TITLE- GUIDELINE FOR USE OF PROGESTERONE THERAPY IN PREGNANCY



TARGET AUDIENCE	Secondary Care
PATIENT GROUP	Pregnant women with threatened miscarriage, previous recurrent miscarriage, previous preterm birth/2 nd trimester loss

Clinical Guidelines Summary

- Progesterone therapy has proven benefits in the treatment of the following conditions in pregnancy:
 - Threatened miscarriage and history of one previous miscarriage (5% improvement in live birth rates).
 - Threatened miscarriage and history of two previous miscarriages (5% improvement in live birth rates).
 - Threatened miscarriage and history of three or more previous miscarriages (15% improvement in live birth rates).
 - o Recurrent miscarriage without bleeding.
 - o Previous preterm birth/2nd trimester miscarriage and short cervix.
- For miscarriage prophylaxis for women fulfilling the criteria, treatment can begin from 6 weeks of gestation after a scan has confirmed the presence of an intrauterine pregnancy.
- For miscarriage prophylaxis, treatment is given in the form of micronised progesterone 400mg twice/day until 12 or 16 weeks of gestation.
- Form C for Unlicensed Medicine completed.
- Standard NHSL unlicensed medicine consent form and detailed consent to progesterone treatment to be completed by patient and prescriber.
- For preterm birth/2nd trimester miscarriage prophylaxis, treatment is given in the form of micronised progesterone 400mg twice/day until at least 34 weeks of gestation.



Guideline Body

This section will normally contain more detailed information to support summary and should have a **content** list to start with in his section.

Content

Introduction.

Use of progesterone in threatened miscarriage.

Use of progesterone in recurrent miscarriage.

Use of progesterone in prophylaxis for preterm birth/2nd trimester miscarriage.

References.

Introduction

There is some evidence that progesterone therapy may be of benefit for some women with complications of threatened miscarriage, recurrent miscarriage and prophylaxis against preterm birth or 2nd trimester miscarriage. This guideline outlines clinical situations where progesterone treatment may be of benefit and subsequent management of these patients.

Use of progesterone in threatened miscarriage.

- Telephone triage all women who present with bleeding in early pregnancy for supportive care or face-to-face review +/- ultrasound scan.
- If a scan is indicated, then arrange appointment in EPAS.
 - If the scan is inconclusive, manage according to Guideline for Inconclusive Scans.
 - If the scan confirms intrauterine pregnancy with fetal heartbeat, manage as follows:

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IU pregnancy with fetal heartbeat and vaginal bleeding

No previous history of miscarriage

- Reassure
- Return if bleeding worsens or persists >14 days for scan/assessment
- Routine antenatal care if bleeding stops

One or more previous miscarriages

- Offer all eligible women the Tommy's leaflet on progesterone treatment in early pregnancy and the NHSL consent form.
- Offer vaginal micronised progesterone 400mg twice/day until 12 or 16 weeks of gestation.
- Can be offered if there is confirmed ultrasound evidence of an intrauterine pregnancy with yolk sac prior to appearance of fetal heartbeat. In this circumstance offer further scan in 7 – 14 days to confirm viability.
- Complete consent forms (unlicensed medication and consent to progesterone) with patient decision on treatment and scan into Badger.
- There is evidence in the medical literature¹⁻⁴ for the use of progesterone in reducing the risk of miscarriage in women with early pregnancy bleeding and a history of miscarriage.
- Women with bleeding with any previous miscarriage using progesterone, increases the likelihood of a livebirth by 5%.
- Women with bleeding with three or more previous miscarriages using progesterone, increases the likelihood of a livebirth by 15%.
- There is no robust evidence to continue progesterone beyond 12weeks gestation though 16 weeks is the recommendation in NICE guideline. Duration of treatment to be confirmed with the woman during consent process.

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Use of progesterone in recurrent miscarriage.

