

Curriculum 2024 Guide for Subspecialty Training (SST): Maternal and Fetal Medicine

June 2024 V1.0



Version Control		
Version	Modifications	Date
1.0	Final version for publication	June 2024



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1. Maternal and Fetal Medicine SST

Doctors who undertake Maternal and Fetal Medicine (MFM) SST will develop the skills needed to provide the highest level of care for women with pregnancies potentially, or actually, affected by the full range of medical and/or fetal problems. Trainees will learn how to lead and co-ordinate care for these women, in association with a wide range of other general and maternity care providers. They will become leaders for these services at a local, regional and potentially even national level, with key roles in education, training, innovation, quality management and improvement, research and governance, pertinent to high-risk pregnancy services.

Through this SST, trainees will learn how to be excellent communicators who can cooperatively reach complex and often difficult decisions with individuals, couples and their families, and other healthcare providers. To achieve this, they will develop an extensive knowledge base, a logical mind, objectivity, empathy and advanced listening skills. Trainees are expected to be non-judgemental, free from bias, and able to negotiate and compromise. They should be kind, decisive when called upon, reflective and supportive. They will have the opportunity to develop a high level of technical expertise, to safely and effectively perform the procedures required in their subspecialty consultant post.

There are three main components to MFM SST. The first element is the clinical knowledge and skills required to become an MFM subspecialist, described by the Fetal Care, Prenatal Diagnosis, Pregnancy Care and Maternal Medicine SITMs and MFM Capabilities in Practice (CiPs). The clinical CiPs show the practical procedures a learner will become proficient in by the end of training. The second element is the generic skills required for all consultants, namely those of clinical governance, teaching, research, leadership and management. However, these must be acquired and developed in a working environment that cares for pregnant women. These skills run through both the subspecialty CiPs and stage 3 of the Core Curriculum. The third element is the subspecialty research CiP, which builds on the research CiP in the Core Curriculum 2024 and will provide the learner with research skills specific to the maternal and fetal medicine field.

As a learner progresses through the subspecialty training, they will learn how to handle a variety of scenarios. Learners will also participate in educational events to further develop their training. Throughout training, learners will need to reflect on whether a project has gone well, learn from positive and negative experiences, and use this to improve their own skills.

Before signing off on this SST, the Subspecialty Training Programme Supervisor (STPS) will decide the level of supervision required for each MFM CiP. If this and the final subspecialty assessment is satisfactory, subspecialty training accreditation will be awarded. More detail

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is provided in Section 6 of the Definitive Document for Maternal and Fetal Medicine SST 2024.

Design of the SST Programme

The 2024 MFM SST is made up of the four CiPs from the Special Interest Training Module (SITM) in Pregnancy Care (CiPs 1-4), the two CiPs from the Maternal Medicine SITM (CiPs 1-2), the three CiPs from the Fetal Care SITM (CiPs 1-3), the three CiPs from the Prenatal Diagnosis SITM (CiPs 1-3) and three corresponding subspecialty-specific CiPs that develop the knowledge and skills from the SITM curricula to a subspecialty level (SST MFM CiPs 1–3).

In addition to the clinical CiPs, a new CiP addressing skills in research and innovation has been developed as a generic CiP for all subspecialty trainees (SSTR).

Pregnancy Care SITM	Pregnancy Care CiPs 1–4
Maternal Medicine SITM	Maternal Medicine CiPs 1–2
Fetal Care SITM	Fetal Care CiPs 1–3
Prenatal Diagnosis SITM	Prenatal Diagnosis CiPs 1–3
SST-specific CiPs	SST MFM CiPs 1–3
SST Research CiP	SSTR CiP

Trainees need to complete all 16 CiPs to achieve subspecialty accreditation. The SST MFM CiPs can only be completed as part of an accredited subspecialty training programme in Maternal and Fetal Medicine. If a trainee has completed part or all of the Pregnancy Care SITM (CiPs 1-4), Maternal Medicine SITM (CiPs 1-2), Fetal Care SITM (CiPs 1-3) or Prenatal Diagnosis SITM (CiPs 1–3) before starting the MFM SST, they will not need to repeat them.

Learners with previous research experience, such as the Special Interest Professional Module (SIPM) in Clinical Research, can use this as evidence toward the SSTR CiP.

The 'indicative' duration of MFM SST is 24 months. If a learner undertakes the programme as a full-time trainee from the beginning of ST5, entering with minimal relevant skills or experience, it is expected to take longer than 24 months. They will still have 36 months left in their training programme. However, this timeframe is a guide only, as training is entirely competency based. Evidence of skills and competencies acquired before starting the MFM SST can be used, where appropriate, as evidence supporting sign-off within the subspecialty curriculum.

The majority of trainees will be still working toward their 'Certificate of Completion of Training' (CCT), although some will have already obtained this. Trainees who are pre-CCT will also need to continue accumulating skills and evidence as described for the Core Curriculum CiPs. Further detail can be found in the Definitive Document for Core Curriculum 2024 and the Essential Curriculum Guide.

Here is the GMC-approved MFM SST:



3. Capabilities in Practice (CiPs)

Pregnancy Care CiPs 1-4

Pregnancy Care CiP 1: The doctor demonstrates the skills needed, and can apply their knowledge, to manage antenatal care for a pregnant person with common medical problems.

problems:			
Key skills	Descriptors		
Able to take a thorough medical history from the pregnant person	 Demonstrates the ability to take a thorough medical history and considers how pregnancy may affect the medical problem presentation and how the condition may affect the pregnancy. Demonstrates the ability to record significant family history, drug history (including interactions and pregnancy safety), past medical history and systemic enquiry, including red flags. 		
Risk assesses the pregnant woman with co-existing medical conditions and plans for her pregnancy, in conjunction with specialist services	 Is able to risk assess women with medical problems and stratify them into low, medium or high-risk groups: those who can be managed using local expertise (category A); those who need clinical review and ongoing advice and guidance from the Maternal Medicine Centre (category B); and those whose care in pregnancy is best led by the Maternal Medicine Centre (category C). Knows the limits of their knowledge and can communicate effectively with other specialities locally, and with the Maternal Medicine Network, to best manage the care of a pregnant person. Working within guidance and thresholds determined by the local Maternal Medicine Network, is able to: assess a woman with a pre-existing medical condition preparing for pregnancy, and work with her to put together an appropriate plan. evaluate and advise on drug therapy for medical conditions and tailor treatment when this would have a detrimental effect on pregnancy. assess conditions that will have a significant impact on the outcome of pregnancy for a mother and her baby. assess conditions where pregnancy may significantly deteriorate the health of a woman with a pre-existing medical condition and the surveillance required to limit risk. 		



Diagnoses and provides	 access additional information needed to best manage complex medical conditions. put together a delivery plan that minimises risk to a mother and her baby. work in partnership with the woman to plan her care and delivery. Refers to other medical and maternal medicine specialists, in line with local guidance. Understand what investigations are needed to explore
initial management for	common medical presentations, including shortness of
common acute medical	breath, chest pain, headache, collapse, abdominal pain and
presentations in pregnancy	fever/sepsis.
	 Constructs a differential diagnosis and requests appropriate investigations.
	 Initiate appropriate emergency management and liaise with allied specialities for an ongoing plan of care.
	Understands the impact of, and interplay between, mental
	health conditions and maternal medicine conditions, and
Diagraph and manager	addresses this in management plans.
Diagnoses and manages	Is able to assess and counsel women with hypertensive disorders, or at risk of a programmy indused hypertensive
hypertensive disorders in	disorders, or at risk of a pregnancy-induced hypertensive disorders, pre-conceptually.
pregnancy	 Understands and recognises the diverse aetiology of
	hypertension in pregnancy, whether pre-existing or arising in pregnancy.
	 Understands the risks that hypertensive disorders pose to
	pregnant people and can plan safe surveillance and management in the antenatal period.
	 Understands the risks that hypertensive disorders pose to a
	baby and can plan safe surveillance in the antenatal period.
	 Safely manages the hypertensive disorders in a woman in labour.
	Understands and can create a safe management plan for a
	woman with severe pre-eclampsia and the complications of this condition.
	Liaises with the multidisciplinary team (MDT), including the
	tertiary centre, where appropriate, to optimise the care of a woman with hypertensive disorders.
	Works in partnership with the woman with a hypertensive
	disorder to plan her care and delivery.
	 Understands the long term implications of hypertensive
	disorders of pregnancy on the health and wellbeing of
	mother and baby.



	 Plans appropriate follow-up for a woman with a hypertensive disorder during pregnancy.
Diagnoses and manages disorders of glucose metabolism in pregnancy	 Assesses and agrees a plan for the woman who has preexisting diabetes to prepare for pregnancy. Demonstrates knowledge of the risk that pre-existing diabetes has on a mother and her baby. Works effectively in the MDT to provide the best possible care for a pregnant woman with pre-existing diabetes during pregnancy and in labour. Refers to the tertiary centre in more complex cases to access specialist care for a pregnant person with diabetes during pregnancy Diagnoses and can counsel a woman who develops diabetes during pregnancy. Devises a safe plan for maternal and fetal surveillance during pregnancy. Can recognise and manage the acute complications of diabetes in pregnancy e.g. diabetic ketoacidosis. Plans for a woman with diabetes to safely give and is able to adapt the plan to changing circumstances. Safely manages the delivery of a woman with diabetes. Works in partnership with a woman to plan her care and delivery. Understands the long term implications of disorders of glucose metabolism in pregnancy on the health and wellbeing of the mother and her baby. Plans appropriate follow-up care for a pregnant person with diabetes.
Diagnoses and manages common endocrine disorders in pregnancy	 Assesses and agrees a plan for the woman with hypothyroidism. Assesses and agrees a plan for the woman with hyperthyroidism. Manages the woman with micro- and macroprolactinoma safely through pregnancy.
Supports the health and wellbeing of a morbidly obese pregnant woman	 Is able to risk assess and plan for pregnancy and delivery, including women who have undergone bariatric surgery. Is able to work with the woman to manage weight gain and create a suitable plan that encourages healthy nutrition. Discusses and negotiates the most appropriate mode of delivery, taking into account patient choice and the safest delivery option.



