Bowel Function, Sexual Function, and Symptoms of Pelvic Organ Prolapse in Women with and Without Urinary Incontinence

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Abstract

Aims—Bowel symptoms, pelvic organ prolapse, and sexual dysfunction are common, but their frequency among women with lower urinary tract symptoms (LUTS) has not been well described. Our aims were to describe pelvic floor symptoms among women with and without urinary incontinence (UI) and among subtypes of UI.

Methods—Women with LUTS seeking care at six U.S. tertiary care centers enrolled in prospective cohort study were studied. At baseline, participants completed the Pelvic Floor Distress Inventory (PFDI-20), Pelvic Organ Prolapse/Incontinence Sexual Questionnaire (PISQ-IR), and PROMIS GI Diarrhea, Constipation, and Fecal Incontinence Scales.

Results—Mean age among the 510 women was 56.4 ± 14.4 years. Women who reported UI (n=420) had more diarrhea and constipation symptoms (mean scores 49.5 vs 46.2 [p=0.01] and

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51.9 vs 48.4 [p<0.01], respectively) at baseline. Among sexually active women, mean PISQ-IR subscale scores were lower among those with UI (condition specific: 89.8 vs 96.7, p<0.01; condition impact: 79.8 vs 92.5, p<0.01). Women with mixed urinary incontinence (MUI) (n=240) reported more prolapse symptoms, fecal incontinence, and worse sexual function compared to those with stress urinary incontinence (SUI) and urgency urinary incontinence (UUI).

Conclusions—Women presenting with LUTS with UI reported significantly worse constipation, diarrhea, fecal incontinence, and sexual function compared to women without UI. In women with UI, sexual function and pelvic organ prolapse (POP) symptoms were worse in those with MUI compared to SUI and UUI.

Introduction

Symptoms of pelvic-floor disorders including constipation, diarrhea, and fecal incontinence as well as pelvic organ prolapse and sexual dysfunction are common in women. These disorders occur concurrently with urinary incontinence (UI); however, the relationship has not been well described. Pelvic floor dysfunction is common to all of these disorders which is why aging and childbirth which both affect the pelvic floor are factors that concurrently affect multiple pelvic floor organs in adult women. It has been reported that women with difficult defecation have more urinary urgency and frequency, but not UI; however, this is controversial[1]. Women with obstetrical anal injury are at increased risk not only for fecal incontinence but also stress urinary incontinence (SUI)[2].

Pelvic organ prolapse has consistently been associated with urinary urgency and urgency incontinence (UUI), with the relationship possibly being causal since correction of the prolapse or placement of a pessary can relieve these bladder symptoms[3]. Also, the relationship between prolapse and SUI is complex, as SUI often occurs concurrently with prolapse, but prolapse may also be protective as correction of prolapse often unmasks occult SUI[4].

The association between UI and sexual activity remains uncertain. Prior studies have provided conflicting results, with patient age and partner status as possible significant factors[5,6]. Mixed urinary incontinence (MUI) has the greatest negative impact on sexual function compared to SUI and UUI[7].

The aims of this study were: 1) to determine the relationships between bowel symptoms including constipation, diarrhea, and fecal incontinence, prolapse symptoms, and sexual function among women seeking care for lower urinary tract symptoms (LUTS); and 2) to evaluate whether the presence of UI, or UI subtype, is associated with the severity of these symptoms.

Material and Methods

Study Design and Population

We report on women enrolled in a one-year, multi-center, prospective observational cohort study from the NIH/NIDDK-sponsored Symptoms of Lower Urinary Tract Dysfunction Research Network (LURN). Details of this cohort study have been previously reported [8]. Briefly, participants were at least 18 years of age, presented to a LURN physician for the first time seeking care for their LUTS, and reported at least one LUTS Tool[9] symptom using a one-month recall screening period. We modified the LUTS Tool, with permission from Pfizer, to capture a recall period of one month for the LURN study. Data collection at the baseline visit for women included a standardized clinical examination including pelvic examination with Pelvic Organ Prolapse Quantitation (POP-Q), assessment of pelvic floor muscle strength with the Oxford scale, urinalysis, and measurement of post-void residual. Medical history including functional comorbidity index (FCI), patient-reported symptoms of LUTS, pelvic floor symptoms, and psychological symptoms was also collected. Quality of life was obtained by validated questionnaires [10–13].

