



The Royal College of Radiologists

Radiotherapy consent form for pancreatic 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:

Date of birth:

Patient unique identifier:

Name of hospital:

Responsible consultant oncologist or consultant therapeutic radiographer:

Special requirements: eg, transport, interpreter, assistance

Details of radiotherapy

Radiotherapy type:

External beam radiotherapy

Site:

Pancreas

Aim of treatment:

(Tick as appropriate)

- Neo-adjuvant** – treatment given before surgery to shrink the tumour
- Adjuvant** – treatment given after surgery to reduce the risk of cancer coming back
- Disease control** – to help you live longer but not to cure your cancer
- Palliative** – to improve your symptoms but not to cure your cancer

Concurrent systemic
anti-cancer therapy:

(Tick as appropriate)

- Yes with** _____
- No**

(A separate consent form will cover the possible side-effects of this treatment)

You may have questions before starting, during or after your radiotherapy.

Contact details are provided here for any further queries,
concerns or if you would like to discuss your treatment further.

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% 	<input type="checkbox"/> Tiredness <input type="checkbox"/> Nausea – feeling sick <input type="checkbox"/> Abdominal discomfort or bloating
Common 10%–50% 	<input type="checkbox"/> Diarrhoea <input type="checkbox"/> Vomiting <input type="checkbox"/> Indigestion or heartburn <input type="checkbox"/> Loss of appetite <input type="checkbox"/> Weight loss <input type="checkbox"/> Abdominal pain or cramping
Less common Less than 10% 	<input type="checkbox"/> Ulcers in the stomach or bowel <input type="checkbox"/> Bleeding from the stomach or bowel <input type="checkbox"/> Skin soreness, itching and colour changes in treatment area – white/lighter skin: pink, red, darker than surrounding area; brown skin: maroon or darker than surrounding area; black skin: darker than surrounding area, yellow/purple/grey colour changes <input type="checkbox"/> Hair loss in the treatment area
Rare Less than 1% 	<input type="checkbox"/> Bowel perforation – a hole in your 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

May happen many months or years after radiotherapy and may be permanent.
Frequencies are approximate.

Expected 50%–100%	
Common 10%–50%	<input type="checkbox"/> Diabetes – new onset diabetes or complications from existing diabetes <input type="checkbox"/> Malabsorption – difficulty digesting or absorbing nutrients from food
Less common Less than 10%	<input type="checkbox"/> Ulcers in the stomach or bowel <input type="checkbox"/> Bleeding from the stomach or bowel <input type="checkbox"/> Bowel narrowing or obstruction (blockage of the bowel) <input type="checkbox"/> Bowel perforation – a hole in your bowel <input type="checkbox"/> Reduced spleen function leading to increased risk of infection <input type="checkbox"/> Skin colour change in the treatment area – usually lighter or darker for any skin tone <input type="checkbox"/> Telangiectasia in the treatment area – small visible blood vessels which look like spidery marks
Rare Less than 1%	<input type="checkbox"/> A different cancer in the treatment area <input type="checkbox"/> Long term decline in kidney function
Specific risks to you from your treatment	
I confirm that I have had the above side-effects explained.	
	Patient initials <input type="text"/>

Patient name:

Patient unique identifier:

Statement of health professional

(to be filled in by health professional with appropriate knowledge of proposed procedure)

- I have discussed what the treatment is likely to involve, the intended aims and side-effects of this treatment.
- I have also discussed the benefits and risks of any available alternative treatments including no treatment.
- 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

- I have had the aims and possible side effects of treatment explained to me and the opportunity to discuss alternative treatment and I agree to the course of treatment described on this form.
- I understand that a guarantee cannot be given that a particular person will perform the radiotherapy. The person will, however, have appropriate expertise.
- I have been told about additional procedures which are necessary prior to treatment or may become necessary during my treatment. This may include permanent skin marks and photographs to help with treatment planning and identification.
- I agree that information collected during my treatment, including images and my health records may be used for education, audit and research. All information will be anonymised. I am aware I can withdraw consent at anytime.

Tick if relevant

- I confirm that there is no risk that I could be pregnant.
- 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/health professional before your treatment as this can cause significant harm to an unborn fetus. Testosterone and other hormone treatments are not contraception.

- I understand that I should not conceive a child or donate sperm or eggs during the course of my treatment and I will discuss with my oncologist when it will be safe for me to conceive a child after radiotherapy.

- I understand that if I were to continue to smoke it could have a significant impact on the side-effects I experience and the efficacy of my treatment.

- I do not have a pacemaker and/or implantable cardioverter defibrillator (ICD).

or

- I have a pacemaker and/or implantable cardioverter defibrillator (ICD) and I have had the risks associated with this explained to me.

Signature:

Patient name:

Date:

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:

Name:

Date:

Patient confirmation of consent

(To be signed prior to the start of radiotherapy)

I confirm that I have no further questions and wish to go ahead with treatment.

Patient initials

Date: