Does successful treatment of Urinary Urgency or Sleep Disordered Breathing improve NOCTURIA and centrally-driven comorbidities?

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INTRODUCTION

Nocturia is more than an isolated lower urinary tract symptom, being significantly associated with dysfunction of sleep quality and duration, cardiovascular morbidity, mental health and mortality [1]. The medical conditions co-existing with nocturia may share central neural control areas in the brainstem [2,3]. The aim of this study was to investigate whether improvement in one comorbid variable, in patients with nocturia, may regulate other co-morbid dysfunctions toward a more normal state.



METHODS

A prospective, 2-arm repeated measures study was performed in 2017. Participants were recruited from both Continence and Sleep Medicine Services.

Inclusion criteria:

- were ≥40 years of age
- experienced nocturia of ≥1 per night
- reported urinary urgency/urge incontinence (UUI) severe enough to require pharmacotherapy <u>OR</u> sleep disordered breathing (SDB) with an apnoeahypopnoea index ≥30 and requiring CPAP

Data collected included:

- · Demographic information
- Overactive Bladder Symptom Score (OABSS)
- Nocturia-related Quality of Life instrument (NQoL)
- Epworth Sleepiness Scale (ESS)
- · Pittsburg Sleep Quality Index (PSQI)
- Hospital Anxiety and Depression Scale (HADS)
- EuroQol Health Questionnaire (EQ-5D-5L)
- · 2-day bladder diary
- Actigraphy parameters
- Blood pressure

The study intervention was either an anticholinergic agent or beta-3 agonist to treat urgency/urgency incontinence or CPAP to reduce apnoea episodes. SDB participants were not treated for any urinary tract symptoms; UUI participants did not commence CPAP therapy during the study.

REFERENCES

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RESULTS

	Continence (pre → post)	Sig (p)	Sleep (pre → post)	Sig (p)
LUTS				
OAB	8.8 (3.0) -		5.5 (3.3) -	
Symptom	6.8 (3.1)	0.004	4.0 (3.4)	0.009
Score	0.0 (3.1)		1.0 (3.1)	
Nocturia	2 (1.0-3.0) -		1 (1.0-2.4) -	
Frequency	1 (1.0-2.0)*	0.004	1 (0.4-2.0)*	0.065
(from FVC)	1 (1.0 2.0)		1 (0.1 2.0)	
Avg Total	697 (294) -		716 (271) -	
Nocturnal Urine	573 (276)	0.089	384 (163)	0.002
(mL)	373 (273)		30 : (200)	
Average	237 (103) -		287 (112) -	
Nocturnal Vol	273 (129)	0.087	266 (165)	0.641
Voided (mL)	273 (223)			
Nocturnal	38.6 (13.0) -		41.1 (16.2) -	
Polyuria	36.3 (14.9)	0.628	30.4 (15.9)	0.058
Index (%)	30.3 (14.3)		30.4 (13.3)	
WELLBEING				
Nocturia	40.2/12=1		24 5 (42 1)	
QoL	19.2 (10.7) -	0.120	21.5 (12.4) -	0.007
Score	15.4 (9.4)		11.9 (11.0)	
HADS	F /0 46'		7 (2.0)	
Anxiety	5 (3-10) -	0.294	7 (2-9) -	0.929
Score	3 (1-10)*		6 (1-9)*	
HADS			- ()	
Depression	4 (1-9) -	0.428	5 (2-7) -	0.858
Score	4 (2-7)*		6 (1-9)*	
CARDIOVASCULAR				
Systolic BP				
(mmHg)	129 (25) -	0.888	137 (13) -	0.069
All patients	130 (17)	0.000	143 (14)	0.003
Systolic BP				
(mmHg)	167 (17) -	0.035	150(5) -	0.318
Baseline >140	140 (8)	0.033	156(8)	0.510
Diastolic BP				
(mmHg)	65 (59-75) -	0.069	85(76-88) -	0.051
l ' ",	74 (63-80)*	0.003	87 (81-97)*	0.031
All patients SLEEP				
First	3 (1.2) -	0.00-	2.8 (1.6) -	0.001
Uninterrupt.	4 (1.6)	0.005	3.9 (2.4)	0.061
Sleep Time				
Sleep	1 (1.0-3.0) -	0.115	1 (0.3-2.0) -	0.537
Latency (mins)	1 (0.0-2.0)*	0.115	1 (0.0-2.0)*	0.527
All patients				
Sleep Latency	3 (2.0-3.0) -	0.004	2 (2.0-3.0) -	0.45-
(mins) 1<30, 2 is	2 (1.0-2.8)*	0.034	2 (1.5-2.5)*	0.157
30-60; 3 is >60			-	
PSQI	8.6 (4.2) -	0.00-	8.7 (4.0) -	0.000
Global	6.7 (3.0)	0.027	6.8 (4.7)	0.036
Score			-	
PSQI	87.3 (13.5) -		82.5(7.9) -	
Habitual Sleep	94.6 (3.2)	0.454	91.6(3.1)	<0.001
Efficiency (%)	` ′		` ′	
PSQI	1 (1.0-2.0) -		1 (1.0-2.0) -	
Daytime	1 (1.0-2.0)*	1.000	1 (0.0-1.0)*	0.034
Function	, , , , , ,		, , , , , ,	
NOTE: results are presented as mean (SD); otherwise as * median (IQR).				

NOTE: results are presented as mean (SD); otherwise as * median (IQR). LUTS = Lower Urinary Tract Symptoms; BP = Blood pressure.

CONCLUSIONS

In a sample of relatively healthy individuals, treatment targeted to the presenting dysfunction resulted in less nocturia, improved sleep quality and longer duration of undisturbed sleep. This early work appears to suggest that in both groups treatment may induce change toward a more normal state in selected variables known to have control areas in the brainstem.