

#97 The Impact of Frailty on the Treatment of Overactive Bladder in Older Adults

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Introduction

- Frailty is a measure of physiologic vulnerability that manifests as increased susceptibility to adverse events such as falls, disability, loss of independence, increased risk of postoperative complications, and death.¹⁻⁴
- The Timed Up and Go Test (TUGT) is a sensitive, specific, and efficient measure of frailty that has a strong independent correlation with poor surgical outcomes.^{5,6}
- The impact of frailty on treatment of overactive bladder (OAB), however, is poorly understood.
- The aim of this study is to examine the impact of frailty on treatment outcomes for OAB in older adults starting pharmacotherapy, onabotulinumtoxinA, and sacral neuromodulation.

Methods

- This is a prospective study of men and women ≥ 60 years of age starting pharmacotherapy, onabotulinumtoxinA, or sacral neuromodulation.
- Subjects were administered questionnaires at baseline, 1- and 3-months.
- Covariates:** Frailty was assessed at baseline using the TUGT, whereby a TUGT time of ≥ 12 seconds was considered to be slow, or frail.
- Outcomes:** Response to treatment was assessed using the overactive bladder symptom score (OABSS) and the Overactive Bladder Questionnaire Short Form (OAB-q SF) Bother and Health Related Quality of Life (HRQOL) subscales. Side effects were also assessed via questionnaire.
- Analyses:** Mixed effects linear modeling was used to model changes in OAB questionnaires over time both within and between groups (frail vs. non-frail).

Results

Figure 1. CONSORT diagram depicting study enrollment.