	 Advises on modifications to birth that can enhance safety and the experience of the woman with morbid obesity. Liaises with midwifery and anaesthetic colleagues to 	
	provide the best possible care.	
Supports a pregnant	 Is able to risk assess the pregnant person with an eating 	
woman with an eating	disorder and make plans for her pregnancy.	
disorder	Can safeguard the wellbeing of both the mother with an	
	eating disorder and her baby.	
Evidence to inform decision – examples of evidence (not mandatory requirements)		

•	Reflective practice	•	RCOG Learning
•	NOTSS	•	Local and deanery teaching
•	TO2	•	Attendance at appropriate courses and
•	CbD		conferences
•	Mini-CEX	•	Attendance at specialist diabetes
			antenatal clinics
		•	Attendance at maternal medicine clinics
		•	Log of cases and outcomes

Mandatory requirements

No mandatory evidence

- Awareness and understanding of local maternal medicine networks and when to make referrals and involve the MDT
- The pathophysiology, definition, diagnosis, associated acute and long-term maternal and fetal complications, and best practice for managing pre-eclampsia and its variants
- The pathogenesis and classification, prevalence and complications of pre-existing diabetes (e.g. metabolic, retinopathy, nephropathy, neuropathy and vascular disease)
- Monitoring and optimisation of glucose control during labour
- Management of hypoglycaemia and ketoacidosis in pregnancy and labour
- The pathophysiology, presentation and implications for maternal and/or fetal health of common maternal conditions present at booking or that occur during pregnancy
- The aetiology, incidence, diagnosis, management; obstetric, medical and neonatal complications, and recurrence chance of each condition
- The interpretation of electrocardiograms (ECG), chest x-rays and blood gases analysis and how they are influenced by pregnancy
- How pregnancy alters physiology and what impact this has on medical conditions that are present, and how results of investigations should be interpreted during pregnancy
- The impact of drug treatment on the health of the mother and her babe
- The incidence, associated obstetric, medical and neonatal complications of the pregnant obese woman
- The endocrinology of obesity



- Weight reduction strategies and appropriate nutrition in managing the pregnant obese woman
- The risks associated with increased body mass index (BMI) in pregnancy and postpartum, and how these may be minimised
- The steps that can be taken before pregnancy to reduce the risks of morbid obesity during pregnancy

Pregnancy Care CiP 2: The doctor demonstrates the skills needed, and can apply their knowledge, to manage the care of a pregnant woman whose pregnancy is complicated by infection which may affect the health of her baby.

Key skills **Descriptors** Manages the care of a Demonstrates a knowledge of the implications for pregnancy of variety of infections: HIV, syphilis, pregnant woman with cytomegalovirus (CMV), toxoplasmosis, hepatitis B and C, infections that can affect herpes simplex virus (HSV), parvovirus and chicken pox their health and that of their (varicella). baby Is able to interpret laboratory results for each infection by working closely with virology. Explains the potential effects of infections on the baby, newborn and long-term effects of fetal infections. Recognises when to refer a pregnant person with an infection and understands how best to share care and monitor them. Works with the tertiary centre and MDT. Works with the MDT to create a plan for medications for the mother during the birth and for the baby postnatally. Gives appropriate advice to minimise the risk of vertical transmission. Evidence to inform decision – examples of evidence (not mandatory requirements) Reflective practice **RCOG Learning** NOTSS Local and deanery teaching TO2 Attendance at appropriate courses and CbD conferences Log of cases and outcomes Mini-CEX **Mandatory requirements** No mandatory evidence

Knowledge criteria

The clinical features, prevention, vertical transmission risk and ultrasound features of CMV, toxoplasmosis, parvovirus and varicella. Understands the short- and longer-term



implications for the baby and newborn of contracting these infections, as well as the laboratory investigation that are needed and how to manage them during pregnancy

• The role of the clinical virologist and the limitations of any antenatal treatment options

Pregnancy Care CiP 3: The doctor demonstrates the skills needed, and can apply their knowledge, to manage the postnatal care of a pregnant person with common medical problems.

Key skills	Descriptors
Manages the care of a woman with medical conditions in the postnatal period – evidence for a variety of conditions but must include diabetes Evidence to inform decision –	 Discusses plans for contraception, tailored to the woman's needs. Makes sure that the woman receives follow-up care in an appropriate setting. Can discuss the long-term implications of medical conditions on the woman's health and wellbeing. Supports the woman to limit the effect of her medical conditions on future pregnancies. examples of evidence (not mandatory requirements)
a Doflostivo prostico	a DCOC Learning

•	Reflective practice	•	RCOG Learning
•	NOTSS	•	Local and deanery teaching
•	TO2	•	Attendance at specialist diabetes antenatal
•	CbD		clinics
•	Mini-CEX	•	Attendance at maternal medicine clinics
		•	Log of cases and outcomes

Mandatory requirements

No mandatory evidence

- Contraception in the postnatal period
- Provision of long-acting contraceptives
- Implications of medical conditions on the wellbeing of mother and baby, and understands the impact on further pregnancies

Pregnancy Care CiP 4: The doctor provides holistic care to a pregnant person.		
Key skills	Descriptors	
Is able to apply legal and ethical principles in pregnancy care, where this is needed	 Is able to screen for and organise safeguarding of a woman at risk of domestic violence. Can screen for and organise safeguarding of the neonate at risk of harm. 	



	 Is able to counsel and complete an advance directive (recording decisions on healthcare in preparation for a future event) for the woman who declines blood products.
Provides the best possible	 Is aware of the effect of social deprivation on pregnancy
outcomes for a pregnant	outcomes.
person who is socially	 Understands the prevalence of domestic violence, the
vulnerable	need to screen all women for this and agree a plan to
	safeguard the pregnant person and their children.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- NOTSS
- TO2

- CbD
- Mini-CEX
- Attendance at pre-birth planning meetings with the safeguarding team

Mandatory requirements

No mandatory evidence

Knowledge criteria

- How social disadvantage can cause medical and neonatal complications, and legal consequences of social disadvantage with respect to: domestic violence, teenage pregnancy and asylum seekers
- The influence of ethnic and religious background on obstetric expectations and outcome
- The law in relation to seeking asylum
- When and how to use different agencies involved in processing claims for asylum seekers and meeting their practical needs
- The role of different agencies (social services, police and voluntary groups) in investigating suspected domestic violence and protecting vulnerable women and children
- The law in relation to physical and sexual assault, bodily harm and rape
- Female genital mutilation (FGM) procedures and their consequences, including for pregnancy and birth
- Child protection issues associated with FGM
- Religious beliefs and customs that may affect healthcare or consent for medical interventions

3.2 SST Maternal Medicine CiPs 1–2

Maternal Medicine CiP 1: The doctor is able to work with others to provide high quality care to the woman with medical conditions in pregnancy or planning a pregnancy.

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Descriptors



Effectively expensive extens	D. Halana dha la an d'Hanfida Barana a Cara CITA an d
Effectively communicates	Builds on the key skills of the Pregnancy Care SITM and Section 1.
with the team providing care	uses them when working on the full range of medical
	problems which may complicate pregnancy.
	Works collaboratively across specialties and, where Albarant managed spinises are through MDT.
	relevant, manages clinical networks through MDT
	meetings to construct pre-pregnancy, antepartum,
	intrapartum and postpartum management plans to
	ensure that high quality care is available locally to women with complex medical problems.
	·
	 Plans care for women with complex medical problems in collaboration with other specialties. Makes appropriate
	referral to a regional maternal medicine clinic, where
	relevant. (In England this will be through the Maternal
	Medicine Networks either through MDTs for category B
	medical problems or referral for ongoing care to a
	Maternal Medicine Centre for category C medical
	problems).
	 Is aware of a possible genetic diagnosis that may not have
	been diagnosed to date. Refers to clinical genetics as
	appropriate.
Provides tailored pre-	Can advise the person with complex medical conditions of
pregnancy counselling	the impact of pregnancy on their condition.
	Is able to advise the person with complex medical
	conditions on the impact of the condition on their
	pregnancy.
	Is able to advise on modifications that will optimise her
	health before embarking on pregnancy.
	Is able to adjust medication to the safest regime for
	pregnancy.
	Is able to put together a plan so the person knows what
	to expect once they become pregnant.
	Is able to advise on the timing of pregnancy.
	Is able to advise someone against conception in
la abla ta cancidantha	circumstances where the risk of pregnancy is too great.
Is able to consider the	Is familiar with the anaesthetic considerations for the
anaesthetic implications of	person with a variety of medical conditions.
maternal conditions, liaise	 Is able to work with anaesthetic colleagues to assess pregnant persons with complicated medical conditions
with anaesthetic colleagues	and put together a plan to keep the person and the baby
and plan according to	safe during pregnancy, delivery and the postnatal period.
someone's needs	 Demonstrates familiarity with the effect of different
	intrapartum analgesia to make sure persons with
	complex medical conditions are safe in labour.



	Participates in obstetric anaesthesia clinics.	
Can perform a risk benefit analysis of investigations and treatments that could be	Knows which investigations and medications are appropriate and can discuss the safety of these for the mother and fetus. And the training and the safety of the	
used during pregnancy	 Is able to interpret tests e.g. chest x-ray, artificial blood gas (ABG) and electrocardiogram (ECG), lung function tests and echocardiogram. Demonstrates understanding of the effects of drugs used for maternal indications on the fetus. 	
	 Understands and accommodates the physiological effects of pregnancy on interpreting laboratory results and the pharmacokinetics of any drugs used. 	
Evidence to inform decision – examples of evidence (not mandatory requirements)		

•	Reflective practice	•	RCOG Learning
•	NOTSS	•	Local and deanery teaching
•	TO2	•	Attendance at obstetric anaesthesia
•	CbD		clinics
•	Mini-CEX	•	Attendance at maternal medicine

Mini-CEX network meetings

Mandatory requirements

No mandatory evidence

- Local team structures, networks and guidelines for the management of medical conditions in pregnancy and outside of pregnancy.
- Awareness and understanding of local Maternal Medicine Networks and regional thresholds when to make referrals and include the MDT. Knows when it is appropriate to manage locally, or to manage locally with input from the regional maternal medicine clinic/the Maternal Medicine Centre and when referral to regional clinics/centres is advised.
- Criteria for referral to Maternal Medicine Centres/regional clinics.
- Structure of the Maternal Medicine Networks/regional clinics.
- In England categories for level of care within the Maternal Medicine Networks i.e. category A, B and C.
- When to seek specialist input.
- The structure and organisation of high dependency, intensive care and outreach teams.
- Indications for high dependency and intensive care.
- Methods of invasive monitoring for oxygenation, acid base balance, intra-arterial pressure, cardiac output, preload and contractility.
- The principles and practice of palliative care.



Maternal Medicine CiP 2: The doctor has a high level of understanding of the impact that medical conditions have on pregnancy and is able to optimise care for the affected woman.

Kov skills	Descriptors	
Key skills	Descriptors	
Is able to manage care for the pregnant person who has renal problems in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to minimise the risks the woman's medical condition. Can construct a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby. Is able to recognise the presentation of renal disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions. Understands which tests are appropriate in pregnancy for diagnosis and monitoring, and which are not valid or have different reference ranges in a pregnant woman. Understands when tests pose an additional risk to the mother or fetus, and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high. Has a good working knowledge of medical treatments for renal conditions that are safe in pregnancy, and can modify treatments when they are not safe. Knows how to 	
Is able to manage care for someone who has haematological problems in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated with the woman's medical condition. Can construct a plan for pregnancy, delivery and the neonatal period to reduce the risk to the fetus/baby. Is able to recognise the presentation of haematological disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions. Understands which tests are appropriate in pregnancy for diagnosis and monitoring of haematological disorders, and which are not valid or have different reference ranges in a pregnant woman. Understands when tests pose an additional risk to the mother or fetus, and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high. 	



