



# **RADICAL CYSTECTOMY WITH BLADDER REPLACEMENT (IN MEN)**

Information about your procedure from  
**The British Association of Urological Surgeons (BAUS)**

This leaflet contains evidence-based information about your proposed urological procedure. We have consulted specialist surgeons during its preparation, so that it represents best practice in UK urology. You should use it in addition to any advice already given to you.

To view the online version of this leaflet, type the text below into your web browser:

[http://www.baus.org.uk/\\_userfiles/pages/files/Patients/Leaflets/Rad cyst male neobladder.pdf](http://www.baus.org.uk/_userfiles/pages/files/Patients/Leaflets/Rad%20cyst%20male%20neobladder.pdf)

## **Key Points**

- The aim of cystectomy is to remove your bladder and the cancer that it contains
- This usually involves removal of your bladder, prostate and local lymph glands to give us the best chance of removing all the cancer
- We construct a reservoir for your urine from a section of bowel which is connected to your urethra (waterpipe) to form a “neobladder”
- This procedure can be done using an open or keyhole approach (often with robotic assistance)

## **What does this procedure involve?**

Removal of your bladder, prostate and local lymph nodes. We construct a reservoir for your urine from a section of bowel which is connected to your urethra (waterpipe) to form a “neobladder”.

## **What are the alternatives?**

- **[Radical radiotherapy](#)** – if, for any reason, a decision has been made that you should not undergo surgery
- **[BCG instillation into the bladder](#)** – only indicated for high-grade non-muscle invasive cancer; not for muscle-invasive disease
- **[Radical cystectomy with formation of a urostomy \(ileal conduit\)](#)** or **[construction of a catheterisable pouch](#)** – radical bladder

removal but with, respectively, an external urostomy or a catheterisable reservoir instead of a new bladder

- [Systemic chemotherapy](#) – drugs given by mouth or into your bloodstream; sometimes used before radiotherapy or radical cystectomy

## **What happens on the day of the procedure?**

Your urologist (or a member of their team) will briefly review your history and medications, and will discuss the surgery again with you to confirm your consent.

An anaesthetist will see you to discuss the options of a general anaesthetic or spinal anaesthetic. The anaesthetist will also discuss pain relief after the procedure with you.

We may provide you with a pair of TED stockings to wear, and we may give you a heparin injection to thin your blood. These help to prevent blood clots from developing and passing into your lungs. Your medical team will decide whether you need to continue these after you go home.

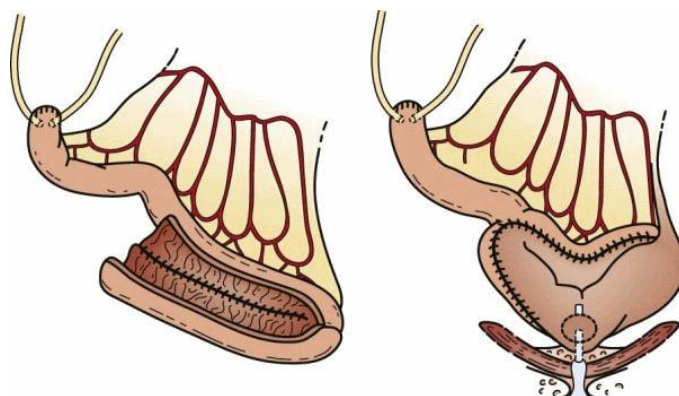
You will normally see a specialist nurse whose job it is to teach you how to manage your neobladder. He/she will also show you how to self-catheterise and perform bladder washouts. We usually mark your abdomen to position a urostomy stoma in case, for any reason, a neobladder cannot be made and you need to have a urostomy (ileal conduit). Try not to wash off this mark before your procedure.

Some urology units will admit you the day before your procedure to administer laxatives to clear your bowel; others will admit you on the day of surgery and will not use laxatives. Your urologist will advise on which protocol will be used.

