

Dear {{customText[Dr|Prof|A|Prof|Mr|Mrs]}} {{accFname}} {{accLname}},

Over 10 years of experience supports BETMIGA™ for alleviating debilitating symptoms of overactive bladder (OAB) and improving patient quality of life



BETMIGA™ is a well-established, first-in-class β 3-adrenoceptor agonist approved for the symptomatic treatment of urgency, increased micturition frequency and/or urgency incontinence as may occur in adult patients with OAB syndrome, providing patients with similar efficacy to antimuscarinics but with a **more favourable tolerability profile**¹⁻⁷

2013

The launch of
BETMIGA™,⁶

~23 mil

patients treated
with BETMIGA™,⁷

69

countries with
access to
BETMIGA™,⁷

Over **15,900 patients across 19 clinical studies** and **19,200 patients in extensive real-world data worldwide** have been treated with BETMIGA™, supporting its established safety profile^{1-4,8-43}

Effective reduction of OAB symptoms



Extensive data from **46,666 patients** across **64 RCTs** (from an SLR and NMA) show that **BETMIGA™ reduces key OAB symptoms** significantly better than placebo and similar to a range of common antimuscarinics⁴⁴

Established safety profile, including elderly and male patients



Clinical trials demonstrate that **BETMIGA™ has low overall incidence of TEAEs**, with most being mild to moderate^{24,25,45}

BETMIGA™ has an **established safety profile in ≥65-year-olds and male patients**^{25,26,46}

No increased risk of MACE



In a pooled analysis of 13 studies and 13,396 patients, BETMIGA™ had **comparable cardiovascular safety profile** vs placebo and antimuscarinics, with **no increased risk of MACE**⁴⁷

Positively impacts patient QoL



BETMIGA™ **significantly improves patients' QoL**, with improvements in sleep quality, coping with symptoms or concerns about incontinence⁴⁸



BETMIGA™ is recommended by the EAU 2023 guidelines as **first-line** pharmacological treatment for symptomatic treatment of urgency, increased micturition frequency and/or urgency incontinence as may occur in adult patients with OAB syndrome⁴⁹



The effectiveness and tolerability of BETMIGA™ is confirmed in **real-world clinical practice**⁸⁻¹³



Extensive data from five real-world studies showed that **BETMIGA™ consistently improves bladder symptoms**⁸⁻¹³



BETMIGA™ maintains an improvement in the symptoms of OAB **over 1 year**, helping to reduce the impact of OAB on **patients' HRQoL in the long term**⁵⁰

EAU, European Association of Urology; HRQoL, health-related quality of life; MACE, major adverse cardiovascular events; NMA, network metaanalysis; OAB, overactive bladder; RCT, randomised clinical trial; QoL, quality of life; SLR, systematic literature review; TEAE, treatment emergent adverse event.



Effective management of OAB symptoms requires **long-term treatment** and **patient persistence**, given the **chronic nature** of the condition⁵⁰



Persistence with **antimuscarinics** is very **low (8% after 2 years)**, primarily due to **side effects**⁵¹⁻⁵⁴

Over **10 years of real-world data** show that **BETMIGA™ consistently demonstrates higher persistence rates** when compared to placebo and antimuscarinics^{10, 55-57}

BETMIGA™ persistence ensures **long-term OAB management**, resulting in **better symptom control** and **improved economic and QoL outcomes**⁵⁰

BETMIGA™

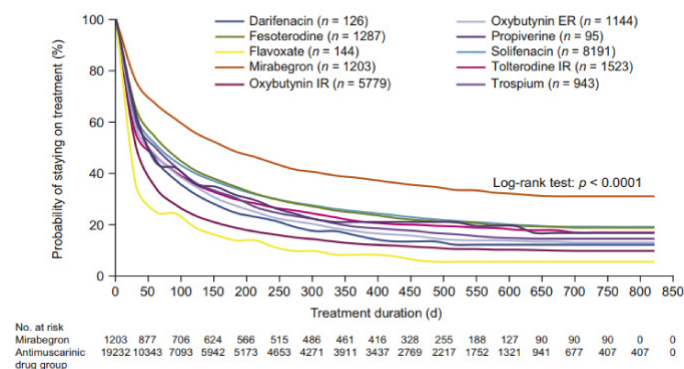
169

Tolterodine*

56 ↓

Other antimuscarinics†

30-78 ↓

Time to discontinuation for BETMIGA™ vs antimuscarinics⁴²

BETMIGA™ is **cost-saving** and **cost-effective** compared with antimuscarinics across multiple countries, due to lower overall treatment costs and HCRU⁵⁷⁻⁶⁵



€69.2 billion

the economic burden of UI in 2024, with the highest cost attributed to:⁶⁶



Productivity losses: absenteeism, presenteeism, caregiver costs (40% in men and 43% in women)



Incontinence pads (28% in men and 22% in women)

First-line treatment with BETMIGA™ is associated with **lower overall annual costs** and **reduced treatment switches, GP and urologist visits, and UTI-related expenses** compared to antimuscarinics⁶²⁻⁶⁴

Key drivers of cost-effectiveness of BETMIGA™ vs antimuscarinics include **utility gains, improved tolerability, better persistence and decreased HCRU^{57,65}**

*(HR 1.55; 95% CI 1.41, 1.71; $p < 0.0001$); †($p < 0.001$ for all comparisons).

GP, general practitioner; HCRU, healthcare resource utilization; OAB, overactive bladder; QoL, quality of life; UI, urinary incontinence; UTI, urinary tract infection.

Click here for BETMIGA™ Abbreviated Prescribing Information

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Yours sincerely,
{userName}

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