



Standard Urotherapy

Behavioural/Lifestyle changes

Fluid intake

Adequate urination intervals

Adequate positioning for voiding and defecation

Avoidance of caffeine, chocolate and citrus foods

Minimum 2 months

ConstipationHydration and fibre intake

Treatment with laxatives

Non-neurogenic

Inadequate response to urotherapy



Neurogenic

May already undergoing intermittent catheterisation

Pharmacological Treatment with Anticholinergics as per **BNSSG Formulary**

Trial duration minimum 1 month at maximum tolerated/licensed dose and review

If ineffective or not tolerated, try alternative formulary option for minimum 1 month at maximum tolerated/licensed dose and review



Contraindicated, not tolerated or ineffective

Mirabegron (TLS blue) as per NICE TA290

Trial duration <u>minimum 6 weeks</u> at maximum tolerated/licensed dose

Consider combination antimuscarinic therapy such as Solifenacin + Mirabegron⁶ (TLS Blue)

Trial duration <u>minimum 1 month</u> at maximum tolerated dose



Contraindicated, not tolerated or ineffective

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 compliance despite maximal medical therapy

Botulinum Toxin A 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 injections after 6-9 months according to duration of action in individual patient [based on return of DO +/- wetting]



Use of Botulinum Toxin A for Overactive Bladder Adults



Re-treatment

Following a minimum of a 3 month interval after Botulinum Toxin A 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. https://www.nice.org.uk/guidance/cg148
- 2. NICE Clinical Guideline NG123 urinary incontinence and pelvic organ prolapse
- 3. NICE Technology Appraisal Guidance TA290 Mirabegron for treating symptoms of overactive bladder. https://www.nice.org.uk/guidance/ta290/chapter/2-The-technology [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)