SleepFlex™ Treatment of Obstructive Sleep Apnea NCT04726514

June 15, 2023

Study Protocol (Summary)

Study participants with mild to moderate obstructive sleep apnea (OSA) underwent 12 weeks of active treatment with the SleepFlex™ program of head and neck exercises.

Statistical analysis includes paired t-tests to evaluate primary and exploratory outcomes in participants who completed the study.

Results Summary

Enrolled participants: 10

Participants with data for analysis: 7

Outcomes are presented as baseline vs. 12-week measures

Total AHI: 15.5±5.1 vs. 12.2±5.5 events/hour (p=0.11)

Supine AHI: 21.0±10.8 vs. 15.3±6.0 events/hour (p=0.048)

Non-supine AHI: 7.3±7.4 vs. 6.8±6.7 events/hour (p=0.064)

Maximal inspiratory pressure: 93±35 vs. 114±34 kPa (p=0.02)

Maximal expiratory pressure: 108±33 vs. 137±14 kPa (p=0.04)

Anterior tongue strength: 53±13 vs. 56±10 kPa (p=0.20)