Appendix 1. Information for Patients and Health Care Professionals

Information for Patients		Information for Health Care Professionals	
•	Individuals with BRCA1/BRCA2 gene alterations have an 11-68% lifetime risk of developing ovarian	•	Patients with inherited BRCA1 and BRCA2 gene alterations should be offered risk-reducing surgery at
	cancer, compared with approx. 2% in the general population.		appropriate ages.
•	Risk-reducing salpingo-oophorectomy reduces the lifetime risk of ovarian cancer by at least 90%.	•	Patients should be fully counseled about the risks of surgery, infertility, iatrogenic menopause and the need for HRT.
•	In order to be eligible for risk-reducing surgery, individuals should have completed their family and be		
	aged over 35 years with BRCA1 or over 40 years with BRCA2 alterations.	•	Ideally, cases should be discussed at a familial cancer multidisciplinary team meeting to confirm
			surgery is appropriate.
•	There is a theoretical possibility of greater blood loss with risk-reducing surgery at the time of cesarean		
	delivery because of large pelvic blood vessels in pregnancy. However, there is no evidence to suggest	•	If the individual has had breast cancer, her oncology team should be consulted and be happy that
	this is a significant risk.		surgery is appropriate and that her likely prognosis justifies the negative consequences of undergoing
			RRSO.
•	Risk-reducing surgery, whenever performed, will result in infertility. However, assisted reproduction		
	techniques with the option of pre-implantation genetic diagnosis, may be feasible as long as eggs are	•	Patients should ideally have an up-to-date ultrasound scan of the pelvis to confirm no macroscopic
	harvested prior to risk-reducing surgery.		tubo-ovarian disease. The adnexa should be visualised and commented on at each pregnancy scan until
			it is no longer possible to do so. Pre-operative serum CA125 is unreliable as it can be elevated in
•	Risk-reducing surgery will result in early menopause. Hormone replacement therapy is recommended		pregnancy.
	until 51 years of age, unless there are contraindications (e.g. previous breast cancer).		
		•	Careful choice of appropriate surgical equipment is recommended for performing RRSO because of the
			engorged pelvic blood vessels in pregnancy.

Barker VE, Vlachodimitropoulou E, O'Brien P, Iskaros J, Rosenthal AN, et al. Combined bilateral salpingo-oophorectomy and cesarean delivery in *BRCA1/2* alteration carriers: a case series. Obstet Gynecol 2023;142.

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• The pathologist should be made aware that the patient is a <i>BRCA</i> -carrier, and the SEE-FIM protocol should be followed to minimise the chance of missing an occult tubal/ovarian cancer.
• Patients should be counseled that if histopathological assessment identifies cancer, they will likely need further staging surgery and chemotherapy.
• Transdermal continuous combined HRT should be started 6 weeks postnatally (or later in those able and desiring to breast feed) and continued to the age of 51 years unless there are contraindications.
• In patients with a history of breast cancer, there is a theoretical risk of HRT stimulating micro- metastases. The patient's oncology team should be consulted pre-operatively and a decision made regarding HRT usage based on risks and benefits.
• Advice for emergency obstetric teams in the event of patient requiring emergency cesarean delivery prior to her planned cesarean delivery: Take intra-operative peritoneal washings for cytology using warm saline instilled around adnexae prior to uterine incision; to avoid contamination with liquor, suction bottle should then be changed. Place left and right adnexae in separately labelled formalin containing specimen pots and request SEE-FIM histopathology protocol.

Key: SEE-FIM = Sectioning and Extensively Examining the FIMbriated end, HRT = hormone replacement therapy.

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