Measures

Seven questions regarding UI on the LUTS Tool [9] were used to determine presence of UI. Women who reported "rarely" or "never" to these questions were classified as "without UI", and responses of "sometimes" or greater on at least one symptom of UI during exercise, laughing/sneezing/coughing, feelings of urinary urgency, sleep, sex, or for no reason were classified as "with UI." Participants were further classified as having SUI if they answered "sometimes" or more on at least one of two questions related to experiencing leakage while exercising or during a laugh, cough, or sneeze. Those who responded "sometimes" or more to leakage due to a sudden feeling of needing to rush to urinate were classified as having UUI. Those with both SUI and UUI were classified as MUI. Those participants with UI who did not meet criteria for SUI, UUI, or MUI were classified as Other UI.

A continuous UI severity measure was also calculated using the 7 LUTS Tool UI questions. For each study participant, the weighted Euclidean length (square root of sum of squared responses) was calculated to form a UI severity score (range 1.84–9.44). Questions were weighted by the ratio of the average correlation between a given question and all other questions to the total average correlation so that less weight was given to questions that had high correlation with other questions (e.g., multiple questions assessing SUI)[14].

Participants also completed the Pelvic Organ Prolapse/Incontinence Sexual Questionnaire, IUGA Revised (PISQ-IR)[10], Genitourinary Pain Index (GUPI)[11], Pelvic Floor Distress Inventory-20 (PFDI-20) [12], and Patient-Reported Outcomes Measurement Information System (PROMIS) GI Constipation, Diarrhea, and Fecal Incontinence scales[13]. The PFDI-20 is a condition-specific quality of life measure that assesses bother related to pelvic floor symptoms and includes three scales, the Urinary Distress Inventory (UDI-6), Pelvic Organ Prolapse Distress Inventory (POPDI-6), and Colo-Rectal-Anal Distress Inventory (CRADI-8). Each scale is scored 0–100 with a higher score indicating greater bother. The PISQ-IR measures sexual function in women with pelvic floor disorders separately for

sexually and not sexually active women. A higher score on the subscale indicates better sexual function. There is no summary score for the PISQ-IR.

PROMIS measures used short forms to derive T-scores normalized to the U.S. population as a reference (by definition, mean=50, standard deviation [SD] = 10). One exception was fecal incontinence, which uses a raw score as the metric. Higher scores on PROMIS measures indicate more symptoms.

Statistical Methods

Characteristics of the participants are shown as means, standard deviations, frequencies, and percentages. Tests for differences by group were performed using chi-square tests and Wilcoxon two-sample tests or Kruskal-Wallis tests. Descriptive statistics were calculated for sexual function, pelvic floor, and bowel symptom measures; differences between groups were tested using one-way ANOVA and Cohen's *d* was used to calculate effect sizes. Urinary subscales (GUPI urinary subscale and UDI-6 subscale of PFDI-20) and summary scores including urinary subscales were excluded due to similarity to LUTS Tool questions.

Multivariable linear regression was used to test for associations between incontinence status (UI vs. non-UI) and sexual function, pelvic floor, and bowel measures. Candidate covariates included age, race, ethnicity, body mass index (BMI), education, employment status, smoking status, diabetes, sleep apnea, functional comorbidity index (FCI), menopausal status (with and without hormone use), history of psychiatric diagnosis, previous brain or spinal surgery, more than two urinary tract infections in the past year by self-report, hysterectomy, any vaginal births, and alcohol consumption. Best subset selection guided covariate selection for all models. Similar models were created to test for associations between the outcomes and UI subtype (SUI, UUI, and MUI, with MUI as the reference category due to its prevalence and increased severity) and UI severity. Data on POP-Q and pelvic floor strength (Oxford scale) were each missing in 20% of participants and were excluded as potential covariates; however, they were tested in separate sub-analyses. All p-values were adjusted for multiple comparisons using the false discovery rate (FDR) correction. All analyses were conducted using SAS 9.4 (SAS Institute, Cary, NC).

Results

Among 545 women recruited from the six sites, 510 with complete responses to the 7 LUTS Tool questions required for UI subtyping were included in the present analyses. Mean age was 56.4 ±14.4 years; most were Caucasian (82%) (Table 1). Mean BMI was 30.6 ±7.8 kg/m^2, and 15% reported a history of diabetes. A median of two vaginal births was reported. Sixty-four percent were post-menopausal and 17% used estrogen treatment (topical or systemic). Sixty-three percent had a stage 0 or 1 pelvic organ prolapse on physical exam, 30% had stage 2, and 6% had stage 3 or 4. At baseline, few study participants reported taking an anti-muscarinic drug (2%) or medication to relieve constipation (6%). Thirty percent had prior hysterectomy, and 14% had undergone surgery for UI and/or prolapse. One-half of the women were sexually active (51%). The mean functional comorbidity index was 2.4 ±2.2 for the group, and mean post-void residual was 44.8 ±58.7 ml.

