseaseCrohn's disease or ulcerative colitis	<u>Gastro-intestinal/Renal</u>	<u>Gastro-intestinal/Renal</u>
<u>Previous pregnancy</u> <ul style="list-style-type: none">Previous Molar pregnancy3 or more consecutive miscarriageMid trimester (12-22 week) lossPrevious HELLP syndromeBaby with neonatal encephalopathyGestational DiabetesPre-eclampsiaPre term birth <34/40Placenta abruptionUterine rupturePervious caesarean birthPrimary PPH 500-999mls requiring additional treatment for uterine atony.Primary PPH ≥1000 mls.Retained placentaShoulder dystociaCervical tears3rd degree tears with ongoing concern or continence issues4th degree tears	<u>Previous pregnancy</u> <ul style="list-style-type: none">Previous SGA below 10th centile at birthPrevious baby >4.5kgPrevious 3rd degree tear with no ongoing concerns around pelvic floor health	<u>Previous pregnancy</u> <ul style="list-style-type: none">PPH 500-999mls not linked to uterine atony or requiring additional treatment for uterine atony or hypovolaemia (confirmed via previous birth records).
<u>Current pregnancy</u> <ul style="list-style-type: none">Screening anomaly including low PAPP-A (<0.415 MOM)Multiple pregnancyGestational diabetesPlacenta praeviaPre-eclampsia /pregnancy induced hypertensionPre-term pre-labour rupture of membranesAPH of placental origin or 2 or more episodes after 24 weeks.Alcohol dependencyMaternal age ≥40 at bookingPrimiparous BMI ≥35Multiparous BMI ≥40Eating disorderGrand multiparty, P5 or >Concerns with fetal growth or placental function.Baby with structural/ Chromosomal abnormalityPolyhydramnios/OligohydramniosEFW via USS ≥97th centile on GROW chart.Gestation >41+6Administering antenatal thromboprophylaxisBreech/malpresentation after 36/40	<u>Current pregnancy</u> <ul style="list-style-type: none">Smoker (Serial USS in line with GAP/GROW)Multiparous women BMI 35-39.9 with a previous vaginal birth in accordance with local criteria for serial USS and GDM screen.Recurrent1 Altered Fetal Movement with normal investigations .	<u>Current pregnancy</u> <ul style="list-style-type: none">Para 4 or <BMI at booking of 30 to 34.9kg/mMaternal age at booking 35-39 inclusive.2 episodes of AFM, which are more than 3 weeks apart, with normal investigations.EFW >90 <97th centile.1 episode of PV bleeding of unknown origin >24/40.BMI <18 with no history of eating disorder.
<u>Skeletal complications</u> <ul style="list-style-type: none">Previous fractured pelvis	<u>Skeletal complications</u> <ul style="list-style-type: none">Spinal Issues (for consideration as to whether this will impact on birth or epidural/spinal anaesthesia).	<u>Skeletal complications</u>
<u>Previous gynaecological history</u> <ul style="list-style-type: none">MyomectomyHysterotomyCone biopsyLLETZ x 2Any uterine perforation resulting from previous STOP or surgery.	<u>Previous gynaecological history</u> <ul style="list-style-type: none">LLETZ X1 (for review of depth of excision)History of significant cervical excisional eventi.e. LLETZ where >10mm depth removed, OR >1 LLETZ procedure carried out OR cone biopsy (knife or laser, typically carried out under general anaesthetic).	<u>Previous gynaecological history</u>
<u>Mental health</u> <ul style="list-style-type: none">Known serious mental health illness such as bi-polar disorder	<u>Mental health</u> <ul style="list-style-type: none">Women taking psychotropic prescribed by GP (may need additional support plan	<u>Mental health</u> <ul style="list-style-type: none">History of mental health problems/depression not currently taking medication and stable

*It is noted that the above is not exhaustive and clinicians should exercise clinical judgment
1= Definition of recurrent altered fetal movement = at least 2 episodes of altered fetal movements are reported within a 21-day period (All Wales Altered Fetal Movement Guideline, WMNN 2021).