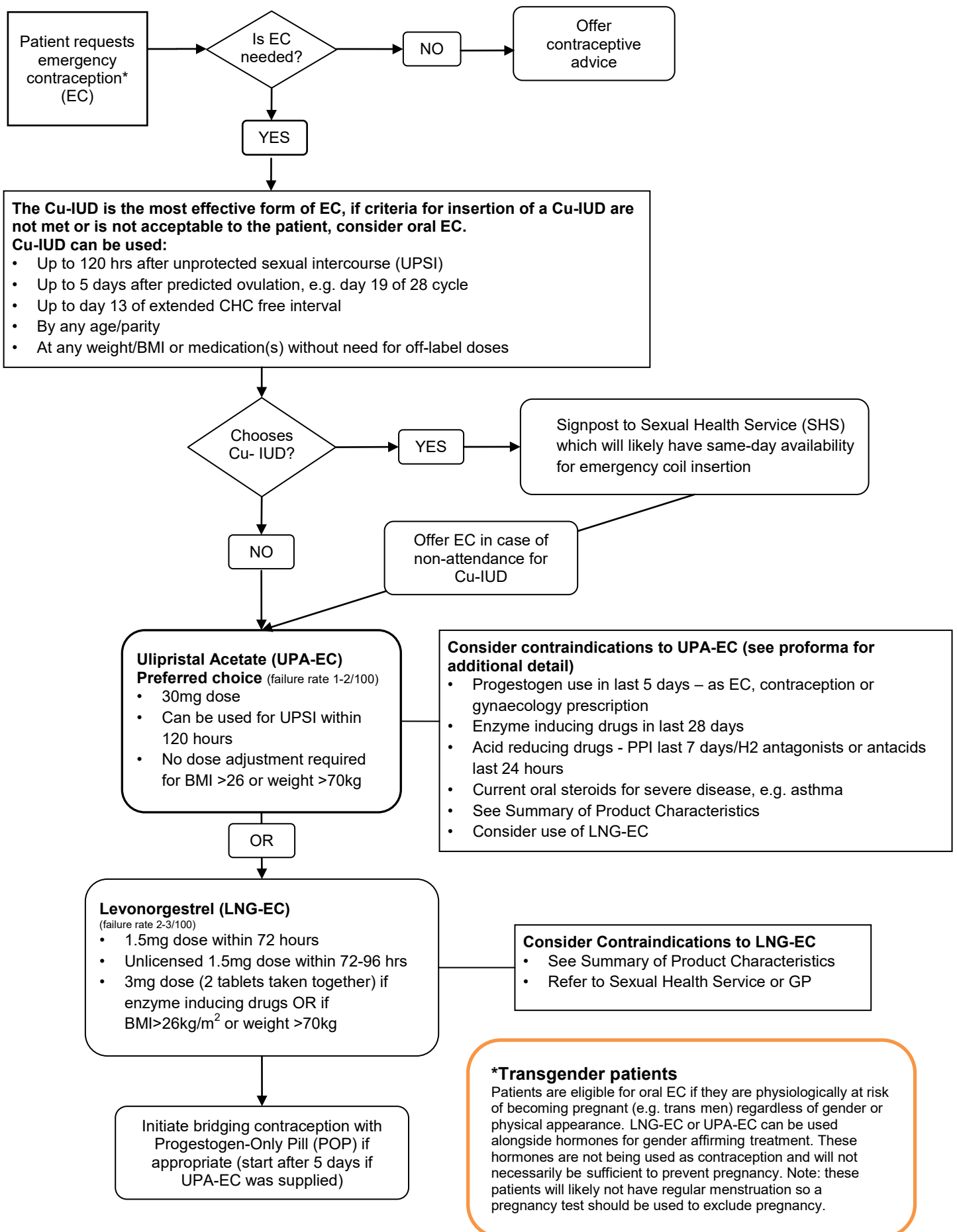


Flowchart for Oral Emergency Contraception (EC): Ulipristal Acetate (UPA-EC) versus Levonorgestrel (LNG-EC) **if** Cu-IUD is not appropriate or acceptable.

NB: Oral EC is unlikely to be effective if **taken after ovulation.**



Emergency Contraception Pro forma

Consultation Details

Healthcare Professional Name (PRINT):	Date of Consultation:	
Patient Name:	Date of Birth:	Age:
Patient aged 13 years or over and competent to consent? Yes <input type="checkbox"/> No <input type="checkbox"/>		
If No or child protection issues: follow local child protection guidance and refer to local service		

Circumstances Leading to EHC Request

UPSI
Time since UPSI? <input type="checkbox"/> <72 hrs <input type="checkbox"/> 72-96 hrs <input type="checkbox"/> 96-120 hrs <input type="checkbox"/> >120 hrs (may warrant referral to local SHS)

History		Action/Information
Day 1 of last menstrual period (LMP)	/ /	This allows calculation of place in the cycle. Oral EC after ovulation (days 9-16) can be given but is likely to be ineffective and a Cu-IUD should be used.
LMP regular? (See info for trans patients on flowchart)	Yes <input type="checkbox"/> No <input type="checkbox"/>	
Given birth within the last 3 weeks?	Yes <input type="checkbox"/> No <input type="checkbox"/>	If yes EC is not required. Note: Early pregnancy loss does require EC.
Any other episodes of UPSI since last menstrual period?	Yes <input type="checkbox"/> No <input type="checkbox"/>	Consider possibility of pregnancy and if necessary signpost to GP/SHS.
Has the patient taken LNG-EC or another progesterone within the last 7 days?	Yes <input type="checkbox"/> No <input type="checkbox"/>	If yes then UPA-EC is less effective, advise Cu-IUD or use LNG-EC.
Are there any concerns in regard to unsafe relationships/child abuse/adult protection concerns?	Yes <input type="checkbox"/> No <input type="checkbox"/>	If yes contact relevant local service.