	 Has a good working knowledge of medical treatments for haematological conditions that are safe in pregnancy, and can modify treatment when they are not safe. Knows how to access advice on safety.
Is able to manage care for someone who congenital and acquired cardiac conditions in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated with the woman's medical condition. Can construct a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby. Is able to recognise the presentation of cardiac disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions. Understands which tests are appropriate in pregnancy for diagnosis and monitoring of cardiac disorders, and which are not valid or have different reference ranges in the pregnant woman. Understands when tests pose an additional risk to the mother or fetus and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high. Has a good working knowledge of medical treatments for cardiac conditions that are safe in pregnancy and is able to modify treatment when they are not safe and knows
Is able to care for someone who has inflammatory conditions (connective tissue disorders, inflammatory bowel disease and dermatological problems) in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated with the woman's medical condition. Can construct a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby. Is able to recognise the presentation of inflammatory or dermatological conditions in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions. Understands which tests are appropriate in pregnancy for diagnosis and monitoring of inflammatory disorders, and which are not valid or have different reference ranges in a pregnant woman. Understands when tests pose an additional risk to the mother or fetus and is able to discuss the relative risk and



	benefits of this. Can support a woman who deems the risk too high.
	 Has a good working knowledge of medical treatments for inflammatory disorders that are safe in pregnancy, including biologics. Is able to modify treatment when they are not safe and knows how to access advice on safety.
Is able to manage care for someone who has epilepsy and other common neurological problems in	Can construct an appropriate plan for pregnancy, delivery and the postnatal period for women with epilepsy, multiple sclerosis, idiopathic intracranial hypertension and chronic headache.
pregnancy	 Can put together a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby.
	 Is able to recognise the presentation of neurological disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions.
	Can counsel a woman with epilepsy and other
	neurological problems to safeguard her baby.
Is able to care for someone who has liver disorders in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated with the woman's medical condition. Can construct a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby.
	 Is able to recognise the presentation of liver disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions.
	 Understands which tests are appropriate in pregnancy for diagnosis and monitoring of liver disorders, and which are not valid or have different reference ranges in the pregnant woman.
	 Understands when tests pose an additional risk to the mother or fetus, and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high.
	 Has a good working knowledge of medical treatments for liver conditions that are safe in pregnancy, and is able to modify treatment when they are not safe. Knows how to access advice on safety.



Is able to manage care for	Con construct on annualists along for a construct of the
someone who has HIV in	Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated
pregnancy	 with the HIV in pregnancy. Can construct a plan for pregnancy, delivery and the neonatal period to minimise the risk to the fetus/baby. Understands which tests are appropriate in pregnancy for diagnosis and monitoring, and which are not valid or have different reference ranges in a pregnant woman. Understands when tests pose an additional risk to the mother or fetus and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high. Has a good working knowledge of medical treatments for HIV conditions that are safe in pregnancy and is able to modify treatment when they are not. Knows how to access advice on safety and the criteria for commencing
	treatment during pregnancy.
Is able to care for someone who has respiratory compromise in pregnancy	 Can construct an appropriate plan for pregnancy, delivery and the postnatal period to reduce the risks associated with the woman's medical condition. Can put together a plan for pregnancy, delivery and the neonatal period to reduce the risk to the fetus/baby. Is able to recognise the presentation of respiratory disorders in pregnancy, can construct a differential diagnosis and work with the MDT to put together a suitable management plan for pre-existing or new onset conditions. Understands which tests are appropriate in pregnancy for diagnosis and monitoring, and which are not valid or have different reference ranges in a pregnant woman. Understands when tests pose an additional risk to the mother or fetus and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high. Has a good working knowledge of medical treatments for respiratory conditions that are safe in pregnancy and is able to modify treatment when they are not safe. Knows how to access advice on safety.
Is able to manage care for someone who has current or past malignancy in pregnancy	When malignancy is diagnosed in pregnancy, is able to support a woman through a tailored plan for treatment during pregnancy and provide them with reassurance of
	the suitability of this plan during.



- Is able to weigh up the timing of delivery around someone's treatment needs.
- When malignancy has been treated prior to pregnancy, is aware of the implications for maternal health during pregnancy and is able to mitigate against these.
- Is mindful of the fetal considerations when managing malignancy in pregnancy.
- Understands which tests are appropriate in pregnancy for diagnosis and monitoring of cancer, and which are not valid or have different reference ranges in a pregnant woman.
- Understands when tests pose an additional risk to the mother or fetus and is able to discuss the relative risk and benefits of this. Can support a woman who deems the risk too high.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- NOTSS
- TO2
- CbD
- Mini-CEX

- **RCOG Learning**
- Local and deanery teaching
- Attendance at appropriate courses and conferences (e.g. BMFMS, MOMS)
- Log of cases with outcomes
- Attendance at non-obstetric specialist medical clinics
- Attendance at maternal medicine **MDTs**

Mandatory requirements

No mandatory evidence

- The normal functional and anatomical changes of the different body systems during pregnancy (e.g. cardiovascular, respiratory, gastrointestinal, endocrine and haematological)
- The pathological changes in the function of these body systems in pregnancy
- Renal conditions understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of renal conditions predating and arising in pregnancy, and the effect of labour and birth on these conditions:
 - o acute renal impairment
 - o hydronephrosis
 - renal disease and hypertension
 - o glomerulonephritis
 - reflux nephropathy
 - renal transplant



- Haematological understands the risk factors, presentation, investigation, differential diagnosis management and outcomes of renal conditions predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - o sickle cell disease and crisis
 - o thalassaemia
 - o thromboembolic disease
 - bleeding disorders
 - disorders of platelets
- Cardiac understands the risk factors, presentation, investigation, differential diagnosis management and outcomes of cardiac conditions predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - o congenital cardiac disease
 - o ischaemic cardiac disease
 - mechanical and tissue valve replacements
 - peripartum cardiomyopathy
- Connective tissue disorders understands the risk factors, presentation, investigation, differential diagnosis management and outcomes of connective tissue disorders predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - System lupus erythematosus (SLE)
 - rheumatoid arthritis
 - o autoimmune lymphoproliferative syndrome (APLS)
- Gastrointestinal understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of gastrointestinal conditions predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - acute fatty liver
 - o Crohn's disease
 - ulcerative colitis
 - obstetric cholestasis
 - hyperemesis gravidarum
 - immune and infective hepatitis
 - liver transplant
- Dermatological conditions understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of dermatological conditions predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - psoriasis
 - o eczema
 - pemphigoid
 - o polymorphic eruption of pregnancy
 - o prurigo
 - o pruritic folliculitis
- Neurology understand the risk factors, presentation, investigation, differential diagnosis, management and outcomes of neurological conditions predating and arising



in pregnancy and the effect of labour and birth on these conditions:

- o multiple sclerosis
- epilepsy
- bell's palsy
- o migraine
- o stroke
- cerebral palsy
- HIV infection understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of HIV predating and arising in pregnancy and the effect of labour and birth on these conditions
- Current pharmacological management of HIV, and drug side effects
- Respiratory disease understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of respiratory conditions predating and arising in pregnancy and the effect of labour and birth on these conditions:
 - o asthma
 - cvstic fibrosis
- Malignancy understands the risk factors, presentation, investigation, differential diagnosis, management and outcomes of malignancy predating and arising in pregnancy and the effect of labour and birth on malignancy:
 - breast cancer
 - o leukaemia
 - lymphoma
- Genetics and disease inheritance of medical disorders the risk to the mother and to the fetus and screening options e.g. haemoglobinopathy
- How pregnancy can influence the findings of investigations and may alter treatment effects
- How the medical problem may deteriorate during pregnancy, how this might present, and how it would be managed
- Paediatric network guidelines for the management of newborn problems, including frameworks around extreme prematurity and antenatal parallel care planning
- The pharmacology of drugs used to manage these conditions
- The pregnancy and breastfeeding safety profile of drugs, chemotherapy and radiotherapy used to manage these medical conditions
- Recurrence risks for future pregnancies
- The best forms of contraception for women with these specific medical disorders

3.3 Fetal Care CiPs 1–3

Fetal Care CiP 1: Uses ultrasound skills to recognise, monitor and manage compromise to fetal wellbeing.

Key skills Descriptors



Uses ultrasound to screen, diagnose and manage fetal	 Understands the principles of transabdominal and transvaginal scanning, using ultrasound safely. Able to measure fetal biometry to monitor the fetus at 		
compromise	Able to measure fetal biometry to monitor the fetus at risk of growth restriction.		
	Able to recognise and manage early and late severe fetal		
	growth restriction (FGR), referring cases of early FGR to		
	tertiary services.		
	Able to recognise disorders of amniotic fluid volume and		
	plan accordingly.		
Uses Doppler studies to	Able to perform uterine artery Dopplers to assess the risk		
screen, diagnose and manage	of placental dysfunction.		
fetal compromise	 Able to perform umbilical artery Dopplers to assess fetal resilience. 		
	Able to perform middle cerebral artery (MCA) Dopplers to		
	evaluate fetal compromise.		
	Able to perform ductus venosus Dopplers to evaluate		
	fetal compromise.		
Uses ultrasound to assess	Able to use transvaginal scanning to diagnose and		
placental location	manage low-lying placenta.		
Discusses their findings with	Demonstrates the ability to communicate their findings		
the pregnant woman	and the degree of risk effectively so that the woman can		
, 3	be involved in an informed decision-making process.		
Assesses and plans the	Provides ongoing assessment of fetal biometry over time		
management and delivery of	when severe FGR is identified.		
a fetus with severe growth	Able to use fetal Dopplers – umbilical, MCA and ductus		
restriction	venosus – to assess fetal wellbeing and plan the timing of delivery.		
	Able to discuss gestation-related risk of delivery versus		
	continuation of pregnancy with the pregnant woman and		
	facilitate informed decision-making.		
Provides support and	Provides follow up after the birth and accesses support		
counselling post birth and for	services for the parents, where outcomes are		
future pregnancies	complicated or poor.		
	Explains additional information learned after the birth The control birth learned after the birth		
	e.g. placental histology.Able to make a plan for future pregnancies, outlining		
	recurrence risks and preventive strategies.		
Evidence to inform decision –	examples of evidence (not mandatory requirements)		
• NOTSS	Reflective practice		
• TO2	Attendance at appropriate courses e.g.		
• CbD	ultrasound theory/practice		
Mini-CEX	 Log of cases with outcomes 		

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Mandatory requirements

- **OSATS**
 - o fetal biometry and liquor volume
 - transvaginal placental localisation
 - umbilical artery Doppler
 - middle cerebral artery Doppler
 - o ductus venosus Doppler
 - uterine artery Doppler

- The risks associated with the different ultrasound modalities and how to limit them mechanical index (MI) and thermal index (TI)
- How to use machine controls to optimise the image, including: power, gain, focal length, magnification, sector width, frame rate, pulse repetition frequency, colour and power Doppler modes.
- The difference between small for gestational age (SGA) and fetal growth restriction (FGR)
- The differential diagnosis for fetal growth restriction
- How Doppler assessments are used to monitor growth restriction, timing of birth and detect fetal anaemia
- National guidance on monitoring for FGR, the timing of birth and signs that a referral need to be made to a subspecialist when managing FGR
- How fetal anomalies may influence the Doppler waveforms (e.g. cardiac arrhythmias, fetal anaemia, hydrops and twin-to-twin transfusion syndrome (TTTS))
- Definition of low-lying placenta and how to make the diagnosis using ultrasound
- Management of placenta praevia
- The risk factors for abnormal placental invasion (AIP) and vasa praevia and how to diagnose them using ultrasound, and/or when to refer to a regional AIP service
- Definition of oligohydramnios and polyhydramnios and the differential diagnosis, investigation and management

Fetal Care CiP 2: The doctor demonstrates the skills and attributes required to as	sess the
fetus at risk of red cell alloimmunisation.	