## **Details of the procedure**

- we use a general anaesthetic for the procedure
- your anaesthetist may also use an epidural or spinal anaesthetic to minimise any post-operative pain
- we usually give you an injection of antibiotics before the procedure, after you have been checked for any allergies
- we normally perform the procedure through an incision in your lower abdomen (tummy) or with a laparoscopic (keyhole) approach with or without robotic assistance
- we will remove your bladder, prostate, seminal vesicles (sperm sacs) and, if necessary, your urethra (waterpipe)

- we may be able to preserve the nerves which control your erections but this is not always possible; you should discuss this with your urologist before the procedure



- we stitch the ureters (the tubes which drain urine from the kidneys to the bladder) to a separated piece of small bowel which is then fashioned into a bladder substitute and joined to your urethra (pictured)
- we re-join the ends of the small bowel from which the urostomy has been made
- we close the wound with stitches or staples
- we normally place one or more drains where the bladder has been removed and leave two small splints in the ureters protruding from your urethra or through the skin of your abdomen
- we normally put two catheters in your bladder, one through your urethra and the other through the skin of your abdomen
- the procedure can take anything from two to six hours to perform, depending on how complex the procedure is and what approach (open or keyhole) is used
- you should expect to be in hospital for seven to 14 days

Following radical cystectomy, some urology units have introduced [Enhanced Recovery Pathways](#). These actually start before you are admitted to hospital. After your surgery, they are designed to speed your recovery, shorten your time in hospital and reduce your risk of re-admission.








We will encourage you to get up and about as soon as possible. This reduces the risk of blood clots in your legs and helps your bowel to start working again. You will sit out in a chair shortly after the procedure and be shown deep breathing/leg exercises. We will encourage you to start drinking and eating as soon as possible.












We normally remove your drains when they have stopped draining. The splints in your bladder substitute will be removed 10 to 14 days after the





procedure when an X-ray along the splints has shown that the ureters and bowel have joined securely.

## Are there any after-effects?

The possible after-effects and your risk of getting them are shown below. Some are self-limiting or reversible, but others are not. We have not listed very rare after-effects (occurring in less than 1 in 250 patients) individually. The impact of these after-effects can vary a lot from patient to patient; you should ask your surgeon's advice about the risks and their impact on you as an individual:

| After-effect   | Risk   |
|--|--|
| Inability to ejaculate (or father children) because your sperm-carrying mechanism has been removed   |  All patients  |
| Inability to get an erection (impotence)   |  Almost all patients (unless the nerves can be preserved) |
| Need for self-catheterisation to wash out and empty your neobladder (the frequency of this usually decreases as time goes on)  |  Almost all patients                                     |
| Your cancer may not be cured by the procedure (this possibility will have been discussed with you before the operation)  |  Between 1 in 2 & 1 in 10 patients                       |
| Need for blood transfusion   |  Between 1 in 2 & 1 in 10 patients                       |
| Paralytic ileus (failure of your bowel to work) for several days which causes nausea, bloating & vomiting, and needs an intravenous drip and a stomach drainage tube through your nose until it recovers |  Between 1 in 2 & 1 in 10 patients                       |
| Incontinence of urine is common in the early weeks and improves in most men but can persist, especially at night   |  Between 1 in 3 & 1 in 7 patients (15 to 30%)            |

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| Anaesthetic or cardiovascular problems possibly requiring intensive care (including chest infection, pulmonary embolus, stroke, deep vein thrombosis, heart attack and death) |    | Between 1 in 10 & 1 in 50 patients (your anaesthetist can estimate your individual risk) |
| Infection in your wound or an abscess in your abdominal (tummy) cavity  |    | Between 1 in 10 & 1 in 50 patients   |
| Decrease in kidney function over time   |    | Between 1 in 10 & 1 in 50 patients   |
| Need to abandon a keyhole procedure and convert to open surgery because of operative difficulties   |    | Between 1 in 10 & 1 in 50 patients   |
| Difficulty emptying your neobladder requiring <a href="#">intermittent self-catheterisation</a>   |   | Between 1 in 10 & 1 in 50 patients   |
| Long-term bladder stone formation   |  | Between 1 in 10 & 1 in 50 patients   |
| Long-term recurrent urinary infections  |  | Between 1 in 10 & 1 in 50 patients   |
| Biochemical imbalance due to reabsorption of acid from your neobladder requiring regular treatment with bicarbonate tablets   |  | Between 1 in 10 & 1 in 50 patients   |
| Need for return to theatres for re-operation due to bleeding, bowel leakage, urine leakage or bowel obstruction   |  | Between 1 in 50 & 1 in 250 patients  |
| Hernia of the incision (or keyhole ports) requiring further surgical repair   |  | Between 1 in 50 & 1 in 250 patients  |
| Scarring, narrowing or hernia formation around your urostomy requiring revision   |  | Between 1 in 50 & 1 in 250 patients  |