Overall, 90 (18%) women reported no more than rarely having incontinence on any of the seven incontinence questions (16a-g) on the LUTS tool and were considered "without UI." Compared to the 420 women with UI, those without UI had lower mean BMI (28.2 vs. 31.1, p=0.002), were less likely to have recurrent urinary tract infections in the prior year (37% vs. 50%, p=0.029), and had fewer self-reported comorbidities (FCI 1.9 vs. 2.5, p=0.012). Of the 420 women with UI, most had MUI (57%), 20% had UUI, and 17% SUI. Six percent (n=25) had Other UI. Women reporting MUI were older, with higher BMI, higher prevalence of smoking and sleep apnea, and more comorbidities compared to women with UUI or SUI only. Women with MUI had significantly higher UI severity compared to women with SUI or UUI (Figure 1). Average UI severity in the MUI group was 5.39±1.54 compared to 3.98±1.48 (SUI) and 3.31±1.13 (UUI) (all p<0.001).

Associations between sexual functioning and UI

In terms of sexual dysfunction, only a few of the PISQ subscales were significantly different between groups. Among sexually active women only, the PISQ-SA Condition Impact and Condition Specific subscales were lower (worse function) in women with UI (mean scores 79.8 [UI] vs. 92.5 [non-UI], p<0.001 and 89.8 [UI] vs. 96.7 [non-UI], p=0<0.001) compared to women without UI, and these differences remained significant after adjusting for BMI, smoking status, and parity.

Among women with UI who reported they were sexually active, those with MUI reported lower mean scores on PISQ SA-Condition Impact subscale (worse function) (average 73.3) compared to women with SUI (average 86.3) and women with UUI (average 87.4). These differences remained significant after covariate adjustment (Supplemental Table 1). Urinary incontinence occurring during sexual intercourse was more common in the SUI and MUI groups, with 17% and 18%, respectively, compared to only 4% in the UUI group (p=0.004). Women who reported more severe UI, regardless of subtype, had significantly worse sexual function on the PISQ SA-Condition Specific (on average 3.05 point reduction in PISQ score per unit increase in UI severity, p<0.001) and Condition Impact measures (on average 2.64 point reduction in PISQ score per unit increase in UI severity, p=0.01).

Associations between prolapse symptoms and UI

Bother associated with prolapse was marginally higher in women with UI (average POPDI-6 scores 17.1 vs. 13.8, p=0.10, Table 2), although measures of anterior and posterior vaginal wall descent did not differ between the two groups. Among those with UI, women with MUI reported more bother associated with prolapse (average POPDI-6 scores 20.6 compared to 10.7 [UUI] and 12.4 [SUI], p<0.001, Table 3), and these differences remained statistically significant after adjustment. In all women, as UI severity increased, women reported worse pelvic floor distress on the POPDI-6.

Associations between bowel function and UI

Compared to those without UI, women with UI had higher PROMIS Constipation (51.9 vs. 48.4, p=0.003), and Diarrhea scores (49.5 vs. 46.2, p=0.008), indicating worse bowel function (Table 2). On multivariable linear regression of these bowel function measures adjusted for other statistically significant covariates (Supplementary Table 2), these results

were only minimally changed. CRADI-8 scores were also higher for women with UI (21.0 vs. 14.8, p=0.02), but this difference did not remain statistically significant after adjustment for comorbidities and parity.

Among those with UI, the MUI group experienced more bowel symptoms and had higher (worse) PROMIS scores for diarrhea and fecal incontinence (Table 1). There were no significant differences in PROMIS Constipation scores among the various UI subtypes with MUI as the reference category. CRADI-8 scores were also higher (more bother) among the MUI group compared with SUI and UUI groups. After adjustment, the differences remained between the MUI and SUI groups, but not between MUI and UUI. As UI severity increased, bowel function worsened on all three PROMIS bowel measures; CRADI-8 scores measuring bother related to bowel function also increased. Full models are available in Supplemental Table 3.

Discussion

We report the relationships between UI and other pelvic floor symptoms and quality of life measures, including bowel, prolapse, and sexual function, in over 500 women seeking care for LUTS. Overall, our results show that in women with LUTS: (1) the presence of UI is associated with constipation and poor sexual function; (2) MUI is associated with worse fecal incontinence, diarrhea, pelvic organ prolapse symptoms, and sexual function compared to SUI; and (3) more severe UI symptoms, regardless of UI subtype, are associated with worse bowel function (fecal incontinence, diarrhea, constipation), pelvic organ prolapse symptoms, and sexual function.