- See also guideline on recurrent miscarriage.
- Progesterone can increase the livebirth rates for women who experience recurrent miscarriage^{1-3,5}.
- Consider micronised progesterone 400mg twice daily after ultrasound confirmed intrauterine pregnancy in women with bleeding and recurrent miscarriage. Start treatment at time of bleeding and continue until 12 or 16 weeks of gestation.
- Women with recurrent miscarriage and previously successful pregnancy using
 progesterone treatment, may attend requesting similar plan in a new pregnancy in
 the absence of bleeding. There is evidence that routine progesterone
 supplementation for women with recurrent miscarriage in the absence of bleeding
 may be of benefit but this is less robust. Progesterone therapy (micronised
 progesterone 400mg twice daily till 12 or 16 weeks), can be offered after discussion
 with the woman about the risks and benefits.
- Routine supplementation should be used with caution in asymptomatic women with unexplained recurrent miscarriage (>=3 previous miscarriages, previous successful pregnancy after miscarriage using progesterone). There is some evidence in the literature of benefit, but the Promise trial did not show any benefit. Women requesting this treatment should be referred for further discussion about the risks and benefits of treatment to the duty EPAS/Day care/Triage consultant.

Use of progesterone in prophylaxis for preterm birth/2nd trimester miscarriage.

- See also guideline on preterm birth.
- Woman with a history of 2nd trimester spontaneous loss or a history of spontaneous preterm birth up to 34 weeks of gestation, with transvaginal ultrasound evidence (between 16 and 24 weeks) of short cervix (<=25mm) should be offered the choice of prophylactic vaginal progesterone or prophylactic cervical cerclage⁶.
- Women choosing vaginal progesterone should be offered 400mg progesterone twice daily until at least 34 weeks of gestation.

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References/Evidence

Any content in your guideline that is either guoted, paraphrased and/or borrowed from an external source must be attributed to the original.

For published papers, Harvard referencing style is preferable

1. Quenby S et al. Miscarriage matters: the epidemiological, physical, psychological, and economic costs of early pregnancy loss. Lancet 2021; 397:



lancetrecommendat 1658-1667. ionsmiscarriage.pdf

- 2. Coomarasamy A et al. Sporadic miscarriage: evidence to provide effective care. Lancet 2021; 397: 1668-1674.
- 3. Coomarasamy A et al. Recurrent miscarriage: evidence to accelerate action. Lancet 2021; 397: 1675-1682.
- 4. NICE Guideline NG126, 2023. Ectopic pregnancy and miscarriage: diagnosis



and initial management.

nice miscarriage ectopic.pdf



GTG recurrent miscarriage.pdf

5. RCOG GTG17, 2023. Recurrent miscarriage.



nice preterm birth.pdf

6. NICE Guideline NG25, 2022. Preterm labour and birth.



Consent to

7. Consent to Progesterone Treatment.



Progesterone Pessaries Form-C---l

8. Form C for progesterone treatment.



Early pregnancy

9. Tommy's Patient Information Leaflet. bleeding and proge

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Appendices

1. Governance information for Guidance document

Lead Author(s):	Evelyn Ferguson
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Distribution	Maternity Clinical Effectiveness Group UHW Early Pregnancy Assessment Service Multi-Disciplinary
	Team
	Women and Neonatal Services DMT
	Consultant Obstetricians/Gynaecologists.
	Senior midwifery team.
	Head of Pharmacy
	Clinical Pharmacist Women and Children's Team

CHANGE RECORD

Date	Lead Author	Change	Version No.
		e.g. Review, revise and update of policy in line with	1
		contemporary professional structures and practice	
			2
			3
			4
			5

2. You can include additional appendices with complimentary information that doesn't fit into the main text of your guideline, but is crucial and supports its understanding.

e.g. supporting documents for implementation of guideline, patient information, specific monitoring requirements for secondary and primary care clinicians, dosing regimen/considerations according to weight and/or creatinine clearance

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