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|---|--|
| Diarrhoea/vitamin deficiency due to shortened bowel requiring specific dietary supplements or other treatment |  Between 1 in 50 & 1 in 250 patients |
| Late stricturing (scarring) of the bowel or ureters requiring further surgery                                 |  Between 1 in 50 & 1 in 250 patients |
| Rectal injury at the time of surgery requiring a temporary colostomy (bowel opening on your abdomen)          |  Between 1 in 50 & 1 in 250 patients |
| Spontaneous rupture of your neobladder with urinary leakage requiring further surgery                         |  Between 1 in 50 & 1 in 250 patients |

### What is my risk of a hospital-acquired infection?

Your risk of getting an infection in hospital is between 4 & 6%; this includes getting *MRSA* or a *Clostridium difficile* bowel infection. This figure is higher if you are in a “high-risk” group of patients such as patients who have had:

- long-term drainage tubes (e.g. catheters);
- bladder removal;
- long hospital stays; or
- multiple hospital admissions.

### What can I expect when I get home?

- you will be given advice about your recovery at home
- you will be given a copy of your discharge summary and a copy will also be sent to your GP
- any antibiotics or other tablets you may need will be arranged & dispensed from the hospital pharmacy
- by the time you get home, you should be able to perform “daily living” activities such as making a cup of tea, preparing food and emptying your stoma bag
- you will feel tired for several weeks and may need to take regular “cat naps” during the day
- we will arrange for the stitches or clips in your wound to be removed after 10 to 12 days
- if you get a fever or start vomiting, especially if associated with unexpected pain in your abdomen, you should contact your GP immediately



- a follow-up appointment will be made for you six to 12 weeks after your surgery

## **General information about surgical procedures**

### ***Before your procedure***

Please tell a member of the medical team if you have:

- an implanted foreign body (stent, joint replacement, pacemaker, heart valve, blood vessel graft);
- a regular prescription for a blood thinning agent (e.g. warfarin, aspirin, clopidogrel, rivaroxaban, dabigatran);
- a present or previous MRSA infection; or
- a high risk of variant-CJD (e.g. if you have had a corneal transplant, a neurosurgical dural transplant or human growth hormone treatment).

### ***Questions you may wish to ask***

If you wish to learn more about what will happen, you can find a list of suggested questions called "[Having An Operation](#)" on the website of the Royal College of Surgeons of England. You may also wish to ask your surgeon for his/her personal results and experience with this procedure.

### ***Before you go home***

We will tell you how the procedure went and you should:

- make sure you understand what has been done;
- ask the surgeon if everything went as planned;
- let the staff know if you have any discomfort;
- ask what you can (and cannot) do at home;
- make sure you know what happens next; and
- ask when you can return to normal activities.

We will give you advice about what to look out for when you get home. Your surgeon or nurse will also give you details of who to contact, and how to contact them, in the event of problems.

### ***Smoking and surgery***

Ideally, we would prefer you to stop smoking before any procedure. Smoking can cause bladder cancer, increase the risk of cancer recurrence and make complications more likely after surgery. We strongly advise anyone with bladder cancer to stop smoking. For advice on stopping, you can:

- contact your GP;

- access your local [NHS Smoking Help Online](#); or
- ring the free NHS Smoking Helpline on **0300 123 1044**.

### ***Driving after surgery***

It is your responsibility to make sure you are fit to drive after any surgical procedure. You only need to [contact the DVLA](#) if your ability to drive is likely to be affected for more than three months. If it is, you should check with your insurance company before driving again.

### **What should I do with this information?**

Thank you for taking the trouble to read this information. Please let your urologist (or specialist nurse) know if you would like to have a copy for your own records. If you wish, the medical or nursing staff can also arrange to file a copy in your hospital notes.

### **What sources have we used to prepare this leaflet?**

This leaflet uses information from consensus panels and other evidence-based sources including:

- the [Department of Health \(England\)](#);
- the [Cochrane Collaboration](#); and
- the [National Institute for Health and Care Excellence \(NICE\)](#).

It also follows style guidelines from:

- the [Royal National Institute for Blind People \(RNIB\)](#);
- the [Information Standard](#);
- the [Patient Information Forum](#); and
- the [Plain English Campaign](#).

### **Disclaimer**

We have made every effort to give accurate information but there may still be errors or omissions in this leaflet. BAUS cannot accept responsibility for any loss from action taken (or not taken) as a result of this information.

#### **PLEASE NOTE**

The staff at BAUS are not medically trained, and are unable to answer questions about the information provided in this leaflet. If you do have any questions, you should contact your urologist, specialist nurse or GP.