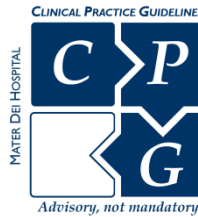


Urinary Retention Management Guidelines

Department of Urology

**EXPLANATORY
NOTES AND
REFERENCES**



Mater Dei Hospital – July 2018

Urinary Retention Guideline Development Group

Guideline Co-ordinator: **Mr John Sciberras**
Consultant Urologist

Guideline Developer: **Mr Keith Pace**
Basic Specialist Trainee, Department of Surgery

GDG members:

Mr Gerald Busuttil	Resident Specialist, Urology			
Dr Tonio Piscopo	Consultant Physician			

Annex

Introduction

Acute Urinary Retention (AUR) is defined as a painful, palpable or percussable bladder, when the patient is unable to pass any urine¹. Whilst Chronic Urinary Retention (CUR) is said to occur when there is a non-painful bladder, which remains palpable or percussable after the patient has passed urine¹. CUR is often defined by the volume of post void residual (PVR). However, there is no consensus on the cut off PVR and whilst some have defined it as a volume of >300mls², others have defined other PVRs as the cut off volumes or given no cut off PVR at all^{3,4}.

Acute Urinary Retention

AUR constitutes 45% of all lower urinary tract (LUT) consultations⁵ and will be encountered by most physicians whatever their specialty.

AUR may be spontaneous or may be preceded by a trigger factor⁶. In spontaneous AUR there is no trigger identified, with AUR being part of the natural history of Benign Prostatic Hyperplasia (BPH), usually after a long period of LUT symptoms (LUTS). Triggered AUR may be preceded by bladder over-distension, surgery with general or regional anaesthesia, excess fluid intake, alcohol consumption, urinary tract infection, prostatic inflammation, faecal impaction, ano-rectal pain or use of drugs with sympathomimetic, anticholinergic or anti-histamine effects⁷.

The risk of developing AUR is directly related to prostate size (>30mls associated with three-fold increased risk)⁸, increased age, increased LUTS, poor flow rate, and larger PVR⁹.

Acute Urinary Retention Management

1. Catheterise the patient. Urethral catheterisation is indicated for the relief of urinary retention. If urethral catheterisation fails contact the urologist for consideration of catheterisation using a hydrophilic guidewire or suprapubic catheterisation^{10,11}.
2. Measure and record the residual volume^{10,11}.
3. Test Urine and take Serum Renal Profile^{10,11}.
4. Start Alpha Blocker – There is moderate quality evidence to suggest that the rate of successful trial without catheter (TWOC) increases with use of alpha blockers and the incidence of



recurrent acute urinary retention is lower in groups treated with an alpha blocker. This data was statistically significant for alfuzosin, tamsulosin and silodosin¹².

Alfuzosin is given as a 10mg daily dose; tamsulosin is given as a 400mcg daily dose; whilst the dose for silodosin is 8mg daily¹³. Alfuzosin and tamsulosin may cause orthostatic hypotension, whilst silodosin is highly urospecific and less likely to cause this adverse effect. However, it is more likely to cause retrograde ejaculation, albeit with preserved orgasm¹⁴.

Prior to starting an alpha blocker, the patient should be asked about planned cataract surgery. If such surgery is planned the initiation of alpha blockers should be avoided until after the surgery is complete in view of the risk of developing floppy iris syndrome intraoperatively¹⁵.

5. Trial Without Catheter (TWOC) - There is no consensus on when this should be performed. In the UK the majority of patients undergo TWOC at 2 days post insertion of catheter; in France the majority undergo TWOC at 3 days. A few factors are known to influence success of TWOC. Age <70, detrusor pressure >35 mmH₂O, initial residual <1L, prolonged catheterisation and an identified trigger are associated with higher success rates of TWOC. However, one should keep in mind that prolonged catheterisation is associated with higher morbidity including urosepsis, haematuria and urine leak¹⁶.
6. If TWOC is successful the patient will need follow up of LUTS and medication review. If unsuccessful re-TWOC or surgery should be considered^{10, 11}.

Chronic Urinary Retention

CUR is invariably linked to an increased PVR; however, The European Association of Urology (EAU), American Urology Association and NICE guidelines on LUTS do not define threshold values for pathological PVR¹⁷.

CUR is classified as High Pressure (HPCUR) or Low Pressure (LPCUR) based on urodynamic findings. Those with a bladder end filling pressure of <25cmH₂O are described as LPCUR, while those with higher end filling pressures are classified as HPCUR¹. Intravesical pressure remains high throughout the voiding cycle in HPCUR.

The two groups also defer in signs and symptoms at presentation with LPCUR patients complaining of hesitancy, slow stream and incomplete emptying. These patients are usually followed up in an outpatient setting and only require catheterisation if symptomatic¹¹.

HPCUR patients complain of urgency. Their serum creatinine tends to be elevated and imaging reveals dilation of the upper urinary tracts¹⁸. HPCUR patients require immediate catheterisation with careful recording of residual volumes and close monitoring of electrolytes as inpatients¹¹.

Post Obstructive Diuresis (POD)

Urine production exceeding 200 mL per hour for 2 consecutive hours or producing greater than 3 L of urine in 24 hours is diagnostic of POD¹⁹.

