

# RETAINER Data Collection

---

Record number (automatic)

---

---

Study ID  
(Centre ID followed by 3 Digit subject number, e.g.  
UHL001, UHL 002...)

---

---

Has patient consented to be contacted in the future  
as part of RETAINER II (patient reported outcome  
measures following hernia repair?)

- Yes  
 No

---

Gender

- Male  
 Female

---

Patient Age

---

---

Patient BMI

---

---

ASA Grade

- 1  
 2  
 3  
 4  
 5

---

Current Medications (Tick all that apply)

- None  
 Alpha-blocker (e.g. Tamsulosin, Silodosin,  
Doxazosin, Alfuzosin, Terazosin)  
 5-Alpha reductase inhibitor (e.g. Dutasteride,  
Finasteride)  
 Dutasteride/Tamsulosin (Combodart)  
 Anti-androgen therapy (hormone manipulation, e.g.  
for prostate cancer)  
 Urological Anti-Cholinergic (e.g. Solifenacin,  
Fesoterodine, Darifenacin, Trospium, Propiverine,  
Oxybutynin patch/tablet)  
 Non-urological medication with powerful  
anti-cholinergic properties (e.g. certain  
psychiatric and anti-Parkinson medications)  
 Mirabegron (Betmiga)  
 Solifenacin/Tamsulosin (Vesomni)  
 Intravesical Botox in past 12 months

---

Known Neurological Condition with Potential to Affect  
Voiding

- None  
 Spinal Cord Injury  
 Spina bifida  
 Multiple Sclerosis  
 Other neurological condition with potential to  
affect voiding

---

Has the patient ever been diagnosed with one of the following urological conditions?

- None
- BPH (benign prostatic hyperplasia)
- Prostate cancer
- Urethral stricture
- Bladder neck stenosis
- Hypospadias or other congenital anomaly of the lower genitourinary tract
- Detrusor failure
- Overactive bladder

---

Has the patient ever had surgery on the bladder or prostate? (Purely diagnostic procedures e.g. cystoscopy not included)

- Never
- TURP (trans-urethral resection of prostate)
- Bladder neck incision or dilation
- TURBT (transurethral resection of bladder tumour)
- Prostatectomy (removal of prostate) for cancer or benign disease via any approach
- Urethral stricture surgery (urethroplasty/urethral dilation/any)
- Botox injection to bladder within past year
- Bladder augmentation or other complex reconstruction

---

Has patient had previous episode of urinary retention requiring a catheter?

- Yes
- No

---

IPSS Score

- N/A - female patient
- 0
- 1
- 2
- 3
- 4
- 5
- 6
- 7
- 8
- 9
- 10
- 11
- 12
- 13
- 14
- 15
- 16
- 17
- 18
- 19
- 20
- 21
- 22
- 23
- 24
- 25
- 26
- 27
- 28
- 29
- 30
- 31
- 32
- 33
- 34
- 35

---

Post void residual measurement pre-operatively, in millilitres (If possible to obtain. A reading from previous 12 months is acceptable)

- Could not obtain
- < 50ml
- 50-100ml
- 100-150ml
- 150-200ml
- 200-250ml
- 250-300ml
- 300-350ml
- 350-400ml
- >400ml

---

When did the patient's bowels last open prior to surgery?

- Day of surgery
- Day before surgery
- 2 days before surgery
- >2 days before surgery

---

Time of surgery

- 7am - 1pm
- 1pm-5pm
- After 5pm

---

Laterality of hernia repair

- Left
- Right
- Bilateral

Approach	<input type="radio"/> Open <input type="radio"/> Laparoscopic <input type="radio"/> Robotic <input type="radio"/> Minimally Invasive Converted to Open
Bladder involved in hernia?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Unsure
Type of anaesthesia used? Please tick all that apply.	<input type="checkbox"/> General <input type="checkbox"/> Spinal <input type="checkbox"/> Epidural <input type="checkbox"/> Local
If general anaesthesia, were any of the following agents used?	<input type="checkbox"/> No <input type="checkbox"/> GA not used or agents not known <input type="checkbox"/> Glycopyrrolate <input type="checkbox"/> Atropine <input type="checkbox"/> Diazepam <input type="checkbox"/> Thiopentone <input type="checkbox"/> Propofol <input type="checkbox"/> Isoflurane <input type="checkbox"/> Sevoflurane <input type="checkbox"/> Enflurane
If spinal or epidural anaesthesia, which of the following agents were used? (please tick all that apply)	<input type="checkbox"/> N/A (no spinal/epidural used) <input type="checkbox"/> Bupivacaine <input type="checkbox"/> Levobupivacaine <input type="checkbox"/> Fentanyl <input type="checkbox"/> Sufentanil <input type="checkbox"/> None of these <input type="checkbox"/> Not sure
If local anaesthetic used, which drug?	<input type="radio"/> N/A (no local used) <input type="radio"/> Levobupivacaine (Chirocaine / Marcaine) <input type="radio"/> Lidocaine (Lignocaine) <input type="radio"/> Mixture of Levobupivacaine / Lidocaine <input type="radio"/> Other <input type="radio"/> Not sure
Perioperative analgesics administered (Intraoperatively or within 8 hours, tick all that apply)	<input type="checkbox"/> Paracetamol <input type="checkbox"/> NSAIDs (non steroidal anti-inflammatories) <input type="checkbox"/> Opioids <input type="checkbox"/> Steroids <input type="checkbox"/> Clonidine
Perioperative IV fluids, volume infused (millilitres)	_____
Was a urinary catheter inserted during or prior to the procedure and removed prior to reversal of anaesthesia?	<input type="radio"/> Yes <input type="radio"/> No
Duration of surgery, skin-to-skin (minutes)	<input type="radio"/> < 30 minutes <input type="radio"/> 30-60 minutes <input type="radio"/> 60-120 minutes <input type="radio"/> >120 minutes <input type="radio"/> Unknown

