

#### Initial Assessment to identify underlying cause

#### **Standard Urotherapy**

# Behavioural/Lifestyle changes

Modify Fluid intake + adequate urination intervals Adequate positioning for voiding and defecation Avoidance of caffeine, chocolate and citrus foods

Non-neurogenic
Inadequate response to urotherapy



#### Constipation

Hydration and fibre intake Treatment with laxatives



#### Neurogenic

May already undergoing intermittent catheterisation

# First Line Pharmacological Treatment with Anticholinergics

Solifenacin 5mg OD (TLS green)

Increased to 10mg OD if beneficial but not curative

Trial duration <u>minimum 1 month</u> at maximum tolerated dose and review Requires annual review. Review 6monthly if frail or cognitive impairment.



Contraindicated or not tolerated

# **Alternative Pharmacological Options with Anticholinergics**

**Trospium Chloride 20mg BD** (standard release) (TLS blue) See BNSSG Formulary for alternative formulary options

Trial duration <u>minimum 1 month</u> at maximum tolerated dose and review Requires annual review. Review 6monthly if frail or cognitive impairment.



Contraindicated, not tolerated or ineffective

## **Second Line Pharmacological Option**

NB: Can be used first line after bladder retraining if anticholinergics are contraindicated or the patient/clinician does not want to trial anticholinergics due to anticholinergic burden

Mirabegron 50mg OD (TLS blue) as per NICE TA290

Trial duration minimum 6 weeks at maximum tolerated dose.

Requires annual review including BP check. Review 6 monthly if frail, cognitive impairment or hypertension.

Consider combination antimuscarinic therapy such as Solifenacin + Mirabegron<sup>6</sup> (TLS Blue)

Trial duration minimum 1 month at maximum tolerated dose





### Referred for detrusor activity assessment using urodynamics and MDT decision

# Non-neurogenic detrusor over activity (DO)

Persistent wetting despite maximal medical therapy

#### **Neurogenic detrusor over activity (DO)**

Persistent neurogenic detrusor overactivity and/or poor bladder

### **Botulinum Toxin A (Botox®) administration**

**Non-neurogenic** 100 units under local/general anaesthetic **Neurogenic** 200 units under local/general anaesthetic

Contraindicated, not tolerated, ineffective, or not willing or able to self-catheterise

# Surgical Intervention

Sacral neuromodulation/ Ileal conduit/ augmentation cystoplasty using intestinal segment

Telephone consultation with Nurse in 6 weeks.
Reassess in clinic after 3 months to assess
outcomes and for suitability of repeated
Botulinum toxin A (Botox®) injections after 6-9
months according to duration of action in
individual patient [based on return of DO +/-

# Re-treatment

Following a minimum of a 3 month interval after Botulinum Toxin A (Botox®) treatment, if urinary incontinence symptoms return, patients can self-refer for repeated treatment.

#### **Stopping criteria**

If Botox treatment is contra-indicated, not tolerated, causes adverse events or does not improve symptoms.

#### References

- 1. NICE Clinical Guideline CG148 Urinary incontinence in neurological disease: assessment and management. <a href="https://www.nice.org.uk/guidance/cg148">https://www.nice.org.uk/guidance/cg148</a>
- 2. NICE Clinical Guideline NG123 urinary incontinence and pelvic organ prolapse
- 3. NICE Technology Appraisal Guidance TA290 Mirabegron for treating symptoms of overactive bladder. <a href="https://www.nice.org.uk/guidance/ta290/chapter/2-The-technology">https://www.nice.org.uk/guidance/ta290/chapter/2-The-technology</a> [Accessed 13.2.20]
- 4. Summary of Product Characteristics BOTOX® 100 Units (last updated on the eMC: 12.06.19) https://www.medicines.org.uk/emc/product/859/smpc [accessed 20 August 2019]
- 5. Summary of Product Characteristics BOTOX® 200 Units (last updated on the eMC: 12.06.19) https://www.medicines.org.uk/emc/product/436/smpc [accessed 20 August 2019]
- 6. Jefferson, K (North Bristol Trust) (2020) Evidence Review: Solifenacin and Mirabegron Combination Therapy for Overactive Bladder. (attached supporting document)