

## Oral Contraceptives Risk Assessment Form

Title: Mr. ☐ Mrs. ☐ Miss ☐ Ms. ☐ Other ☐	D.o.B.	:/	/	Age:					
Name:	Home Address:								
Surname:	-								
	Name	& Addres	ss of GP						
Email:	-								
Telephone:									
Please answer the following questions									
Do you have any recent or past medical history of note?  If yes, please provide details	Yes□	No□	Is the contraceptive for your	own use?	Yes No				
Do you take any current or repeat medicines?  If yes, please provide details	Yes	No 🗆	Is there a possibility you may	y be pregnant?	Yes No				
Are you over 35 years of age and a smoker?	Yes	No 🗆	Are you overweight? Or have If yes, please provide details healthcare professional can be	below (if you are unsure, your	Yes No No				
Do you have a family history of blood clots or thrombosis?  If yes, please provide details	Yes	No 🗆	Do you have any of the follo migraine headaches, cancer, disease? If yes, please provide details	HIV, high blood pressure, liver	Yes No No				
Have you been taking your current contraceptive pill for more than a year?	Yes	No 🗆	Have you been prescribed the more than 9 months?	ne same contraceptive pill for	Yes No No				
Have you had a check up with your doctor / nurse about your contraceptive pill in the last year?  If yes, please provide details			Are you having any problem contraceptive pill such as irrulf yes, please provide details						
Please write below any further information which may be relevant e.g. medicines, conditions									
Confidential sexual health patient helplines									

FPA (formerly the Family Planning Association) national helpline – 0845 3101334

Sexual Health Line – 0800 567 123

Brook Clinic – 0800 0815023 or www.brook.org.uk

Sexwise – 0800 282930 or http://www.maketherightdecision.co.uk





## FOR OFFICIAL USE-HEALTHCARE PROFESSIONAL USE ONLY

Retain completed forms for 8 years. Fax or post copy of completed form to GP within 3 months

This form is intended to be used per supply. For additional supplies to the same patient, a new form will be needed.

Product: (e.g. Microgynon)									
Date	Quantity*	Referral required?	Directions	Pharmacist	Signature				
l		Yes ☐ No☐ If yes, see below.	Take once daily.		1				
Reason for referral									
* Maximum of one issue of 3 calendar-packs in each 9 month period.									
Missed pill advice									
Combined oral contraceptive pills (21 active tablets)									
Take the missed pill as soon as you remember then continue as normal, with extra precautions (barrier methods) required for 7 days. These rules									
are applicable no matter where in the packet the patient is.  Progestogen-only pills									
		remembered. If more than o	ne pill has been m	issed, only one pill should be taken	. The next pill should				
The missed pill should be taken as soon as remembered. If more than one pill has been missed, only one pill should be taken. The next pill should be taken at the usual time. This may mean that two pills are taken in 1 day. Additional contraceptive precautions (condoms or avoidance of sex)									
are advised for 2 days (48 hours) after restarting the POP.									
The patient information leaflet that comes with the pill might say to use condoms for the next seven days after remembering to take the pill. This									
is because it takes seven days for the pill to suppress ovulation.									
If the patient has had UPSI and missed pills then consider EC (as per emergency contraception PGD) do not supply COCs/POPs to patients with suspected pregnancy.									
Additional advice									
STIs	Barrier Contraceptives	Sexual Health Helplines	Efficacy	LARCs (IUD, IUS, or Implanon)	Cervical screening				
Record of supply									
Drug brand, ba	tch number and expiry date.	Date Qty	Details Price		Comment				
I confirm that	the patient is not contraindicate	ed based on the information prov	vided by the PGD						
I have explaine	ed the potential warnings and sig	de effects of the treatment to th	e patient, and reque	ested they report them if they occur					
I have provided the patient with an information leaflet (PIL) for the treatment I am administering, and advised them to read it									
					<del>_</del>				
PATIENT CONSENT									
I have received information on the risks and benefits of the medicines recommended and fully understand them. I have also had the opportunity									
to ask questions. I consent to the recommended medicines being given at each appointment.									
Patient signati	ire	Date							
Healthcare pro	fessional signature		Date						
	<u> </u>								