

## Radiotherapy consent form – vaginal vault brachytherapy for gynaecologic cancer

This form should only be used if the patient is over 16 years old and has capacity to give consent. If the patient does not legally have capacity please use an appropriate alternative consent form from your hospital.

Patient details						
Patient name:  Patient unique identifier:		Date of birth:  Name of hospital:				
				Responsible consultant oncologist or consultant theraputic radiographer:  Special requirements: eg, transport, interpreter, assistance		
Details of radioth	erapy					
Radiotherapy type:	Vaginal vault radiotherapy (brachytherapy)					
Aim of treatment: (Tick as appropriate)	Adjuvant – treatment given	the best chance of being cured given after surgery to reduce the risk of cancer coming back liative – to improve your symptoms and/or help you live longer but not to				
Contact details are provide	s before starting, during or after y ed here for any further queries, ke to discuss your treatment further.	our radiotherapy.				

Patient name:	Patient unique identifier:				
Possible early/short-term side-effects					
Start during radiotherapy or shortly after completing radiotherapy and usually resolve within two to six months of finishing radiotherapy. Frequencies are approximate.					
Expected 50%–100%	☐ Tiredness				
<b>Common</b> 10%–50%	☐ Mild pelvic pain				
Less common Less than 10%	<ul> <li>□ Urinary frequency (passing urine more often than normal) and urgency (a sudden urge to pass urine)</li> <li>□ Cystitis/pain when you urinate</li> <li>□ Bowel urgency (a sudden urge to open your bowels)</li> <li>□ Bowel frequency (opening your bowels more often than normal)</li> <li>□ Rectal pain/discomfort</li> <li>□ Looser stools compared to normal</li> <li>□ Vaginal itching, discharge or light bleeding (spotting)</li> <li>□ Moderate pelvic pain</li> </ul>				
Rare Less than 1%	☐ Bleeding from your bladder or bowel				
Specific risks to you from your treatment					

I confirm that I have had the above side-effects explained.

Patient initials

Patient name:		Patient unique identifier:			
Possible late or long-term side-effects					
		d may be permanent. Frequencies are approximate. often referred to as pelvic radiation disease.			
<b>Definite</b> 100%	☐ Vaginal narrowing, shortening or dryness – this may impact vaginal intercourse, and the comfort and quality of a vaginal examination. You may be advised to use vaginal dilators after treatment which may reduce this risk.				
<b>Expected</b> 50%–100%	☐ Bleeding from the vagina after using vaginal dilators or vaginal sexual intercourse				
<b>Common</b> 10%–50%	<ul> <li>☐ Urinary frequency (passing urine rurge to pass urine)</li> <li>☐ Bowel urgency (a sudden urge to operate of the compared to normal)</li> </ul>	nore often than normal) and <b>urgency</b> (a sudden pen your bowels)			
Less common Less than 10%	you receive anal sex.  Bleeding from bowel or bladder  Bowel/bladder damage which r	wels more often than normal) y worsen on opening your bowels. This may also affect your sex life if			
Rare Less than 1%	☐ A different cancer in the treatm	ent area			
Specific risks to you from your treatment					

I confirm that I have had the above side-effects explained.

Patient initials

Patient name:	Patient unique identifier:	
	to be filled in by health professional with	
<ul> <li>I have discussed what the treatment is likely to involve, the in</li> <li>I have also discussed the benefits and risks of any available a</li> <li>I have discussed any particular concerns of this patient.</li> </ul>		his treatment.
Patient information leaflet provided:  Yes / No – Details: Copy of consent form accepted by patient:  Yes / No Signature:	Date:	
Name:	Job title:	
Statement of patient  I have had the aims and possible side effects of treatment opportunity to discuss alternative treatment and I agree to described on this form.  I understand that a guarantee cannot be given that a partice radiotherapy. The person will, however, have appropriate end of the person to be described about additional procedures which are necess become necessary during my treatment. This may include photographs to help with treatment planning and identificated.  I agree that information collected during my treatment, increcords may be used for education, audit and research. All I am aware I can withdraw consent at anytime.  Tick if relevant  I understand that if I were to continue to smoke it could have side-effects I experience and the efficacy of my treatment.	Statement of interpreter witness (where appropriate)  I have interpreted the information contained in this form to the patient to the best of my ability and in a way in which I believe they can understand.  or I confirm that the patient is unable to sign but has indicated their consent.  Signature:	
Patient name:	Date:	Name:  Date:
		Patient confirmation