Key skills	Descriptors
Safely manages the pregnancy where there is a risk of red cell immunisation	 Provides appropriate antenatal care to the woman with a pregnancy at risk. Recognises when there is a risk of fetal anaemia. Explains the potential fetal and maternal risks of red cell antibodies. Liaises with blood transfusion and neonatal services. Classifies the risks for any pregnancy complicated by red cell antibodies. Performs and interprets the findings of a MCA Doppler.



Monitors the pregnancy at risk and understands the thresholds for referral to tertiary units with transfusion services.

Evidence to inform decision – examples of evidence (not mandatory requirements)

• NOTSS	Reflective practice
• TO2	 Evidence of MDT working
• CbD	RCOG Learning:
Mini-CEX	 observation of fetal blood
	transfusion

Mandatory requirements

- OSATS
 - o middle cerebral artery Doppler

Knowledge criteria

- Differential diagnosis for fetal anaemia
- Ultrasound and cardiotocography (CTG) changes secondary to severe fetal anaemia
- Which red cell antibodies may cause haemolytic disease of the fetus and newborn, and threshold antibody levels that carry significant risk
- When and how surveillance for fetal anaemia should be instituted
- How MCA velocities are used to monitor signs of anaemia
- Triggers for referral to a tertiary level unit capable of performing intrauterine transfusion
- Treatment of fetal anaemia
- The role of intravenous immunoglobulin (IVIgG) in haemolytic disease of the fetus and
- Management of the newborn risk of kernicterus

Fetal Care CiP 3: The doctor demonstrates the skills and attributes required to assess complications of twin pregnancies.	
Key skills	Descriptors
Uses ultrasound to monitor twin pregnancies	 Able to determine the chorionicity of a twin pregnancy when scanning in first trimester.
	 Able to assess and monitor a twin pregnancy using biometry and Doppler scanning techniques.
Manages complicated twin pregnancies	 Able to diagnose and make an initial assessment of growth discordancy in twin pregnancies. Able to discuss effectively the timing of delivery with
	parents and facilitate informed decision-making, considering the risk to both twins of delivery or continuing the pregnancy when there is growth

discordancy.



•	Refers to tertiary services when early and severe growth
	discordancy occurs.

- Able to assess and monitor the monochorionic twin pregnancy for presence and evolution of TTTS.
- Refers to tertiary services when there is evidence of TTTS or selective FGR in monochorionic twins.
- Assists with follow up after treatments for TTTS.
- Recognises the possibility of other complications of monozygotic twinning, including selective FGR, discordant anomalies, twin reversed arterial perfusion sequence (TRAP) and single intrauterine death, and refers appropriately to fetal medicine tertiary services.
- Is aware of the principles of management of higher multiples.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- NOTSS
- TO2
- CbD
- Mini-CEX

- Reflective practice
- Attendance at specialist twin clinics
- Log of cases with outcomes
- Observation of advanced procedures in the management of complicated twin pregnancies e.g. fetal reduction and laser ablation

Mandatory requirements

- OSATS
 - multiple gestation chorionicity
 - twin pregnancy assessment

- Definition of significant growth discordance in twin gestations and the importance of
- Management of growth discordancy in twin pregnancies
- The clinical and ultrasound features of TTTS, and referral triggers for fetal medicine subspeciality input
- Short and long-term outcomes from TTTS
- The management of TTTS and follow up regimes, following treatment
- The ultrasound features of TRAP and conjoined twins
- Ongoing management of a pregnancy complicated by co-twin death
- Other complications of multiple gestations that necessitate discussion with, or referral to, a tertiary fetal medicine service, e.g. discordant anomaly



3.4 Prenatal Diagnosis CiPs 1–3

Prenatal Diagnosis CiP 1: The doctor can use ultrasound to recognise where fetal anatomy is not normal.

Key skills	Descriptors
Demonstrates normal structural findings in all trimesters and recognises when they cannot be demonstrated	 Performs and records a detailed, systematic ultrasound of the fetus in line with NHS Fetal Anomaly Screening Programme (FASP) guidance. Understands the strengths and limitations of ultrasound for each organ system within each trimester. Explains normal anatomical views to the pregnant person. Documents and records normal anatomical views. Recognises when image quality is technically poor. Can explain next steps to the pregnant person if normal views cannot be obtained.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- **NOTSS**
- TO2
- CbD
- Mini-CEX
- RCOG Learning
- FASP online training
 - Local and deanery teaching

- Attendance at relevant courses and conferences
- Log of cases and outcomes
- Attendance at fetal medicine clinics
- Attendance at multidisciplinary team (MDT) meetings
- Attendance at specialist neonatal and paediatric clinics
- Examples of anonymised birth plans

Mandatory requirements

- OSATS
 - Fetal anomaly scan
 - Fetal echo

- The normal appearance on ultrasound scans, in all trimesters, of the fetal central nervous system (CNS), face and neck, thorax, cardiovascular system, abdominal wall and gastrointestinal tract, urogenital system, and the fetal skeleton and extremities
- Local protocols for follow up, if any, after an incomplete anatomy scan
- Normal embryology of all body systems, and the common fetal anomalies that can happen when they do not develop in the way they should, as identified by FASP.
- Normal fetal behaviour and activity, and abnormalities of this
- Fetal circulation, and how it adapts at birth
- Diagnostic features of each condition targeted by FASP, their differential diagnosis and chance of structural, chromosomal and syndromic associations. These conditions are Trisomy 21, 18 and 13, anencephaly, spina bifida, congenital diaphragmatic hernia, gastroschisis, exomphalos, renal agenesis, facial cleft, hypoplastic right or left heart and



lethal skeletal dysplasia

- The thresholds for diagnosing mild, moderate and severe ventriculomegaly measurements, and the potential implications of the different severities of ventriculomegaly
- The role of magnetic resonance imaging (MRI) for CNS lesions
- The difference between Dandy-Walker malformation, Dandy-Walker Variant and mega cisterna magna, the implications of each and the pitfalls in prenatal diagnosis
- The common fetal tachycardia and bradycardia arrhythmias and the role of the paediatric cardiologist in their management
- The different types of ventricular septal defect (VSD) and their association with cardiac, extracardiac and chromosomal anomalies. Understand the role of the paediatric cardiologist in their management
- The ultrasound features of transposition of the great arteries, atresia of either outflow tract, stenosis of either outflow tract, double outlet right ventricle or a common outflow tract (truncus arteriosus)
- The association of these conditions with further cardiac, extracardiac and chromosomal anomalies
- The role of the paediatric cardiologist in the management of fetal cardiac problems
- The ultrasound features of gastrointestinal (GI) atresia, associations and surgical options following birth
- The spectrum of ultrasound findings of echogenic bowel and its association with chromosomal anomalies, cystic fibrosis, growth restriction and viral infections
- Urinary tract obstruction and multi cystic dysplastic kidney (MCDK): aetiology, spectrum of severity, postnatal investigation and the likely short- and long-term impact of these conditions
- The local pathway for postnatal referral for talipes and the Ponseti approach to treatment
- Limb reduction defects: associations and aetiology
- Findings suggestive of lethal skeletal dysplasia and the features of the more common non- lethal dysplasias, particularly certain types of osteogenesis imperfecta and achondroplasia
- A differential diagnosis for non-immune hydrops, the need for tertiary referral and the range of investigations likely to be offered

Prenatal Diagnosis CiP 2: The doctor can assess and investigate a pregnancy where there are concerns about the fetus.	
Key skills	Descriptors
Can provide genetic counselling in common prenatal situations	 Takes medical history and constructs, where appropriate, a family tree for people who are pregnant, or have a chance of, genetic conditions.



Provides initial counselling for common fetal structural anomalies and manages people in partnership with tertiary fetal medicine services	 Explains common modes of Mendelian and multifactorial inheritance, and recurrence risks. Counsels for previous trisomy and monosomy X. Counsels for previous neural tube defect. Is experienced in carrying out ultrasound diagnosis and managing pregnancies complicated by fetal anomalies that are covered by the FASP. Discusses other potential prenatal tests such as fetal karyotyping. Recognises when to refer the person who is pregnant to a tertiary centre and how best to share care and monitoring. Liaises with the tertiary centre and the MDT to manage pregnant people with fetal anomalies. Formulates, implements and, where appropriate,
	 modifies management plan, in collaboration with subspecialists. Counsels pregnant people and their partners about the fetal risks, implications for the pregnancy and the long-term outcome. Signposts pregnant people to external sources of information and support. Constructs a follow-up plan for the pregnancy to support the pregnant person and plan next steps. Plans birth and appropriate neonatal support with a fetal medicine specialist.
Counsels and manages pregnancies at risk of fetal infection	 Investigates common fetal infections. Works with virology to interpret laboratory results for each infection. Explains the potential long-term effects of fetal infections on fetuses and newborns. Recognises when to involve other specialists in the care of a pregnant person with a suspected or confirmed fetal infection and plans for the sharing of care and monitoring. Liaises appropriately with the tertiary centre and the MDT to manage fetal infection.
Counsels and manages severe early fetal growth restriction (FGR)	 Is able to produce a differential diagnosis for severe early FGR. Knows when and which further investigations should be offered for severe early FGR. Liaises with the fetal medicine tertiary referral centre about diagnosis of severe early FGR and to manage it.



