

Radiotherapy consent form for lymphoma

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:		Date of birth:		
Patient unique identifier:		Name of hospital:		
Responsible consultant	oncologist or consultant therape	eutic radiographer:		
Special requirements: eg,	transport, interpreter, assistance			
Dataila of vadiath				
Details of radiothe	erapy			
Radiotherapy type:	External beam radiotherapy			
Site:				
Aim of treatment: (Tick as appropriate)		with the intent of curing your cancer		
	Palliative – to prevent, redu	ce or delay the symptoms but not to cure your cancer		
•	before starting, during or after y	our radiotherapy.		
-	ed here for any further queries, ke to discuss your treatment further.			

Possible early/short-term side-effects

These depend on which area of your body is treated, and the dose of radiotherapy, so your team will explain which are relevant for you. They start during radiotherapy or shortly after completing radiotherapy and usually resolve within two to six months of finishing radiotherapy.

	Expected 50%–100%	Common 10%-50%	Less common Less than 10%	Rare Less than 1%	Not applicable to you
General regardless of site: Tiredness					
Skin (in the treated area): Skin soreness, itching, swelling and colour changes*					
Localised hair loss					
Head and/or neck: Eye: redness, watering or light sensitivity					
Mouth: redness, pain, ulceration, dryness or taste change					
Throat: pain or hoarse voice					
Chest: Pain on or difficulty swallowing Shortness of breath or cough					
Abdomen and/or pelvis: Nausea	П				П
Vomiting					
Indigestion					
Diarrhoea					
Urinary frequency (passing urine more often than normal)					
Limb: Limb swelling					

 $^{^{\}star}$ redness in white skin tones and subtle darkness, yellow/purple/grey appearance in brown and black skin tones

Specific risks to you from your treatment

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

Patient initials

Possible late or long-term side-effects

These may happen many months or years after radiotherapy and may be permanent.

	Expected 50%–100%	Common 10%–50%	Less common Less than 10%	Rare Less than 1%	Not applicable to you
Skin (in the treated area): Colour change (lighter or darker for any skin tone) and sensitivity to sun and temperature change					
Poor hair growth					
Head and/or neck: Dry eyes Cataracts Dry mouth Dental problems Hypothyroidism Increased risk of stroke					
Chest: Increased risk of heart disease, especially angina and heart attack Increased risk of lung scarring (fibrosis)					
Abdomen and/or pelvis: Worsening kidney function Reduced spleen function, increased risk of infections Increased risk of diabetes Infertility Early menopause Low testosterone levels					
Limb: Joint stiffness Swelling of the limbs (lymphoedema)					
Increased risk of secondary cancer specifically:					

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		
•	appropriate knowledge of proposed procedure)		
 I have discussed what the treatment is likely to involve, the ir I have also discussed the benefits and risks of any available a I have discussed any particular concerns of this patient. 			
Patient information leaflet provided: Yes / No – Details:			
Copy of consent form accepted by patient: Yes / No			
Signature:	Date:		
Name:	Job title:		
Statement of patient		Statement of:	
I have had the aims and possible side effects of treatment opportunity to discuss alternative treatment and I agree to described on this form.	witness (where appropriate)		
 described on this form. I understand that a guarantee cannot be given that a partic radiotherapy. The person will, however, have appropriate e I have been told about additional procedures which are ne to treatment or may become necessary during my treatme include permanent skin marks and photographs to help with planning and identification. 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. 	☐ 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.		
Tick if relevant			
☐ I confirm that there is no risk that I could be pregnant.	Signature:		
☐ I understand that I should not become pregnant during treatment.			
Note: if there is any possibility of you being pregnant you must tell your hospital doctor your treatment as this can cause significant harm to an unborn fetus. Testosterone and are not contraception	Name:		
I understand that I should not conceive a child or donate sp my treatment and I will discuss with my oncologist when it child after radiotherapy.	Date:		
☐ I understand that if I were to continue to smoke it could have side-effects I experience and the efficacy of my treatment.	Patient confirmation		
I do not have a pacemaker and/or implantable cardioverter or	of consent (To be signed prior to the start of radiotherapy)		
I have a pacemaker and/or implantable cardioverter defibring risks associated with this explained to me. Signature:	I confirm that I have no further questions and wish to go ahead with treatment.		
		Dation	
Patient name:	Date:	Patient initials	
		Date:	