Risk assessment

Performed at booking and at regular intervals throughout pregnancy

Major risk factors for SGA and FGR

Maternal demographics

- Maternal age ≥ 40 years (nulliparous)
- Continued smoking ≥ 16 weeks' gestation (> 10 per day)

Recreational drug use Previous pregnancy history

- Previous FGR pregnancy**
- Previous hypertensive disease of pregnancy*[†]
- Previous stillbirth[†]

Maternal medical history

- Chronic hypertension*†
- Diabetes with vascular disease*†
- Renal impairment*†
- Antiphospholipid syndrome*†

Current pregnancy risk

- Heavy bleeding < 20 weeks' gestation
- Pre-eclampsia or gestational hypertension
- Antepartum haemorrhage or placental abruption

Minor risk factors for SGA and FGR

Maternal demographics

- Nulliparity
- Maternal age ≥ 40 years (multiparous)
- Smoking 1 to 10 cigarettes per day

Previous pregnancy history

 Short (< 6 months) or long (> 60 months) interpregnancy interval

Maternal medical history

- Conception via assisted reproductive technology
- BMI > 30 kg/m² or < 18.5 kg/m²

Current pregnancy risk

- Placenta praevia
- Low gestational weight gain

Antenatal screening

Starting from 20 to 24 weeks' gestation and performed until birth

Low risk: no major and < 3 minor risk factors

Serial fundal height assessment from 26 to 28 weeks' gestation until birth, plotted on customised antenatal growth chart

Suspected FGR

Slowing of customised fundal height > 30 centiles

or

Customised fundal height centile < 10

Increased risk

One or more major risk factors

- Start low-dose aspirin if indicated*
- Monthly serial ultrasound growth assessments starting from 28 to 30 weeks' gestation until birth

Risk factors for early-onset FGR[†]

• Monthly serial ultrasound growth assessments starting from 24 to 26 weeks' gestation until birth **plus** UtA Doppler at 20 to 24 weeks

Three or more minor risk factors

 Consider two ultrasound growth assessments in the third trimester: one at 30 to 32 weeks' gestation and one at 36 to 38 weeks' gestation

Unreliable fundal height measurement[§]

• Two ultrasound growth assessments in the third trimester: one at 30 to 32 weeks' gestation and one at 36 to 38 weeks' gestation



Specialist referral for review within 1 to 2 weeks[‡] **Normal UA Doppler** Abnormal UtA and/or Abnormal CPR (if ≥ 32) weeks' gestation) or EFW < 3rd centile Specialist referral for review within 1 week[‡] • If ≥ 38+0 weeks, review within 48 hours for consideration of planned birth Abnormal UA Doppler Forward flow present Same day referral for specialist review

Abnormal UA Doppler

AEDF or REDF

Urgent inpatient management

Normal UA Doppler

• EFW ≥ 3rd centile

gestation)

Normal UtA Doppler

Normal CPR (if ≥ 32 weeks'

* Low dose aspirin is recommended, starting between 12+0 and 16+6 weeks' gestation, taken at night.

[†]Risk factors for early-onset FGR include previous FGR birth < 32 weeks' gestation, previous hypertensive disease with birth < 34 weeks' gestation, significant maternal medical disease and previous stillbirth (particularly early gestation or FGR).

[‡]Clinical concern may override the recommended timeframes for specialist review (eg, oligohydramnios, significant slowing of growth or reduced fetal movements).

§ Unreliable fundal height measurements may be due to BMI > 35 kg/m² large or multiple fibroids or polyhydramnios.

Abbreviations: AC = abdominal circumference; AEDF = reversed end-diastolic flow; CPR = cerebroplacental ratio; EFW = estimated fetal weight; FGR = fetal growth restriction; PI = pulsatility index; RDEF = reversed end-diastolic flow; SGA = small for gestational age; UA = umbilical artery; UtA = uterine artery.

Management of FGR < 32+0 weeks (early onset)

			ſ	 Perform every two weeks Ultrasound for growth, UA Doppler, liquor Clinical review If fetal growth normalises to > 10th centile with all normal Dopplers over ≥ 1 month, transfer to routine low-risk care 		
view history Confirm gestational age Intenatal combined or Inaternal serum screening esults Intenatal screening for Ineuploidy and other Ineuploidy and Ineuploidy		Isolated SGAEFW and/or AC 3rd to < 10th centile				
	ſ	Normal UA and UtA Dopplers Once > 32+0 weeks'				//
		 FGR EFW or AC < 3rd centile or EFW or AC < 10th centile plus Abnormal UA (forward flow present) or Abnormal UtA Doppler (performed once at diagnosis) Once ≥ 32+0 weeks' gestation, manage as per late-onset flowchart 	At least weekly Every two weeks • Ultrasound for UA Doppler, liquor • Ultrasound for growth • cCTG (or CTG)* • Ultrasound for growth • Clinical review • Ultrasound for growth • Consider antenatal steroids • Ultrasound for growth • Consider antenatal steroids • Increase surveillance and/or consider inpatient monitoring if there is aligohydramnios, so or very poor interval growth or suspected pre-eclampsia. bhormal UA (forward ow present) or bhormal UA Doppler performed once at iagnosis) • Admit for birth planning > a 2 32+0 weeks' attion, manage as per onset flowchart • Antenatal steroids < 35+0 weeks ± magnesium sulphate < 30+ 0 weeks	At least weekly	Every two weeks	
				 Ultrasound for UA Doppler, liquor cCTG (or CTG)* Clinical review 	Ultrasound for growth	$\left \right\rangle$
				 Consider antenatal steroids Increase surveillance and/or consider inp or very poor interval growth or suspected 	patient monitoring if there is oligohydramnios, static d pre-eclampsia.	
Aedical, FGR risk factors consider serology for congenital infection creen especially if < 28 veeks' gestation or severe GR (EFW < 3rd centile) view fetal anatomy Consider referral for ertiary review including unatomical survey,	┥					
				Admit for birth planning In-patient management • Antenatal steroids < 35+0 weeks ± magnes • Twice daily cCTG (or CTG)* • At least daily maternal BP and pre-eclamp • UA and DV Doppler, liquor performed two to	sium sulphate < 30+ 0 weeks osia assessment to three times per week	
especially if < 28 weeks' Jestation or severe FGR		FGR at high risk of deterioration				
		AEDF or REDF		 If absent or reversed DV a-wave If reduced cCTG (or CTG) STV* By pre-labour caesarean 		
TG is preferred in the assessment of being should be assessed using con rig for birth are STV < 2.6 ms (26+0 to	early-o vention o 28+6 v	nset FGR. If cCTG is not available, fetal al CTG and fetal Doppler studies. STV veeks' aestation) and STV < 3.0 ms (29+0				