Counsels pregnant person about prenatal investigations

- Understands both the non-invasive and invasive options and is able to discuss the risks and benefits, facilitating choice.
- Understands the different levels of resolution of genetic testing and can communicate the importance of this to parents.
- Explains the risks and benefits of each procedure to the pregnant person and any alternatives.
- Communicates the scope and limitations of these tests.
- Describes how prenatal samples are processed and when, and how, the results are given.
- Offers genetic counselling where appropriate.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- NOTSS
- TO2
- CbD
- Mini-CEX
- **RCOG Learning**
- Local and deanery teaching
- Attendance at relevant courses and conferences

- Attendance at clinical genetics clinics
- Log of cases and outcomes
- Attendance at fetal medicine clinics
- Attendance at MDT meetings
- Attendance at specialist neonatal and paediatric clinics
- Examples of anonymised birth plans

Mandatory requirements

No mandatory evidence

- The genetic basis for trisomy 21, 18 and 13 and the ultrasound features associated with them
- The range of tests available for screening and testing for the common fetal trisomies and the organisation and quality control of the screening service
- Other aneuploidies: the implications of Turner syndrome (45,XO), Klinefelter syndrome (47,XXY) and Triple X syndrome (47,XXX) and appreciate the approach to managing pregnancies complicated by much rarer and unique chromosomal anomalies
- The underlying genetic inheritance patterns and prenatal testing for cystic fibrosis, muscular dystrophy and Fragile X syndrome, and the need for liaison with clinical genetics
- When it is appropriate to offer invasive testing, and when not to
- The role of non-invasive testing
- The implications for the current pregnancy and the long-term prognosis for each condition, and recurrence risks for future pregnancies
- The limitations of ultrasound in detecting and diagnosing congenital anomalies (e.g. cleft palate) or predicting prognosis (e.g. diaphragmatic hernia)
- Triggers and diagnoses that need to be referred to tertiary services



- Diagnostic features of each condition, their differential diagnosis and the chance of associated structural, chromosomal and syndromic associations
- The role of DNA analysis from maternal plasma

Prenatal Diagnosis CiP 3: The doctor demonstrates the skills and attributes required to provide ongoing support and care to people who have had a problem identified with their pregnancy.

Key skills	Descriptors
Counsels on and organises, or refers onwards for, termination of pregnancy for fetal anomaly	 Raises the option of termination of pregnancy for fetal anomaly appropriately and sensitively. Counsels pregnant person about the different methods of termination, explaining when termination is offered and when feticide is legally mandated. Organises termination of pregnancy for fetal anomaly (or refers appropriately where there is conscientious objection or the need for tertiary involvement). Supports the parent journey from diagnosis to follow up with planning for future pregnancies. Adjusts care around termination of pregnancy in high-risk situations. Manages complications of termination of pregnancy. Is aware of and can signpost to appropriate organisations that provide support.
Supports a pregnant person who wants to continue with their pregnancy where the fetus will not survive to birth, or the baby is expected to die in the neonatal period	 Supports and empowers the parent or parents in their decision. Plans for delivery with the parent or parents and paediatric team to give them the best experience possible in the circumstances, with clarity on intervention and non-intervention in labour. Plans an appropriate end of life pathway with the family and paediatric team.
Provides follow up and counselling after a pregnancy complicated by fetal anomaly	 Explains the role of the post-mortem and any other relevant post-birth tests (e.g. genetic testing, post-mortem MRI). Explains the findings and implications of any additional post-birth investigations. Refers, where appropriate, to the wider MDT, including clinical genetics. Counsels the parent or parents about the chance of recurrence across the range of conditions targeted by



FASP, and arranges genetic counselling where
appropriate.

- Proposes a plan to manage future pregnancies.
- Recognises when tertiary service involvement is appropriate for more complex cases.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- NOTSS
- TO2
- CbD
- Mini-CEX
- RCOG Learning
- Local and deanery teaching
- Attendance at relevant courses and conferences

- Attendance at neonatal unit ward rounds
- Log of cases and outcomes
- Attendance at fetal medicine clinics
- Attendance at MDT meetings
- Attendance at specialist neonatal and paediatric clinics
- Examples of anonymised birth plans

Mandatory requirements

No mandatory evidence

Knowledge criteria

- The antenatal management, intrapartum care and immediate postnatal management of each condition
- The impact of the diagnosis and individual circumstances on the timing, location and mode of birth
- The local prenatal, birth and post-birth pathways for care of the fetus and newborn with these conditions
- The legal framework under which termination of pregnancy by feticide may be offered
- Recognise which conditions are amenable to prenatal treatment (e.g. diaphragmatic hernia and spina bifida)
- The recurrence risk and management plan for future pregnancies for each condition

3.5 MFM CiPs 1–3

SST MFM CiP 1: The doctor is able to lead in providing care to women with pregnancies complicated by the full range of fetal concerns.

Key skills	Descriptors
Manages rare fetal structural anomaly	 Diagnoses, provides a differential diagnosis for, and manages the full range of rare fetal structural anomaly. Demonstrates how ultrasound findings of rare fetal structural anomalies are researched and managed.



	 Can counsel women and their partners about fetal risks, implications for the pregnancy and long term outcomes. Offers other prenatal test as appropriate, and involves Clinical Genetics team. Liaises appropriately with the referring centre and the multidisciplinary team (MDT). In collaboration with paediatric specialists, formulates, implements and, where appropriate, modifies a management plan. Specialists include paediatric cardiology, urology, neurology and members of the surgical team. Signposts to external sources of information and support. Constructs a follow-up plan for the pregnancy. Plans birth and appropriate neonatal support for someone who is pregnant and has concerns about their baby. Formulates a management plan for future pregnancies.
Manages fetal hydrops	 Constructs a differential diagnosis and uses
ivialiages retai ffydrops	appropriate investigations.
	 Treats reversible causes.
	Manages pregnancies where the cause of the
	hydrops remains unclear.
	Pursues the diagnosis post-birth and provides
	counselling for future pregnancies.
Manages rare complications of	Diagnoses and manages twin-to-twin transfusion (TTTC) and the fall and fall are fall as a fall as
multiple gestations	syndrome (TTTS) and provides follow-up care.
	 Manages discordant anomaly, including counselling on the selective termination of
	pregnancy.
	 Recognises and manages twin reversed arterial
	perfusion (TRAP) sequence.
	Refers to quaternary services for high level
	procedures, where indicated.
	Manages monoamniotic twin pregnancies.Manages triplet and higher order multiple
	Manages triplet and higher order multiple gestations, including providing counselling,
	without judgement, on multifetal pregnancy
	reduction.
	Diagnoses and manages severe early onset
	selective fetal growth restriction in monochorionic



	and dichorionic multiple pregnancies.
Manages pregnancies with a high chance of fetal alloimmune disorders	 Explains the potential fetal and maternal risks of red cell antibodies. Provides surveillance for pregnancies complicated by Parvovirus infections. Liaises with blood transfusion and neonatal services. Classifies risks for any pregnancy complicated by red cell antibodies and provides appropriate surveillance for fetal anaemia. Prepares women and their partners for the neonatal care necessary in cases of haemolytic disease of the fetus and newborn (HDFN). Explains the risks of maternal antiplatelet antibodies and knows when they should be tested for. Manages a pregnancy complicated by maternal antiplatelet antibodies, including birth and neonatal care, and offers antenatal treatment, where appropriate.
Offers and provides termination of pregnancy at all gestations appropriately	 Raises the option of termination of pregnancy for fetal anomaly appropriately. Can counsel a pregnant person about the different methods of termination. Organises termination of pregnancy for fetal anomaly. Adjusts care around termination of pregnancy in high-risk situations. Manages complications of termination of pregnancy.
Manages high level procedural skills	 Can counsel on, and take consent for, high-level interventional procedures.
Is able to support non-subspecialist colleagues to manage pregnancies complicated by fetal problems	 Provides subspecialist advice to non-subspecialist colleagues. Works in partnership with referring clinicians to provide joint care. es of evidence (not mandatory requirements)



- Reflective practice
- TO2 (includes SO)
- Mini-CEX (to include feticide).
- Procedural log
- CbD
- RCOG and other learning
- Attendance at regional national meetings and training courses
- Observation of, and reflection on, high-level fetal procedures
- Observation of neonatal surgery
- Attendance at local and regional MDT meetings
- Clinical attachments on tertiary level NNU and/or paediatric ITU
- Attendance at paediatric follow up clinics
- Relevant audit and quality improvement project

Mandatory requirements

- **OSATS**
 - CVS
 - o Amniocentesis
 - fetal echocardiography (ECHO)
 - anomaly scan
 - cervical length scan
 - o therapeutic amniodrainage
 - feticide

- Embryology of all key fetal anatomical systems
- Pathology and epidemiology of all major anomalies affecting each fetal system in addition to those covered in CiPs 2 and 3, including, as a minimum:
 - o encephalocele, holoprosencephaly, microcephaly and intracranial mass
 - o cardiac tumours
 - o renal cystic disease, duplex kidney and bladder/cloacal exstrophy
 - o laryngeal/tracheal atresia, pulmonary sequestration and pleural effusion
 - o meconium ileus, hepatic calcification/mass, abdominal cyst and ascites
 - o cystic hygroma, micrognathia, macroglossia, anophthalmia and neck mass
 - o skeletal dysplasias (early or late onset), polydactyly, sirenomelia, sacral agenesis and hemivertebra
 - o fetal akinesia/hypokinesia sequence
 - sacrococcygeal teratoma
- Diagnostic features of each condition, their differential diagnosis and the chance of associated structural, chromosomal and syndromic associations in the fetus
- Outcomes, prognoses and recurrence risks associated with each of these conditions and abnormalities
- Antenatal management, intrapartum care and immediate postnatal management of each condition
- Conditions that are responsive to prenatal therapy (e.g. fetal arrhythmias, spina bifida and congenital diaphragmatic hernia) and how these treatments are administered and the complications of them.