---

Intraoperative injuries recognised?	<input type="checkbox"/> None <input type="checkbox"/> Bowel injury <input type="checkbox"/> Urological injury <input type="checkbox"/> Ilioinguinal nerve injury/intentional sacrifice <input type="checkbox"/> Other
-------------------------------------	--

---

Mesh used?	<input type="radio"/> Yes <input type="radio"/> No
------------	---

---

Type of mesh used?	<input type="radio"/> None <input type="radio"/> Non absorbable (e.g. polypropylene, PTFE, polyester) <input type="radio"/> Absorbable (e.g. Vicryl) <input type="radio"/> Composite (e.g. PP, Polyglactin 'Vypro', Collagen 'Parietex') <input type="radio"/> Biologic mesh (human dermis, porcine dermis) <input type="radio"/> Bovine (e.g. 'Permacol', 'Alloderm', 'Surgisis')
--------------------	---

---

Drain used?	<input type="radio"/> Yes <input type="radio"/> No
-------------	---

---

Did patient resume normal voiding (passage of urine) postoperatively (on the day of surgery) by clinician's judgement?	<input type="radio"/> Yes <input type="radio"/> No
--	---

---

Was a bladder scan performed postoperatively for a post-void residual measurement (PVR) as part of routine clinical care?	<input type="radio"/> Yes <input type="radio"/> No
---	---

---

If a PVR was measured postoperatively, what was the value?	<input type="radio"/> PVR not measured <input type="radio"/> < 50ml <input type="radio"/> 50-99ml <input type="radio"/> 100-199ml <input type="radio"/> 200-299ml <input type="radio"/> 300ml or more
--	--

---

Was patient admitted to hospital unexpectedly on day of surgery (e.g. from a planned day surgery pathway?)	<input type="radio"/> Yes <input type="radio"/> No
--	---

---

If patient was admitted to hospital on the day of surgery, was urinary retention the primary reason for admission or a significant contributor?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not admitted
---	---

---

Was patient readmitted to hospital (following discharge) within 30 days of surgery?	<input type="radio"/> Yes <input type="radio"/> No
---	---

---

If patient was readmitted to hospital following discharge within 30 days of surgery, was urinary retention the primary reason for admission or a significant contributor?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> No readmission
---	---

---

Was a diagnosis of urinary retention made within 1 week of surgery? (If the answer is no, the data entry for this patient is complete)	<input type="radio"/> Yes <input type="radio"/> No
--	---

---

---

The remaining questions apply only if urinary retention developed.  
If urinary retention occurred, when was the diagnosis made?

- Day of surgery
- Day 1 postoperatively
- Day 2 postoperatively
- Day 3 postoperatively
- Day 4 postoperatively
- Day 5 postoperatively
- Day 6 postoperatively
- Day 7 postoperatively

---

How was the diagnosis of retention made? Please tick all that apply.

- Failure to void over a given period
- Suprapubic pain/pressure
- Palpable bladder
- Bladder scan reading above a given level

---

How did urinary retention manifest itself?

- Patient did not pass urine at all, or passed only small dribbles, postoperatively
- Patient initially seemed to be passing urine normally, but subsequently retention evolved

---

If diagnosis was made by failure to void, at how many hours postoperatively or from the last void was this made?

- < 4
- 4
- 5
- 6
- 7
- 8
- 9
- 10
- 11
- 12
- >12

---

If retention was diagnosed by a bladder scan, at what scan volume was diagnosis made?

- < 300ml
- 300 - 400ml
- 400 - 500ml
- 500 - 600ml
- 600 - 700ml
- 700 - 800ml
- 800 - 900ml
- 900 - 1,000ml
- >1,000ml

---

If urinary catheter placed, what was the residual volume of urine?