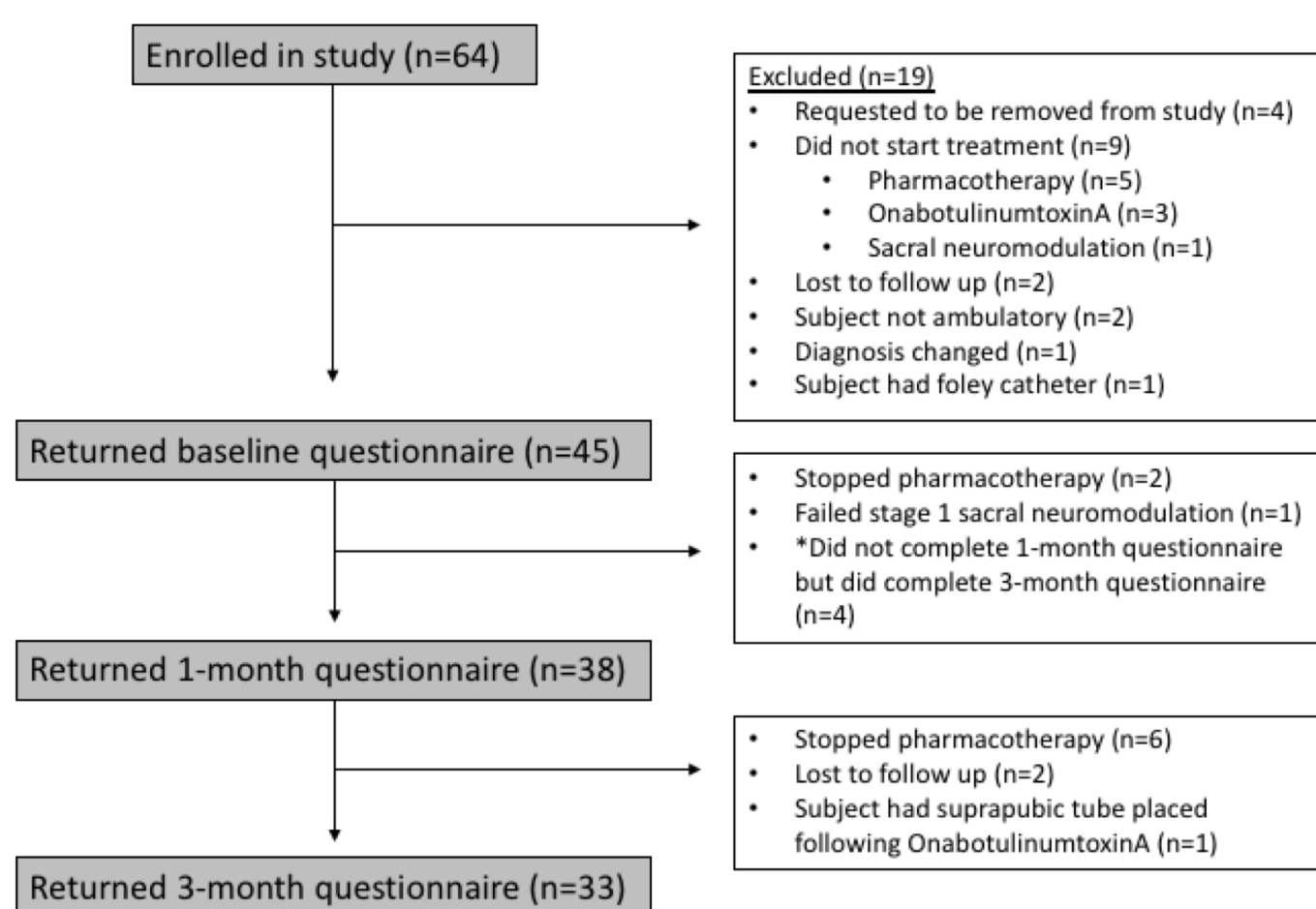


Figure 2. Questionnaire-based treatment responses to OAB treatments stratified by TUGT <12 and TUGT ≥ 12 seconds. Panel A is the OABSS, B is OAB-q SF and C is OAB-q Bother. All figures are adjusted for age and neurogenic bladder.