- The additional information which might be gained by use of 3D imaging and/or fetal MRI
- The differential diagnosis for fetal hydrops, and how to address this systematically
- The differential diagnosis for fetal anaemia
- Which red cell antibodies carry the greatest chance of haemolytic disease of the fetus and newborn, what thresholds there are for commencing surveillance for fetal anaemia, when to refer for fetal blood sampling and transfusion, how this is performed, and how the newborn is managed when there is a chance of haemolytic disease
- How platelet antibody-antigen combinations commonly cause neonatal alloimmune thrombocytopenia and what the outcomes can be, and how the chance of harm can be reduced.
- The embryology of normal twinning and the incidence and pathogenesis of abnormal twinning, resulting in TTTS, selective fetal growth restriction (sFGR), TRAP sequence and conjoined twins.
- When treatment is indicated for these conditions, and the pros and cons of treatment options
- A differential diagnosis for s-FGR and the classification of s-FRG in monochorionic gestations, and the impact that chorionicity has on outcomes and interpreting surveillance
- The differential risks associated with co-twin death in monochorionic and dichorionic multifetal gestations
- The outcomes of higher-order pregnancies, and the impact on these of multifetal pregnancy reduction
- The techniques used for selective termination of pregnancy for discordant anomalies in multiple gestations, and the risks involved
- UK law on termination of pregnancy, including justifying criteria, gestational limits and when to perform feticide, and ethical issues around late amniocentesis/late termination of pregnancy
- The significance of signs of life following a termination
- The various methods of termination of pregnancy, and the pros and cons of each method
- The indications, methods, potential benefits and complications of the following highlevel fetal medicine procedures: vesicocentesis, pleural and vesical shunt placement, placental laser, radiofrequency ablation, cord occlusion and fetal blood transfusion
- The structure of the local paediatric network, including surgical services
- Paediatric network guidelines for the management of newborn problems, including frameworks around extreme prematurity, and antenatal parallel planning

SST MFM CiP 2: The doctor can independently manage pregnancies complicated by the widest range, and most complex, of maternal medicine conditions, alongside specialists from other disciplines, contributing to the design and leadership of the maternal medicine



subspecialty service (in England this will be through the Maternal Medicine Networks either through MDTs for category B medical problems or referral for ongoing care to a Maternal Medicine Centre for category C medical problems).

Key skills	Descriptors
Manages the care of the pregnant woman presenting with any coexisting medical problem, including those with rare disorders and severe manifestations or complications of more common problems	 Extends further the key skills described in the Pregnancy Care and Maternal Medicine SITMs to provide pre-pregnancy counselling, antenatal, intrapartum and postnatal care to women with highly complex medical problems (e.g. category C in England). Can lead on the care provision of a pregnant person who is receiving joint care from a non-subspecialist and tertiary level team. Provides constructive advice to the non-subspecialist obstetrician and physician.
Lead MDT meetings	 Demonstrates the ability to work with other disciplines to make sure women receive the best possible care for their medical conditions, despite their pregnancy status. Understands and advises the MDT on the risks and benefits of different methodologies used in imaging and therapeutics. Understands, advises and signposts the MDT about prescribing in pregnancy.
Provides regional clinical leadership	 Leads on the coproduction of guidelines and standards which aim to best identify, refer and manage pregnant women, or those who wish to become pregnant, with co-existing medical conditions, and those who develop medical illness during their pregnancy. Designs and contributes to the monitoring of the above standards. Coordinates and contributes to the education and training of the multi-disciplinary team working within the Maternal Medicine Network/regional service. Makes sure everyone receives equal access to specialised care. Recognises the increased vulnerability of women from ethnic minorities and socially deprived groups in the design of systems.



Engages with other stakeholders

- Works with relevant networks e.g. maternity and perinatal mental health networks, fetal medicine services and neonatal operational delivery networks.
- Works with high level management organisations and bodies such as local maternity services, integrated care systems, NHSE/I, NHS Education for Scotland (NES), Health Education and Improvement Wales (HEIW), Northern Ireland Medical and Dental Training Agency (NIMDTA) and maternity strategic clinical networks.
- Liaises and works in partnership with other contributors to the care pathway e.g. ambulance trusts, GPs and health visitors.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- CbD
- Mini-CEX
- Reflective practice
- TO2 (includes SO)

- **RCOG Learning**
- Anonymised examples of pregnancy care plans for women with medical disorders
- Attendance at specialist courses and conferences
- Attendance at adult medical clinics
- Attendance at obstetric anaesthetic clinics
- Local and deanery teaching
- Attendance and leading of Maternal Medicine **MDTs**

Mandatory requirements

No mandatory evidence

Knowledge criteria

- Structure of local maternal medicine networks and regional thresholds for referral and MDT involvement
- The pathology, prevalence, presentation and diagnosis of, and risks and best practice management for, women who have significant medical problems (pre-pregnancy, antenatal and postnatal) that predate or arise in pregnancy or in the puerperium. Examples of category C medical problems outlined in the National Maternal Medicine Service Specification include:
 - Heart: pulmonary hypertension, severe left ventricular ejection fraction <45%, complex congenital heart disease (univentricular system, including Fontan, severe aortic and mitral valve stenosis and mechanical valve), aortic dilation, ventricular arrhythmias, new ischaemic heart disease and heart transplant.



- Lung: sickle chest crisis, restrictive lung disease with forced vital capacity <50%, neuromuscular disorder with respiratory muscle involvement e.g. myasthenia gravis, Guillain-Barre syndrome, lung transplant and pulmonary vasculitis
- Gastrointestinal and liver: portal hypertension, complex pancreatitis, active malignancy, cirrhosis, decompensated liver disease and liver transplant
- Endocrine: primary and secondary hyperaldosteronism, phaeochromocytoma, Cushing's syndrome, acromegaly, metabolic disorder (e.g. glycogen storage disorder) and hypo- and hyperparathyroidism
- Kidney: active lupus nephritis, new renal vasculitis, pre-pregnancy chronic kidney disease stage 4 and 5, renal dialysis, simultaneous renal and pancreatic transplant
- o Rheumatological: large and medium vasculitis and vascular Ehlers-Danlos syndrome
- Neurological: All epilepsy without local access to a combined clinic, including specialist neurology and obstetrics, symptomatic raised intracranial pressure, unstable congenital vascular malformation (CVM)/arteriovenous malformation/AVM, recent intracerebral haemorrhage(<2 years), acute stroke, new onset Guillain-Barre syndrome, new diagnosis or recent exacerbation of myasthenia gravis, active central nervous system malignancy and myotonic dystrophy
- Haematological: complex thalassaemia, current extensive venous thromboembolism, active haematological malignancy, clotting deficiency (Factor II, X and combined deficiency), Von Willebrand disease type II and III, carriers of haemophilia with male or unknown gender of fetus (including knowledge of when to offer late amniocentesis, and intrapartum management of such cases), transfusion dependent disease, antiphospholipid syndrome with extensive arterial events, antithrombin deficiency and moderate/severe platelet function disorder or with platelet count <100
- Findings of relevant national reports including MBRRACE report and recommendations
- How pregnancy induces significant changes in all aspects of physiology in pregnancy and the postpartum period. Plus the effect that pregnancy has on additional medical conditions
- How the medical problem may deteriorate acutely during pregnancy, how this might present and how it would be managed

SST MFM CiP 3: The doctor can apply their knowledge of clinical and molecular genetics to manage complex pregnancy.							
Key skills	Descriptors						
Manages a pregnancy at elevated chance of, or affected by, aneuploidy	 Takes an appropriate history and arranges appropriate parental investigations. Communicates effectively with women and their partners/families about risk, screening and testing options. Manages the care of a woman with a personal or family history of a chromosomal anomaly, including assessment of risk, prenatal diagnostic 						



	options, and further management options after
	testing.
	 Manages an ongoing aneuploid pregnancy, including plans for birth and a multidisciplinary approach to the care of the newborn. Recognises when advice from, and referral to, clinical genetics services is needed.
Manages a pregnancy with a chance of a single gene disorder in a structurally normal fetus	 Takes an appropriate history, constructs a family tree and arranges appropriate parental investigations. Communicates effectively with women and their partners/families about risk, screening and testing options. Manages the care of a woman with a personal or family history of a single gene disorder, including assessment of risk, prenatal diagnostic options, and further management options, after testing. Manages an ongoing pregnancy affected by a single gene disorder, including communication and planning with paediatric services. Recognises when advice from, and referral to, clinical genetics services is needed.
Diagnoses and manages genetic and syndromic disorders in a fetus that has a structural anomaly	 Can appropriately counsel and manage in families with a previous child with multiple anomalies or syndromic disorder. Accesses online information, of the highest quality, regarding very rare syndromic and genetic problems. Manages the care of a woman with a personal or family history of a syndromic anomaly, providing information, screening and prenatal testing options. Uses a dysmorphology database to reach a differential diagnosis. Recognises when referral is indicated for more specialised counselling and genetic advice. Provides options to manage an affected pregnancy, including termination of pregnancy, without judgement. Manages an ongoing pregnancy, including planning for birth and a multidisciplinary approach to the care of the newborn.



Requests and uses a wide range of molecular, cytogenetic and biochemical tests for prenatal diagnosis

- Is able to take non-directive informed consent for performing these tests.
- Is able to interpret and communicate the results of these tests and knows when a multidisciplinary approach is required.
- Identifies the role of genomic testing in both maternal and fetal medicine, in pregnancy, or ideally in pre-conception counselling
 - Maternal Medicine Expanding research towards genetic prediction tests of obstetric disease processes, such as pre-eclampsia, gestational diabetes, obstetric cholestasis
 - o Role of genomics in preconception care for women with rare genetic disease and pregnancy care for women with rare genetic disease, especially in affected families seeking IVF, planning surrogacy, or carrier screening.
 - Role of whole genome testing vs whole exome sequencing.

Evidence to inform decision – examples of evidence (not mandatory requirements)

- Reflective practice
- Local and deanery teaching
- TO2 (includes SO)
- Mini-CEX
- CbD

- NOTSS
- RCOG Learning

Mandatory requirements

No mandatory evidence

Knowledge criteria

- Normal chromosome structure and function
- Gene structure and function, including gene control, mechanisms and effects of mutation, and genetic heterogeneity
- Patterns of genetic inheritance and susceptibility, gene expression and penetrance, multifactorial and mitochondrial inheritance
- Cell division (meiosis and mitosis), and abnormalities arising from these processes
- Types of aneuploidy, including structural rearrangements, deletions and common microdeletions, trisomies, sex chromosome anomalies (including monosomy X, Klinefelter syndrome and Triple X), extra markers, mosaicism (fetal and placental), uniparental disomy and triploidy
- The underlying genetic aetiology of the single gene disorders mentioned in CiPs 2 and 3, and the following conditions:
 - myotonic dystrophy
 - Huntington's disease
 - haemoglobinopathies, haemophilia and other common bleeding disorders



- inborn errors of metabolism
- Detailed knowledge of the following syndromes and associations:
 - DiGeorge
 - o Fryns
 - Beckwith-Wiedemann
 - Meckel-Gruber
 - Smith-Lemli-Opitz
 - VATER/VACTERL association
- The pre- and postnatal phenotypes of these common aneuploidies, single gene disorders and syndromes, including prognosis
- Methods of screening for aneuploidy, including ultrasound, biochemical and noninvasive DNA based techniques
- The statistical terms relevant to screening, including sensitivity, specificity, false positive rates, positive predictive rates, and how these are interdependent
- The meaning of likelihood ratios in risk calculations
- Current screening programmes, including national implementation, audit, quality control, the UK National Screening Committee and regional screening coordinators
- How recurrence risks for chromosomal and single gene disorders are derived
- Prenatal testing options, both invasive and non-invasive, including ultrasound, MRI, noninvasive prenatal testing, amniocentesis, CVS and fetal blood sampling.
- Laboratory techniques for analysing parental and fetal samples, including quantitative polymerase chain reaction, fluorescence in situ hybridization, karyotyping, microarray, mutational analysis, sequencing (exome, or whole genome), enzymatic analysis and analyte assessment.

