

Clinical Experience Program Recommendations

Name of clinic: [Clinic/institution na	me]	
Medical director: [Lead physician co	ontact]	
Additional point of contact: [Other	contact if needed, e.g. administrator, lab manager] _	
Local representative: [Sales directo	r or inside sales]	
Clinical specialist: [For any clinical of	questions]	
Goal: [Number]	patients by [Date]	

Patient selection

- Indications: Primary snoring and mild obstructive sleep apnea (AHI <15 events/hour)
- Likely responders:
 - Individuals with a BMI <35 kg/m²
 - Those with a higher Friedman Tongue Position score (see sidebar)
 - Those without significant facial or oropharyngeal abnormalities such as class 2 malocclusion
 - Not currently taking central nervous system depressants or suffering from alcohol use disorder
- Contraindications:
 - 1) Pregnancy 2) Pacemaker or implanted electrodes 3) Dental jewelry in the mouth
 - 4) Suffering from mouth ulcers



Assessing the Friedman Tongue Position

Ask the patient to open their mouth widely while breathing normally with the tongue in the natural position. Repeat a minimum of five times in order to assess the most consistent position of the tongue as follows: I) Uvula and tonsils/pillar visible; IIa) Most of the uvula is visible, but not the tonsils/pillar; IIb) The entire soft palate is visible, but only the base of the uvula; III) Some of the soft palate is visible, with the distal end absent; IV) Only the hard palate is visible. See Friedman *et al.*, Advances in Otorhinolayrngology (2017).

Guidance and resources for patients

- Excessive salivation, a tingling sensation on the tongue, and/or some tooth sensitivity is expected, particularly in the early days of use
- Start with a low level of stimulation and ramp up over time.
 Patients should use therapy at their highest tolerated level on any given day, which will differ person to person.
- Consistency is key! Patients should use therapy every day
 for the first six weeks; after that, the smartphone app will
 automatically move them to the maintenance phase, during
 which patients should use therapy two or more times per week.
- The eXcite^{OSA®} team is here to help patients should use the in-app chat with therapy questions or for technical troubleshooting.

Contact Signifier Medical Technologies if you require any assistance accessing or interpreting the cloud-based data. Our team can also provide slides and other materials for internal presentations.

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Suggested clinical follow-up

- Document patient characteristics that may be related to the treatment response in the medical record for retrospective analysis, such as craniofacial and upper airway anatomy.
- Complete a sleep study at baseline and 6-12 weeks after initiating therapy, ideally with the same sleep diagnostic device type for both pre- and post-therapy.
- Interview the bed-partner, if possible, to gather insights on snoring and witnessed apneas.
- Collect patient-reported outcomes that are sensitive to the milder end of the sleep-disordered breathing spectrum, either instead of or in addition to standard measures such as the Epworth Sleepiness Scale:
 - PROMIS Sleep Disturbance scale
 - PROMIS Sleep-Related Impairment scale
 - Functional Outcomes of Sleep Questionnaire
- Include data from the HIPAA-compliant eXcite^{OSA} cloud-based portal in your analyses. Data available in the portal include:
 - Objective adherence
 - Stimulation level



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