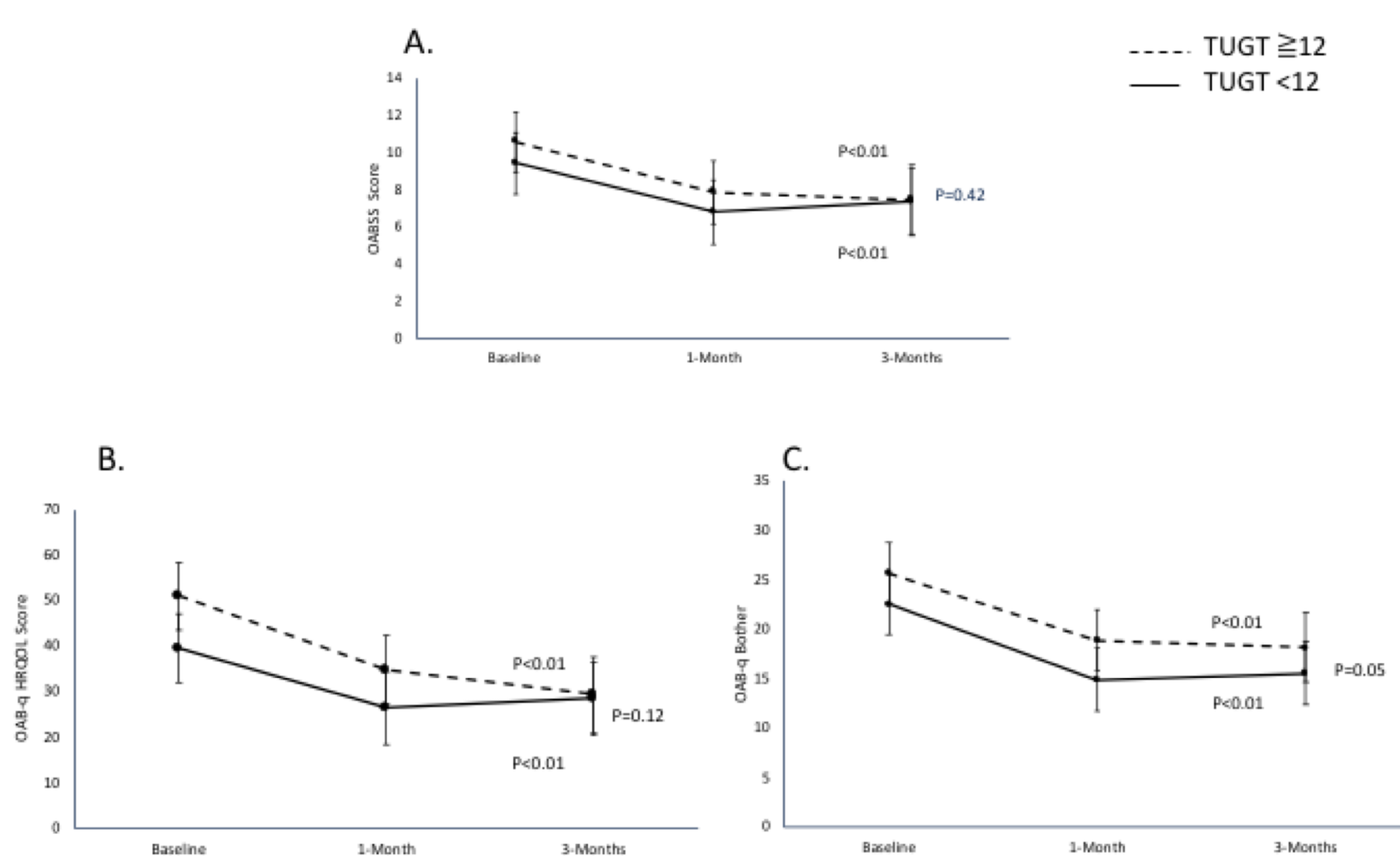


Table 3. Side effects at 1- and 3-months. All p values comparing 1- and 3-month values were >0.05 and are not presented.

Side effect	Overall N=38	1 month		P value	Overall N=32	3 months		P value
		TUG <12 N=21	TUG ≥ 12 N=17			TUG <12 N=19	TUG ≥ 12 N=13	
Any side effect	28 (73.7)	17 (81)	11 (64.7)	0.26	27 (81.8)	17 (85.0)	10 (76.9)	0.56
Headache	5 (13.2)	1 (4.8)	4 (23.5)	0.09	6 (18.2)	3 (15.0)	3 (23.1)	0.56
Dry mouth	18 (47.4)	8 (38.1)	10 (58.8)	0.20	17(51.5)	9 (45.0)	8 (61.5)	0.35
Constipation	13 (34.2)	7 (33.3)	6 (35.3)	0.90	16 (48.5)	8 (40.0)	8 (61.5)	0.23
UTI	3 (7.9)	1 (4.8)	2 (11.8)	0.43	3 (9.1)	1 (5.0)	2 (15.4)	0.31
Nausea	5 (13.2)	1 (4.8)	4 (23.5)	0.09	6 (18.2)	3 (15.0)	3 (23.1)	0.56
Urinary retention	8 (21.1)	5 (23.8)	3 (17.6)	0.64	4 (12.1)	2 (10.0)	2 (15.4)	0.64
Fatigue	12 (32.4)	6 (28.6)	6 (37.5)	0.57	14 (42.4)	10 (50.0)	4 (30.8)	0.27
Confusion	7 (18.4)	3 (14.2)	4 (23.5)	0.46	5 (15.6)	2 (10.5)	3 (23.1)	0.34
Pain	6 (15.8)	2 (9.5)	4 (23.5)	0.24	9 (27.3)	6 (30.0)	3 (23.1)	0.66

Table 1. Baseline clinical characteristics, questionnaires, prior OAB bladder treatments stratified by TUGT <12 and TUGT ≥ 12 seconds.