3.6 The SSTR CiP

The previous requirement for 'research accreditation' (evidenced by a higher degree, the Research Advanced Professional Module or two relevant first author papers) has been removed from the 2024 SST curriculum. In recognition of the important contribution made by, and expected of, most subspecialists, this requirement has been replaced with a subspecialty-specific research CiP. Trainees who have already been involved in research are likely to be able to use evidence of these research skills to support sign-off of this CiP.

SSTR CiP: The doctor is able to engage with research and promote innovation within their subspecialty.

Key skills Descriptors

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Demonstrates research skills	 Is able to demonstrate practice in healthcare research and the different methodologies within their subspecialty. Shows continued engagement in Good Clinical Practice (GCP) and Research and Development (R&D) processes. Engages in ethics and governance processes within research, demonstrating they are able to follow guidelines on ethical conduct and consent for research. Demonstrates involvement in informatics, statistical analysis and emerging research areas within their subspecialty. Shows engagement with national trials within their subspecialty, including patient recruitment, trial monitoring and adverse event reporting. Shows understanding of the role of public and patient involvement within clinical trials. Is able to discuss clinical trials with, and facilitate recruitment of, patients within their subspecialty. Has the ability to translate research into clinical
Demonstrates critical thinking Innovates	 Is able to develop and critically appraise a research protocol. Is able to critically evaluate clinical trial data to establish the clinically significant outcomes and relevance for clinical practice within their subspecialty. Is able to interpret research findings, reflect on the potential impact on their clinical practice and share this with colleagues and patients. Can develop and critically appraise a patient information leaflet. Is able to interpret research findings within their subspecialty and discuss these when taking informed consent for treatment. Demonstrates how their clinical practice has developed from innovative research within their subspecialty. Is able to demonstrate engagement with the introduction of any innovations within their subspecialty, including governance and costs.



- National teaching and courses
- Critical appraisal of protocols/papers
- Subspecialty journal club presentations
- GCP re-certification
- Participation, including recruitment for national multicentre trials
- Preparation of research protocol/grant applications
- Oral, and/or poster presentations at national/international subspecialty meetings

- SIPM in Clinical Research
- Peer reviewed original research publications relevant to their subspecialty
- A higher degree such as a PhD or research MD

Mandatory requirements

No mandatory evidence

4. GMC Generic Professional Capabilities (GMCs)

The key skills in all 16 CiPs also map to a variety of generic professional capabilities (GPCs). When providing evidence of their progress in this programme, learners should make sure that it also displays progress/capability in the GMC GPCs, such as dealing with complexity, teamwork and leadership, and knowledge of patient safety issues.

Mapping to the GPCs

Domain 1: Professional values and behaviours

Domain 2: Professional skills

Domain 3: Professional knowledge

Domain 4: Capabilities in health promotion and illness prevention

Domain 5: Capabilities in leadership and team-working

Domain 6: Capabilities in patient safety and quality improvement

Domain 7: Capabilities in safeguarding vulnerable groups

Domain 8: Capabilities in education and training

Domain 9: Capabilities in research and scholarship



Learners can expect to be assessed on their wider skills as a medical professional, their skills in leadership and teamwork, and their level of clinical competence. Evidence showing progress in these areas will result in the learner progressing through the programme.

To help learners and Educational Supervisors determine what acceptable progress looks like, there is a Statement of Expectations for each of the 16 CiPs.

	Statement of Expectations for the MFM SST
Meeting	Learners can risk stratify women with pre-existing medical conditions,
expectations	determining the most appropriate level of obstetric medical care. They are
for the	able to manage the care of most women with hypertension, common
Pregnancy Care	endocrine disorders, diabetes and extremes of body mass index. They will
CiP1	be confident in the initial assessment and diagnostic pathway of women
	presenting with new onset health concerns.
Meeting	Learners can manage appropriately, in conjunction with associated
expectations	specialists, the care of pregnant people with pre-existing or newly
for the	acquired viral infections that may have affect pregnancy outcome.
Pregnancy Care	
CiP2	
Meeting	Learners can advise women with common medical problems on the best
expectations	contraceptive options, long-term health implications and impact of their
for the	medical condition on future pregnancies (and how to minimise this).
Pregnancy Care	
CiP3	
Meeting	Learners consider the implications of cultural, religious and social factors
expectations	on pregnancy, and work with associated health and social care specialists
for the	to optimise outcomes.
Pregnancy Care	
CiP4	
Meeting	Learners can work with other professionals within their own and other
expectations	specialties to optimise the care and well-being of the pregnant person
for the	with medical conditions, assessing them through focused history taking,
Maternal	examination and choice of investigations. They can work with pregnant
Medicine CiP1	people when considering the risk-benefit analysis of possible treatment
	and management options. They are also able to give effective advice and
	pregnancy plans to people who are considering pregnancy and have pre-
	existing medical conditions.
Meeting	Learners are able to consider the impact of pre-existing disease on
expectations	pregnancy, and pregnancy on pre-existing disease. They can plan with a
For the	pregnant person with specific medical conditions, to optimise their care
Maternal	and birth experience. They are also competent in evaluating the pregnant
Medicine CiP2	person who presents with a new condition during pregnancy.
Meeting	Learners can use ultrasound to confirm normal and abnormal fetal growth
expectations	and wellbeing. Learners can communicate with their patients effectively,



for the Fetal	discuss their findings and plan care. They will engage with a
Care CiP1	multidisciplinary network, and recognise when referral is required to a
	subspecialist.
Meeting	Learners can discuss and plan management with a pregnant person whose
expectations	fetus is at risk from red cell alloimmunisation. They are also able to assess
for the Fetal	the likelihood of fetal anaemia, and liaise with tertiary units when
Care CiP2	intervention is indicated.
Meeting	Learners can provide care for pregnant people with mono- or dichorionic
expectations	twin pregnancies, using ultrasound assessment to monitor growth and
for the Fetal	wellbeing. Learners can screen for twin-to-twin transfusion syndrome and
Care CiP3	other complications of monochorionic twinning. They can also plan
Care CIPS	· · · · · · · · · · · · · · · · · · ·
	effective care with a pregnant person with a multiple pregnancy, and in
Direction	more complex cases, liaise with, and refer to, subspecialty centres.
Meeting	Learners are able to complete a fetal anomaly scan and confirm normal
expectations	views.
for the Prenatal	Learners can recognise when fetal anatomy deviates from normal.
Diagnosis CiP1	
Meeting	Learners understand genetic inheritance and can apply this knowledge to
expectations	counselling of parents where an anomaly has been found. Learners can
for the Prenatal	discuss abnormalities and prepare parents for the next steps in diagnosis.
Diagnosis CiP2	Learners understand the implications of abnormal findings on the anomaly
	scan, and can recognise significant fetal infections and initiate
	management or referral. Learners are able to monitor and intervene
	appropriately in severe early onset growth restriction.
	Learners use a good working knowledge of other tests that may be offered
	to parents where there is a risk of prenatal problem, or where an anomaly
	has been found.
Meeting	Learners are able to continue to provide high-quality care after a diagnosis
expectations	has been reached, including providing counselling about termination of
for the Prenatal	pregnancy.
Diagnosis CiP3	Learners are able to continue to provide high-quality care for parents who
	continue with their pregnancy after an anomaly has been found, can
	support parents after an affected pregnancy and can provide information
	to help them make decisions for the future.
Meeting	Learners are able to investigate and manage pregnancies complicated by
expectations	the full range of fetal structural and genetic differences, however rare and
for the SST	complex, fetal hydrops, pregnancies at high risk of haemolytic disease, and
MFM CiP1	multiple gestations complicated by anomaly and other issues.
	Learners can work effectively with the multidisciplinary team, including
	the referral source, to provide high-quality information, supported choice,
	optimal outcomes for ongoing pregnancies and follow-up care to people
	who have experienced pregnancy loss.
Meeting	Learners are able to lead on the care and management of individual
expectations	pregnant people with the most complex of co-existing medical conditions,
expectations	pregnant people with the most complex of co-existing medical conditions,



for the SST	while working effectively with the multidisciplinary team, including the
MFM CiP2	referral source.
	Learners will contribute locally, regionally and nationally to pathways,
	audits and policies designed to optimise the outcomes for this group of
	pregnant people.
Meeting	Learners will understand the complexity of single gene, chromosomal,
expectations	epigenetic and syndromic disorders, and know what investigations can be
for the SST	performed to determine underlying genetic causes. Working with the
MFM CiP3	multidisciplinary team, they are able to counsel on prognosis and
	recurrence risk.
Meeting	Learners have knowledge, understanding and practical experience of
expectations	research skills pertinent to their subspecialty.
for the SSTR CiP	Learners can demonstrate ongoing engagement with research in their
	subspecialty field, are able to critically appraise their own research
	findings and those of others, and can translate innovation into everyday
	subspecialty practice.

The CiP knowledge criteria show the processes/frameworks a learner should understand and the clinical knowledge they must have if they want to become a subspecialist in maternal and fetal medicine. This is more in-depth than the knowledge base expected for the MRCOG. The key skills and descriptors outline the expected learning outcomes for the MFM SST. However, learners will not experience the entire range of possible scenarios during their training; therefore, after completing the programme they should continue their learning and skill development through their independent practice as a maternal and fetal medicine subspecialist and at MDT meetings.

5. Procedures associated with the clinical **MFM CiPs**

The procedures required to complete this SST are listed below. A learner can show progress in these procedures through OSATS, procedure logs, attendance at courses and other forms of evidence.

Each procedure is assigned a supervision, or entrustability, level (defined in Section 6.4 of the Definitive Document for Maternal and Fetal Medicine SST 2024) recognising that acquisition of safe independent practice in some of the more complex and uncommon procedures may only be achieved as a consultant, working with more experienced colleagues. Level 5 indicates that a learner should be able to perform the procedure independently.



Procedures marked with * must be evidenced with three competent summative OSATS. The others can be evidenced using summative OSATS and other means (e.g. CbD, Mini-CEX, formative OSATS, reflections).

Some of these procedures also feature in a similar table in the Fetal Care and Prenatal Diagnosis SITM curricula guides, but the same supervision level is not necessarily required for completion of the SITM.