Absolute indications for birth

- Abnormal fetal heart rate (eg, repetitive unprovoked decelerations on cCTG or CTG)
- Maternal deterioration (eg, severe pre-eclampsia with uncontrolled hypertension/HELLP syndrome or other end-organ damage)

* cCTG is preferred in the assessment of early-onset FGR. If cCTG is not available, fetal wellbeing should be assessed using conventional CTG and fetal Doppler studies. STV criteria for birth are STV < 2.6 ms (26+0 to 28+6 weeks' gestation) and STV < 3.0 ms (29+0 to 31+6 weeks' gestation). **Abbreviations:** AC = abdominal circumference; AEDF = absent end-diastolic flow; BP= blood pressure; CPR = cerebroplacental ratio; cCTG = computerised cardiotocograph; CTG = cardiotocograph; DV = ductus venosus; EFW = estimated fetal weight; FGR = fetal growth restriction; HELLP = haemolysis, elevated liver enzymes and low platelets; IOL = induction of labour; REDF = reversed end-diastolic flow; UA = umbilical artery; UtA = uterine artery.

Management of FGR should be individualised.

Re

Increased surveillance or expedited birth should occur if there are features of concern (eg, cessation of growth, oligohydramnios, repeated episodes of reduced movements)

Management of FGR ≥ 32+0 weeks (late onset)



Perform every two weeks

• Maternal deterioration (eg, severe pre-eclampsia with uncontrolled hypertension/HELLP syndrome or other end-organ damage)

Management of FGR should be individualised.

restriction; HELLP = haemolysis, elevated liver enzymes and low platelets; IOL = induction of labour; REDF = reversed end-diastolic

flow; UA = umbilical artery; UtA = uterine artery.

Increased surveillance or expedited birth should occur if there are features of concern (eg, cessation of growth, oligohydramnios, repeated episodes of reduced movements).

	For	All Babies			
1) Review fetal growth monitoring	2) Review risk facto	ors for FGR	3) Calculate customised birthweight centile using GROW		
customised birthweight centile ≥ 10 →	Antenatal FGR diagnosis including evi abnormal Doppler studies)	idence of placental insuffi	ficiency (for example, NOT FGR	NO NOT FGR	
Customised birthweight centile ≥ 3 to 0	Calculate z-score for BMI and length ± skin/body fat, where expertise and equipment allow	 Two or more of the foll BMI z-score < -1.3 Length z-score < -1.4 Skin/body fat z-score Antenatal FGR diag Major FGR risk factore Placental insufficient 	lowing? .3 ore < -1.3 gnosis or ncy on histology	YES Neonate with FGI	
customised birthweight centile < 3				Neonate with FGF	
	Management O	f Neonates With FGI	R		
 Vonitor with NOC/NEWS for ≥ 24 hours Screen for hypoglycaemia for 12 to 24 hours Arrange paediatric/neonatal review for any of: Customised birthweight centile < 3 Confirmed or suspected genetic abnormality Confirmed or suspected congenital infection Disproportionate microcephaly or macrocephaly Poor postnatal growth Neonatal diagnosis of FGR with no evidence of placental insufficiency (abnormal Doppler) 	 Paediatric/neonatal review Full examination Review maternal serology and Doppler studies Review newborn hearing screen 	Evidence of plate No further investive No clinical evid first-line investive • FBC • Urine CMV F • Consider pl • Ensure new Suspected corr • CMV serolo • Syphilis serolo • Toxoplasmo Suspected corr • Toxoplasmo Suspected corr • FBC, LFT, tot • Urine CMV F • Ensure new Suspected ger • Molecular k • If aneuploid	acental insufficiency and nestigation needed dence of placental insufficients stigation PCR lacental histology aborn hearing screen is com- ngenital infection: materna ogy (IgG and IgM) ology (EIA screen initially) if ology (IgG and IgM) ology (IgG and IgM) if not c osis serology (IgG and IgM) ngenital infection: neonata cal and conjugated bilirubin PCR aborn hearing screen is com- netic disorder caryotype (EDTA) OR dy is suspected, FISH and sta	no genetic or infection concerns iency (eg, normal Doppler studies): hpleted al first-line investigations f not tested in third trimester clearly immune f first-line investigations hpleted andard karyotype (heparin)	