Characteristic	Overall N=45	TUG <12 N=27	TUG ≥ 12 N=18	P value
Age, mean \pm SD	70.3 \pm 6.3	69.7 \pm 5.3	71.2 \pm 7.6	0.43
Female gender, n (%)	31 (68.9%)	18 (66.7%)	13 (72.2%)	0.69
BMI, mean \pm SD	31.7 \pm 12.5	31.4 \pm 15.5	32.1 \pm 6.7	0.85
Race/ethnicity, n (%)				0.26
White	34 (75.6)	22 (81.5)	12 (66.7)	
Nonwhite	11 (24.4)	5 (18.5)	6 (33.3)	
Highest level of education, n (%)				0.17
Less than high school	4 (8.9)	1 (3.7)	3 (16.7)	
High school graduate or equivalent	1 (2.2)	0	1 (5.6)	
Bachelors' degree	16 (35.6)	12 (44.4)	4 (22.2)	
Graduate degree	24 (53.3)	14 (51.9)	10 (55.6)	
Total household income in the last year				0.16
<25 K	3 (6.7)	1 (3.7)	2 (11.1)	
25-49 K	7 (15.6)	2 (7.4)	5 (27.8)	
50-99 K	7 (15.6)	4 (14.8)	3 (16.7)	
100-149 K	5 (11.1)	4 (14.8)	1 (5.6)	
≥ 150 K	5 (11.1)	5 (18.5)	0	
Prefer not to say	18 (40.0)	11 (40.7)	7 (38.9)	
Neurogenic bladder, n (%)	8 (17.8%)	2 (7.4%)	6 (33.3%)	0.03
Multiple sclerosis	2 (4.4%)	0	2 (11.1%)	
Parkinson's Disease	3 (4.4%)	2 (7.4%)	1 (5.6%)	0.04
Other	3 (6.7%)	0	3 (16.7%)	
Number of medications currently taking, mean \pm SD	12.7 \pm 6.0	12.2 \pm 7.0	13.6 \pm 4.0	0.46
Baseline PVR, mean \pm SD	66.6 \pm 6.0	74.9 \pm 87.3	53.0 \pm 56.5	0.38
Baseline TUGT, mean \pm SD	12.3 \pm 6.9	8.9 \pm 1.5	17.4 \pm 8.5	<0.01
Baseline Animal fluency test, mean \pm SD	18.4 \pm 6.0	19.6 \pm 5.5	16.7 \pm 6.5	0.12
Baseline Katz ADL, mean \pm SD	5.3 \pm 0.6	5.5 \pm 0.5	5.0 \pm 0.6	0.01
Baseline Lawton IADL, mean \pm SD	7.2 \pm 1.4	7.6 \pm 1.2	6.6 \pm 1.5	0.04
Baseline general health rating, n (%)				0.05
Excellent	5 (11.1)	3 (11.2)	2 (11.1)	
Very good	9 (20.0)	9 (33.3)	0	
Good	20 (44.4)	11 (40.7)	9 (50.0)	
Fair	10 (22.2)	4 (14.8)	6 (33.3)	
Poor	1 (2.2)	0	1 (5.6)	
Previous treatments tried, n (%)	28 (62.2)	21 (60.0)	7 (70.0)	0.57
Pelvic floor physical therapy (PFPT)	19 (43.2)	12 (44.4)	7 (41.2)	0.83
Medications	28 (62.2)	16 (59.3)	12 (66.7)	0.62
Percutaneous tibial nerve stimulation (PTNS)	4 (8.9)	3 (11.1)	1 (5.6)	0.52
OnabotulinumtoxinA	3 (6.7)	2 (7.4)	1 (5.6)	0.81
Sacral neuromodulation	3 (6.7)	2 (7.4)	1 (5.6)	0.33

Table 2. Type of OAB treatment selected during the study.

Treatment selected, n (%)	Overall N=45	TUG <12 N=27	TUG ≥ 12 N=18	P value
Medication	22 (48.9)	13 (48.1)	9 (50.0)	0.96
OnabotulinumtoxinA	12 (26.7)	7 (25.9)	5 (27.8)	
Sacral neuromodulation	11 (24.4)	7 (25.9)	4 (22.2)	

Conclusions

- Adults ≥ 60 years of age starting second- and third-line treatments for OAB, regardless of TUGT time, demonstrated improvement in OAB symptoms at 3 months.
- These findings suggest that frail older adults may receive comparable benefit and similar rates of side effects compared to less frail individuals.
- Further studies are needed to explore and confirm these findings.

References

- Buchner DM, Wagner EH. Preventing frail health. *Clinics in geriatric medicine*. 1992;8(1):1-17.
- Fried LP, Tangen CM, Walston J, et al. Frailty in older adults: evidence for a phenotype. *The journals of gerontology Series A, Biological sciences and medical sciences*. 2001;56(3):M146-156.
- Makary MA, Segev DL, Pronovost PJ, et al. Frailty as a predictor of surgical outcomes in older patients. *Journal of the American College of Surgeons*. 2010;210(6):901-908.
- Suskind AM, Walter LC, Jin C, et al. Impact of frailty on complications in patients undergoing common urological procedures: a study from the American College of Surgeons National Surgical Quality Improvement database. *BJU international*. 2016;117(5):836-842.
- Robinson TN, Wu DS, Sauaia A, et al. Slower walking speed forecasts increased postoperative morbidity and 1-year mortality across surgical specialties. *Annals of surgery*. 2013;258(4):582-588; discussion 588-590.
- Savva GM, Donoghue OA, Horgan F, O'Regan C, Cronin H, Kenny RA. Using timed up-and-go to identify frail members of the older population. *The journals of gerontology Series A, Biological sciences and medical sciences*. 2013;68(4):441-446.