Procedures	Level by end of training	Fetal Care CiP1	Fetal Care CiP2	Fetal Care CiP3	Prenatal Diagnosis CiP1	Prenatal Diagnosis CiP2	Prenatal Diagnosis CiP3	SST CiP1
Fetal biometry and liquor volume*	5	Х						
Transvaginal placental localisation*	5	X						
Umbilical artery Doppler*	5	Х						
Middle cerebral artery Doppler*	5	Х	Х					
Ductus venosus Doppler*	5	Х						
Uterine artery Doppler*	5	Х						
Multiple gestation chorionicity*	5			X				
Twin pregnancy assessment*	5			Х				
Fetal ECHO*	5							Χ
Anomaly scan*	5							Χ
Cervical length scan*	5							Х
CVS*	5							Χ
Amniocentesis	5							Χ
Therapeutic amniodrainage*	5							Х
Feticide*	5							Χ
Twin amniocentesis	4							Х

Procedures	Level by end of training	Fetal Care CiP1	Fetal Care CiP2	Fetal Care CiP3	Prenatal Diagnosis CiP1	Prenatal Diagnosis CiP2	Prenatal Diagnosis CiP3	SST CiP1
Drainage of fetal cystic structure	2							X
MPFR/selective termination in dichorionic twins or higher order pregnancy	2							X
Fetal blood transfusion	1							Х
Fetal shunt Placental laser	1							X

OSATS are not assigned a level of entrustability, rather they are assessed as being competent or working toward competence. The entrustability levels here are given to guide the assessor in judging whether the learner has reached the required degree of independence at the end of training.

Evidence required

Discussion and detail on how trainees can evidence the acquisition of key skills and competencies during subspecialty training can be found in Section 4 of the Essential Curriculum Guide.

The table below may be useful for learners to see whether a specific workplace-based assessment can be used as evidence of progress in each CiP:

CiP	OSATS	Mini-CEX	CbD	NOTSS	TO1/TO2	Reflective practice
Pregnancy Care		Χ	Х	Х	Χ	Χ
CiP1: The doctor						
demonstrates the						
skills needed, and						
can apply their						
knowledge, to						
manage antenatal						
care for a pregnant						
person with						



CiP	OSATS	Mini-CEX	CbD	NOTSS	TO1/TO2	Reflective practice
common medical problems.						
Pregnancy Care CiP2: The doctor demonstrates the skills needed, and can apply their knowledge, to manage the care of a pregnant woman whose pregnancy is complicated by infection which may affect the health of her baby.		X	X	X	X	X
Pregnancy Care CiP3: The doctor demonstrates the skills needed, and can apply their knowledge, to manage the postnatal care of a pregnant person with common medical problems.		X	X	X	X	X
Pregnancy Care CiP4: The doctor provides holistic care to a pregnant person.		Х	Х	Х	X	Х
Maternal Medicine CiP1: The doctor is able to work with others to provide high quality care to		Х	Х	Х	Х	Х



CiP	OSATS	Mini-CEX	CbD	NOTSS	TO1/TO2	Reflective practice
the woman with medical conditions in pregnancy or planning a pregnancy.						
Maternal Medicine CiP2: The doctor has a high level of understanding of the impact that medical conditions have on pregnancy and is able to optimise care for the affected woman.		X	X	X	X	X
Fetal Care CiP1: Uses ultrasound skills to recognise, monitor and manage compromise to fetal wellbeing.		X	Х	X	X	X
Fetal Care CiP2: The doctor demonstrates the skills and attributes required to assess the fetus at risk of red cell alloimmunisation.	X	Х	X	X	X	X
Fetal Care CiP3: The doctor demonstrates the skills and attributes required to assess	X	X	X	X	X	X



CiP	OSATS	Mini-CEX	CbD	NOTSS	TO1/TO2	Reflective practice
complications of twin pregnancies.						
Prenatal Diagnosis CiP1: The doctor can use ultrasound to recognise where fetal anatomy is not normal.	Х	Х	X	Х	Х	X
Prenatal Diagnosis CiP2: The doctor can assess and investigate a pregnancy where there are concerns about the fetus.		X	X	X	Х	X
Prenatal Diagnosis CiP3: The doctor demonstrates the skills and attributes required to provide ongoing support and care to people who have had a problem identified with their pregnancy.		X	X	X	X	X
SST MFM CiP1: The doctor is able to lead in providing care to women with pregnancies complicated by the full range of fetal concerns.	X	X	X		X	X
SST MFM CiP2: The doctor can		Х	Х		Х	Х



CiP	OSATS	Mini-CEX	CbD	NOTSS	TO1/TO2	Reflective practice
independently manage pregnancies complicated by the widest range, and most complex, of maternal medicine conditions, alongside specialists from other disciplines, contributing to the design and leadership of the maternal medicine subspecialty service (in England this will be through the Maternal Medicine Networks either through MDTs for category B medical problems or referral for ongoing care to a Maternal Medicine Centre for category C medical problems).						
SST MFM CiP3: The doctor can apply their knowledge of clinical and molecular genetics to manage complex pregnancy.		X	X	X	X	X



6.1 Generic capabilities

All subspecialty doctors will need to provide evidence collected during subspecialty training for the following areas, at the centralised assessments:

- Clinical governance
- Teaching experience
- Research and innovation
- Leadership and management
- Presentations and publications.

This evidence should be uploaded into the 'Other evidence' section of the ePortfolio.

Pre-CCT subspecialty doctors on the Core Curriculum 2024 will be expected to meet the expectations of the core generic and non-clinical specialty CiPs at ST5/6/7 level. They can use their experiences during subspecialty training and emergency duties to help evidence these generic capabilities and skills. The evidence should be linked to the appropriate core generic and non-clinical specialty CiPs, and may need to be supplemented to satisfy their Educational Supervisors and Annual Review of Progression (ARCP) panels that they meet the full range of requirements at ST5/6/7 level.

For each core generic and non-clinical specialty CiP, there is a CiP guide outlining what the level of expectation is for senior doctors in ST5, ST6 and ST7.

CCT holders and overseas doctors undertaking subspecialty training do not need to complete the core generic and non-clinical specialty CiPs, although they can choose to link evidence of their generic skills into the core generic or non-clinical specialty CiPs on the ePortfolio after uploading this evidence into the 'other evidence' section of the ePortfolio.

Assessing progress

A trainee's progress follows the same principles of any other trainee in obstetrics and gynaecology, as detailed in Essential Curriculum Guide. However, the annual subspecialty assessment of progress is performed centrally, coordinated by the RCOG. Before this assessment, the subspecialty trainee and supervisor will need have performed assessments for each of the 16 CiPs, look at the evidence collected so far and give a global judgement on the trainee's progress. Together, they will construct the subspecialty-specific Educational Supervisor's Report (SST ESR), and this, alongside the evidence in the trainee's ePortfolio, will be reviewed by a subspecialty assessment panel. The panel will give a narrative outcome, stating if they judge the trainee to have successfully completed subspecialty training, to be making good progress or if they are behind schedule and may need additional focus or training time.



For pre-CCT subspecialty trainees, this narrative outcome is a major part of the trainee's evidence for their subsequent ARCP, which will also assess their progress through the Core Curriculum. Learners do need to appreciate that satisfactory progression through subspecialty training does not guarantee a satisfactory outcome (outcome 1) at the subsequent ARCP. For this reason, they will need to complete an ESR for their ARCP with their Educational Supervisor; this is separate from the SST ESR they created for their subspecialty assessment. The two different forms of ESRs are clearly marked and easily accessible on the front page of the learner or supervisor ePortfolio log-in for that learner. Learners need to ensure that they are also achieving any Training Matrix of Progression requirements for the Core Curriculum that are additional to those on the MFM subspecialty matrix.

For pre-CCT subspecialty trainees on the Core Curriculum 2024, the key additional areas to focus on are evidencing all of the core generic and non-clinical specialty CiPs to ST5/6/7 level, and sign-off of the core clinical CiPs (9–12) to entrustability level 5 by the completion of training and the final ARCP. All subspecialty learners using the Core Curriculum 2024 should collect evidence to satisfy all four core clinical CiPs to entrustability level 5, but DO NOT need to collect 'ongoing competency' OSATS for core procedures that they have already demonstrated competency in (with three competent summative OSAST), in line with the 2024 core matrix.

It is a GMC requirement that to achieve a CCT in Obstetrics and Gynaecology, training must be undertaken in both aspects of the specialty. Therefore, in addition to providing evidence for the core clinical CiPs 10 and 12, MFM pre-CCT subspecialty trainees also need to provide evidence for the gynaecology core CiPs 9 and 11. These CiPs relate to emergency and nonemergency gynaecology =

Information on the experience and evidence required for the gynaecology core CiPs 9 and 11 can be found in 'Guidance for Subspecialty Training Programme Supervisors and pre-CCT maternal and fetal medicine subspecialty trainees on Core Curriculum 2024 on cross specialty working'.

Career guidance 8.

Learners require two SITMs or subspecialty training for CCT. A learner aspiring to become a maternal and fetal medicine subspecialist should be advised to undertake any of the four SITMs included in the SST Maternal and Fetal Medicine:

- Pregnancy Care (foundation SITM for Maternal Medicine)
- Maternal Medicine (contingent SITM for Pregnancy Care)
- Fetal Care (foundation SITM for Prenatal Diagnosis) and/or
- Prenatal Diagnosis (contingent SITM for Fetal Care).

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This should make learners more competitive for the subspecialty interview, as the MFM SST builds on these SITMs. However, it is not mandated that these SITMs have been started, or completed, for a learner to be eligible for the MFM SST. Any completed CiPs of the Pregnancy Care, Maternal Medicine, Fetal Care or Prenatal Diagnosis SITMs, or all of the SITM(s), can be used for the MFM SST. However, learners' choices will be dependent on training opportunities available for their chosen SITMs.

Advice for learners who aspire to become a maternal and fetal medicine subspecialist is to start preparing for this direction as early as possible. They should have career conversations early in their training with their Educational Supervisor, find a mentor in the field of maternal and fetal medicine, complete the Clinical Research SIPM, achieve publications and start enquiring where posts may come up, as SST posts may be outside their region.

For further careers advice, learners should have a discussion with their Training Programme Director/SITM Director.

9. Further resources

The further resources listed below can be found on the RCOG Curriculum 2024 webpages:

- **Essential Curriculum Guide**
- Definitive Document for Maternal and Fetal Medicine SST 2024
- Curriculum Guide for Pregnancy Care SITM
- Curriculum Guide for Maternal Medicine SITM
- Curriculum Guide for Fetal Care SITM
- Curriculum Guide for Prenatal Diagnosis SITM
- Curriculum Guide for Clinical Research SIPM
- Training Matrix for SST MFM
- Definitive Document for Core Curriculum 2024 •
- Training Matrix of Progression
- British Maternal and Fetal Medicine Society (BMFMS)
- MacDonald Obstetric Medicine Society (MOMS)
- NHS Fetal Anomaly Screening Programme (FASP)

Find out more at rcog.org.uk/curriculum2024

