

European Respiratory Society Annual Congress 2013

Abstract Number: 3313

Publication Number: P4001

Abstract Group: 4.2. Sleep and Control of Breathing

Keyword 1: Sleep disorders **Keyword 2:** Monitoring **Keyword 3:** Quality of life

Title: Efficacy of an oral appliance for the treatment of obstructive sleep apnea in a Brazilian population

Prof. Dr Lilian 19865 Giannasi odontogiannasi@uol.com.br¹, Prof. Dr Fernanda 19866 Almeida falmeida@usp.br², Dr. Sergio 19870 Nacif pro_ar@uol.com.br MD³, Dr. Israel 19873 Santos israel.santos@uninove.br³, Mr. Ismael 19877 Dias ismaelsousa@gmail.com³, Prof. Dr Fernando 19879 Leitao Filho fernandostudart@uol.com.br MD⁴, Dr. Renato 19891 Pasqual remarrachp@gmail.com MD³, Prof. Dr Luciana 26653 Sampaio lucianamalosa@uninove.br³, Prof. Dr Rodolfo 26668 Vieira rodolfo.vieira@uninove.br³ and Prof. Dr Luis 26710 Oliveira oliveira.lvf@pq.cnpq.br³. ¹ Dental School - UNESP, Bioscience Department - UNESP University, Sao Paulo, SP, Brazil, 007-0009 ; ² Dental School, University of British Columbia, Vancouver, BC, Canada, 00786 ; ³ Rehabilitation Sciences Doctoral Degree Pos Graduation Program, Nove De Julho University - UNINOVE, Sao Paulo, SP, Brazil, 05006-000 and ⁴ Medicine School, Fortaleza University - UNIFOR, Fortaleza, CE, Brazil, 0070098 .

Body: Aim: the aim of this prospective study was to validate, in Brazil, the use of an MRA to treat OSA and primary snoring, comparing polysomnographic and Epworth Sleepiness Scale (ESS) data obtained prior to and during MAS treatment. Method: this study was carried out on 63 patients presented with different OSA severity or primary snoring, who were fitted to PMPositioner® between 2009 and 2011. The diagnosis was established by a polysomnogram (PSG) prior treatment and a PSG after 6 month verified the efficacy of MRA therapy. Subjective daytime sleepiness was evaluated by ESS questionnaire prior to treatment and at the follow up. Results: The sample was divided in primary snoring and OSA group. For the primary snoring group, PSG variables did not show significant results, except for snoring decreasing. For the OSA group the mean AHI has reduced from 23.0±11 to 5.3±4.0 (p≤0.001) and median ESS reduced significantly from 13.0(3-24) to 8.5(3-13). Complete response (AHI<5.0) was found in 25 (40%) of the patients; partial response (AHI ≤10) was found in 27 (43%) patients. Conclusion: The findings of the present study validate the efficacy of the adjustable PM Positioner® for the treatment of OSA in Brazilian patients. This appliance can provide safe treatment for OSA.