Diuresis may be a physiological process post obstruction and helps to eliminate excess volume and accumulated solutes. However, this should resolve once haemostasis is achieved. Some patients will continue to have diuresis even after haemostasis is achieved and this is referred to as pathological POD. The pathophysiology is unclear; however, it is likely to be due to a combination of down-regulation of sodium transporters in the ascending loop of Henle, reduction in the glomerular filtration rate with resultant loss of juxtamedullary nephrons and a degree of nephrogenic diabetes insipidus²⁰.

If the patient's urine output exceeds 200mls per hour for 2 consecutive hours:

1. The patient requires admission for close monitoring for conversion to pathological POD.
2. These patients should have close urine output charting.
3. Daily weight should be recorded.
4. Serum renal profile should be monitored every 12 hours.
5. Urine should be collected for urinary sodium, potassium and osmolality.
6. All patients with pathological POD require intravenous fluid replacement run at a negative balance, with fluid type tailored to serum and urinary electrolyte levels and hydration status.
7. These patients will benefit from a consultation with a nephrologist²¹.

References

1. Abrams P, Cardozo L, Fall M et al. Standardisation Sub-committee of the International Continence Society. The standardisation of terminology of lower urinary tract function: report from the Standardisation Sub-committee of the International Continence Society. *Neurourol Urodyn* 2002; 21: 167 – 78
2. Kaplan SA, Wein AJ, Staskin DR , Roehrborn CG , Steers WD. Urinary retention and post-void residual urine in men: separating truth from tradition. *J Urol* 2008 ;180: 47 – 54
3. Ghalayini IF Al-Ghazo MA , Pickard RS . A prospective randomized trial comparing transurethral prostatic resection and clean intermittent self-catheterization in men with chronic urinary retention. *BJU Int* 2005; 96: 93 –7
4. Thomas AW, Cannon A, Bartlett E, Ellis-Jones J, Abrams P. The natural history of lower urinary tract dysfunction in men: the influence of detrusor underactivity on the outcome after transurethral resection of the prostate with a minimum 10-year urodynamic follow-up. *BJU Int* 2004; 93: 745 – 50
5. Roghmann F, Ghani K, Kowalczyk K et al. Incidence and Treatment Patterns in Males Presenting with Lower Urinary Tract Symptoms to the Emergency Department in the United States. *J Urol* 2013; 190: 1798-1804.
6. Desgrandshamps F, De la Taille A, Doublet JD. The management of Acute Urinary Retention in France: a cross sectional survey in 2618 men with Benign Prostatic Hypertrophy. *BJUI* 2016; 97: 727.
7. Fitzpatrick JM, Desgrandshamps F, Adjali K. Management of Acute Urinary Retention: a worldwide survey of 6074 men with benign prostatic hyperplasia. *BJUI* 2011; 109; 88-95.
8. Jacobsen SJ, et al. Natural history of prostatism: risk factors for acute urinary retention. *J Urol*. 1997;158:481-487
9. Jacobsen N, Jacobsen D, Girman C et al. Natural History of Prostatism: Risk Factors for Acute Urinary Retention. *J Urol* 1997; 158: 481-7
10. Lower urinary tract symptoms in men: management (2010 updated 2015) NICE guideline CG97
11. Speakman M in *Oxford Textbook of Urological Surgery*, Hamdy F. et al. 2018
12. Fisher E, Subramonian K, Omar MI. The role of alpha blockers prior to removal of urethral catheter for acute urinary retention in men. *Cochrane Database of Systematic Reviews* 2014, Issue 6.
13. *British National Formulary (2018)*, Royal Pharmaceutical Society of Great Britain.
14. Kobayashi K, Masumori N, Kato R. Orgasm is preserved regardless of ejaculatory dysfunction with selective α 1A-blocker administration. [Int J Impot Res](#) 2009 Sep; 21(5): 306–310.



15. American Urology Association. Management of Benign Prostatic Hyperplasia (2010; reviewed and validity confirmed 2014).
16. Muruganandham K, Dubey D, and Kapoor R. Acute urinary retention in benign prostatic hyperplasia: Risk factors and current management. *Indian J Urol*. 2007 Oct-Dec; 23(4): 347–353.
17. Negro C, Gordon M. Chronic urinary retention in men: How we define it, and how does it affect treatment outcome. *BJUI* 2012; 110, 1590–1594.
18. O'Reilly PH, Broomam PJ, Farah NB et al. High pressure chronic retention. Incidence, aetiology and sinister implications. *Br J Urol* 1986; 58: 644–6.
19. Vaughan ED Jr., Gillenwater JY. Diagnosis, characterization and management of postobstructive diuresis. *J Urol* 1973;109(2):286-92.
20. Loo MH, Vaughan ED. Obstructive nephropathy and postobstructive diuresis. *AUA Update Series* 1985; 4: 1-7.
21. Halbgewachs C, Domes T. Postobstructive diuresis. *Canadian Family Physician* Feb 2015, 61 (2) 137-142;

The Urinary Retention Management Guideline **MDHCPG/SUR01v1.0/2018** is approved.

This is a clinical practice guideline and does not in any way replace or supersede the clinical discretion necessary in its implementation



Signed:

Mr John Sciberrs
Consultant Urologist
Guideline Co-ordinator

Mr Keith Pace
Basic Specialist Trainee, Department of Surgery
Guideline Developer

GDG members

Mr Gerald Busuttil

Dr Tonio Piscopo