- < 300ml
- 300 - 400ml
- 400 - 500ml
- 500 - 600ml
- 600 - 700ml
- 700 - 800ml
- 800 - 900ml
- 900 - 1,000ml
- > 1,000ml

---

Digital rectal exam (DRE) findings (tick all that apply)

- DRE not done
- Small prostate
- Moderate prostate
- Large prostate
- Suspicious / hard / malignant prostate
- Stool in rectum
- "Normal"

---

How was retention managed in the first instance?	<input type="radio"/> Urethral catheter (indwelling) <input type="radio"/> Self-intermittent catheterisation <input type="radio"/> Suprapubic catheter
--	--

---

If urethral catheter was placed or attempted, how many catheterisation attempts were required?	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4 or more <input type="radio"/> No attempts to pass urethral catheter
--	---

---

Was a urology consult performed?	<input type="radio"/> Yes <input type="radio"/> No
----------------------------------	---

---

Was an alpha blocker started? (e.g. Tamsulosin, Silodosin, Alfuzosin) (do not include combination pills e.g. Combodart here)	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Patient was already taking one
--	---

---

Was a 5-alpha reductase inhibitor started? (e.g. Dutasteride, Finasteride) (do not include combination pills e.g. Combodart here)	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Patient was already taking one
---	---

---

Was a tablet combining an alpha blocker and a 5-alpha reductase inhibitor started? (e.g. Dutasteride/Tamsulosin 'Combodart')	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Patient was already taking one
--	---

---

Were any longterm medications stopped due to the urinary retention?	<input type="radio"/> No <input type="radio"/> Anti-cholinergic stopped (including combination tablets e.g. 'Vesomni') <input type="radio"/> Mirabegron (Betmiga) stopped
---	---

---

Were laxatives/suppositories/enemas given as constipation was thought to be contributing to urinary retention?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Takes regularly, not changed
--	---

---

If admitted, how many inpatient nights were involved?	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> 4 <input type="radio"/> 5 <input type="radio"/> >5 <input type="radio"/> Not admitted
---	---

---

Was urinary retention, or complications related to same, the primary reason for this length of stay?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> N/A - not admitted
--	---

---

Did the patient develop an acute kidney injury within the first postoperative week?	<input type="radio"/> Yes <input type="radio"/> No
---	---

---

Was a urinary tract infection detected in this patient?	<input type="radio"/> Yes - on day of admission <input type="radio"/> Yes - between postoperative day 1 and day 7 or discharge <input type="radio"/> No
---	---

---

Was a course of antibiotics used to treat a suspected urinary tract infection?	<input type="radio"/> Yes <input type="radio"/> No
--	---

---

Did patient suffer pain / a frequent sensation of needing to void / bladder spasm with a urinary catheter?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> N/A - did not have an indwelling catheter
--	--

---

Were any additional complications seen related to catheterisation? Please tick all that apply	<input type="checkbox"/> Traumatic catheterisation <input type="checkbox"/> Haematuria <input type="checkbox"/> Catheter dislodged with balloon inflated <input type="checkbox"/> Delirium <input type="checkbox"/> Impaired mobility due to catheterisation
---	--

---

Did patient undergo successful trial without catheter? (i.e. resume normal voiding without an indwelling urethral or suprapubic catheter and without the need to perform self-catheterisation in the longterm)	<input type="radio"/> Yes <input type="radio"/> No
--	---

---

From time of initial catheterisation, on what day was trial without catheter passed?	<input type="radio"/> Day 1 <input type="radio"/> Day 2 <input type="radio"/> Day 3 <input type="radio"/> Day 4 <input type="radio"/> Day 5 <input type="radio"/> Day 7 <input type="radio"/> Day 7-14 <input type="radio"/> > Day 14 <input type="radio"/> Trial without catheter not passed
--	---

---

How many times has a trial of catheter removal been performed in this patient following the postoperative urinary retention?	<input type="radio"/> 1 <input type="radio"/> 2 <input type="radio"/> 3 <input type="radio"/> >3 <input type="radio"/> No trials without catheter to date <input type="radio"/> N/A as catheter was not inserted
--	---

---

Was urological intervention/training performed whilst patient was admitted? (Tick all that apply)	<input type="checkbox"/> Urology service was required to place a urethral catheter <input type="checkbox"/> Cystoscopy was required to place a urethral catheter <input type="checkbox"/> Suprapubic catheter was inserted <input type="checkbox"/> Urethral stricture dilation was performed <input type="checkbox"/> Transurethral resection of prostate was performed <input type="checkbox"/> Patient was trained in intermittent self-catheterisation
---	---

---

What was the patient's status at discharge?	<input type="checkbox"/> Discharged without catheter, voiding by self <input type="checkbox"/> Discharged without catheter, performing intermittent self catheterisation <input type="checkbox"/> Discharged with indwelling urethral catheter, on free drainage <input type="checkbox"/> Discharged with indwelling urethral catheter, with flip-flo valve <input type="checkbox"/> Discharged with suprapubic catheter, on free drainage <input type="checkbox"/> Discharged with suprapubic catheter, with flip-flo valve
---	---

---

Was urology follow-up planned from time of discharge?	<input type="radio"/> Yes <input type="radio"/> No
---	---