Medical History	Yes	No	Action/Information
Allergy to UPA-EC or LNG-EC?			If yes advise Cu-IUD and signpost for fitting. If declined refer to GP or Sexual Health Service (SHS).
Current unexplained vaginal bleeding?			If yes signpost to GP or Sexual Health Service (SHS) or Out of Hours
BMI >26kg/m ² or >70kg in weight			If yes advise Cu-IUD (first line), UPA-EC if suitable or LNG-EC 3000 microgram dose (unlicensed).
Currently breastfeeding?			LNG-EC or IUD*. Advise to discard breast milk for 7 days after UPA-EC use. *Increased risk of intrauterine perforation with IUD – would be for discussion at SHS.
Current severe disease treated with oral glucocorticoids e.g. asthma?			If yes UPA-EC not suitable, consider LNG-EC if UPSI is <96 hours or refer to GP or SHS if greater.
Severe malabsorption syndrome e.g. Crohn's disease or severe diarrhoea?			If yes signpost for Cu-IUD as LNG-EC and UPA-EC may be less effective.
Porphyria?			If yes UPA-EC is not suitable – advise Cu-IUD or use LNG-EC.
Currently taking medicines that increase gastric pH?			UPA-EC will have a reduced effect if PPI taken in the last 7 days or H2 antagonist or antacid taken within the last 24 hours.
Currently taking enzyme inducing medication including St. John's Wort?			If yes UPA-EC is not suitable. The only licensed option is an IUD or consider LNG-EC 3000 microgram dose (unlicensed).
Other significant drug interactions			If interaction cannot be managed then refer to SHS or relevant specialist.

Refer to flowchart for choice of UPA-EC/LNG-EC/Cu-IUD depending on the answers provided above

Counselling Checklist to be Discussed Prior to Treatment

Pregnancy Risk: Days 9-16 of /28 cycle Days 1-8 and >16 of /28 cycle LNG-EC within 96 hours UPA-EC within 120 hours Copper IUD up to 120 hours after UPSI / or ovulation		20-30% risk of pregnancy with x 1 UPSI 2-3% risk of pregnancy with x 1 UPSI 2-3 in 100 women will become pregnant 1-2 in 100 women will become pregnant < 1 in 100 women will become pregnant	
<input type="checkbox"/>	Cu-IUD discussed as most effective 1 st line option	<input type="checkbox"/>	Mode of action, efficacy and failure rates (see above)
<input type="checkbox"/>	Action if vomiting occurs within 3 hours (return for an additional dose)	<input type="checkbox"/>	Explain any common side effects
<input type="checkbox"/>	If EC fails there is no increased risk of foetal abnormality	<input type="checkbox"/>	Next period may be late/early and light bleeding may occur over the next few days (not to be counted as a period)
<input type="checkbox"/>	Return if there is a further episode of UPSI	<input type="checkbox"/>	Patient to read the PIL for the EC
<input type="checkbox"/>	When to seek medical advice i.e. should severe abdominal pain occur	<input type="checkbox"/>	If no normal menstrual period within 3 weeks take pregnancy test

Contraception Advice (when appropriate)

Bridging/Quick Start Contraception Discussed	Yes <input type="checkbox"/>	No <input type="checkbox"/>	Barrier methods discussed	Yes <input type="checkbox"/>	No <input type="checkbox"/>
<input type="checkbox"/> Client declined ongoing contraception/advice			<input type="checkbox"/> Bridging contraception supplied (desogestrel)		

Outcome of consultation (tick all that apply)

<input type="checkbox"/>	Signposted for Cu-IUD insertion	<input type="checkbox"/>	No EC required
<input type="checkbox"/>	LNG-EC 1500 microgram single dose under PGD Batch No:	<input type="checkbox"/>	UPA-EC 30mg single dose Batch No:
<input type="checkbox"/>	LNG-EC 3000microgram single dose under PGD (unlicensed) Batch No:	<input type="checkbox"/>	EC not indicated and declines Cu-IUD (Refer to SHS or GP)
Referral	Signposted to Sexual Health Service <input type="checkbox"/>	Signposted to Out of Hours Service <input type="checkbox"/>	Signposted to GP <input type="checkbox"/>

Consent

Emergency hormonal contraception treatment risks have been fully explained to me and I agree to treatment. I have been informed of how my data will be stored and who will be able to access that information, as well as how it may be used.

Patient Signature		Date	
Healthcare Professional Supplying Signature		Date	