

Medium-Term Outcomes of Transcutaneous Tibial Nerve Stimulation in Patients with Refractory Overactive Bladder

Gubbiotti M.¹, Antonioni A.¹, Zucchi A.², Rosadi S.¹

1. Valdarno Hospital, Dept. of Urology, USL Toscana Sud-Est, Italy
2. University of Pisa, Dept. of Urology, Pisa, Italy

www.ics.org/2026/abstract/

Abstract #654

INTRODUCTION

Transcutaneous posterior tibial nerve stimulation (T-PTNS) is a non-invasive neuromodulation technique. **Aim of the study: to assess the safety and efficacy of T-PTNS in patients with refractory OAB over a medium-term follow-up period.**

METHODS

This single-centre prospective study included **OAB patients** unresponsive to pharmacological therapy. Baseline evaluation included a 3-day bladder diary, urodynamics (UD), a symptom- bother VAS (0= worst; 10= best), and the Overactive Bladder questionnaire-Short Form (OAB-q SF). After in-person training, patients used **T-PTNS** (Tensi+®) at home for **20 minutes daily** (as per protocol). **Assessment were repeated at 1, 3, 6, and 12- months; UD was repeated at 12-months.** Success was defined as a $\geq 50\%$ reduction in urgency episodes (with or without urgency incontinence) or $\geq 30\%$ reduction in 24-hour voiding frequency.

CONCLUSION

Home-based T-PTNS (Tensi+®) was safe and well tolerated, with sustained symptom and quality of life improvements in patients with refractory OAB.



RESULTS

Thirty-two OAB patients (6M, 26F); mean age: 61.4 ± 18.8 y.o. Mean OAB symptoms duration: 5.9 ± 3.9 years. No patient was taking OAB medication at baseline; all had a post-void residual volume (PVR) < 50 mL. **At 3 months, urgency episodes decreased in all patients; 28/32 (87.5%) improved day- and night-time urinary frequency, and 24/32 (75%) reported reduced urgency incontinence episodes.** Benefits persisted at 12 months. UD showed increased maximum cystometric capacity and volume at first uninhibited detrusor contraction ($p < 0.001$), while stable PVR ($p > 0.1$). OAB-q SF scores and VAS satisfaction significantly improved (both $p < 0.001$). Overall treatment success rate was 95%. Pain decreased in the 3 patients with chronic pelvic pain (VAS, $p < 0.001$). No significant adverse events occurred.