Although it is well known that UI adversely affects sexual function in women, less is known about the effects of UI subtypes. Conflicting results were reported in three studies, which identified UI subtypes by urodynamic testing and assessed sexual function (in sexually active women) through PISQ-12 scores[6,7,15]. Two of these studies found SUI patients had worse sexual function than UUI patients[6,7], and one found MUI patients had the poorest sexual function[7]. The third study found no difference in sexual function between all three UI subtypes [16]. All of these studies were limited by small sample sizes and their results were not adjusted for potentially important covariates, such as age, BMI, and comorbidities. In contrast to these prior results, we found poorer sexual function in women with MUI among sexually active women, but no large differences in function between stress and urgency UI subtypes using a much larger sample of women and multivariable analysis.

The association between pelvic organ prolapse and UI has pathophysiological basis. SUI commonly occurs with pelvic organ prolapse, due to similar pelvic floor injury causing urethral hypermobility and/or some degree of intrinsic sphincter deficiency. UUI may also have a strong relationship to pelvic organ prolapse[17] since POP may cause bladder outlet obstruction and overactive bladder symptoms. A large cystocele may also put traction on the urethra, resulting in an open urethra. Surgical repair of pelvic organ prolapse has been shown to improve UUI in the majority of patients[3]. Levator ani pelvic floor muscle injury, sphincteric injury, and/or pudendal nerve injury may be also present in a subset of patients who have concomitant UI, pelvic organ prolapse, and fecal incontinence[218,19]. Thus, our

finding that increasing UI severity was associated with more distress from prolapse symptoms was not surprising.

Regarding bowel function, past studies have similarly demonstrated that constipation, difficult defecation, and fecal incontinence occur commonly in women with LUTS[20–22]. Studies performed in specialty clinic populations have demonstrated associations between functional constipation and overactive bladder[22], and a high prevalence (19%) of fecal incontinence in women seeking care for urinary incontinence[20]. However, there have only been a few studies that have examined the effects of UI subtypes on bowel dysfunction. Meschia et al. suggested that anal incontinence is more prevalent among patients with MUI and UUI than SUI (28.8%, 28.7%, and 21.8%, respectively)[23]. We have also demonstrated increased bowel dysfunction (fecal incontinence and diarrhea) in women with mixed UI, compared to SUI. Our results strengthen these prior findings with the use of validated PROMIS questionnaires (rather than a non-validated screening questionnaire) to assess the bowel symptoms and multivariable analysis to adjust for potential confounding variables.

There are several theoretical explanations for the association between bowel and bladder dysfunction. Both the bladder and bowel originate embryologically from the same cloaca and, given the proximity of the bowel and bladder in the pelvis, a distended rectal vault could have a mass effect on the bladder. Both the distal bowel and bladder share afferent nerves, as well, explaining why sacral neuromodulation is used to treat both bowel and bladder incontinence. Studies on the treatments of one organ resulting in a positive impact on other pelvic organs are lacking in adults. However, it has been clearly demonstrated that aggressive treatment of constipation in children with dysfunctional elimination without any bladder intervention frequently results in resolution of UI[12]. These theories may explain why women with MUI have worse bowel function since they likely have combined anatomic (loss of support) and neurologic deficits.

Our study has several important strengths. First, we used a condition-specific questionnaire (PISQ-12) to assess sexual function in women with pelvic floor disorders. The questionnaire has undergone validation in this patient population and assesses both sexually active and non-active women[10]. Second, bowel symptoms were assessed using validated PROMIS questionnaires[13]. Finally, unlike many previous studies that were typically from a single center with small sample size, we have recruited a large number of women prospectively across several sites; this may enhance the generalizability of our findings to other care-seeking women in different clinical care settings.

Our study has several limitations. As it entails cross-sectional comparisons, the causal relationship of one symptom to another cannot be inferred. The classification into SUI, UUI, and MUI was based on self-reported symptoms on the LUTS Tool questionnaire, and not urodynamic findings. Also, patients were recruited at tertiary academic centers with expertise in managing LUTS. Thus, our results may be less generalizable to women who seek treatment with community urologists, gynecologists, or primary care physicians. On the other hand, relatively few women reported taking anticholinergic medications (2%) or had had prior pelvic surgery for UI at study entry (14%), suggesting our participants did not include many complex or refractory cases. Finally, the UI severity measure reported here has

not been validated and therefore results regarding UI severity may not be reproducible in other populations.

Conclusion

Among women seeking care for LUTS, those with UI symptoms, mixed UI, and/or more severe UI were more likely to report poorer bowel dysfunction, prolapse symptoms, and worse sexual function. Our findings suggest that health care providers should question their patients seeking care for LUTS to identify and manage co-occurring pelvic floor dysfunctions.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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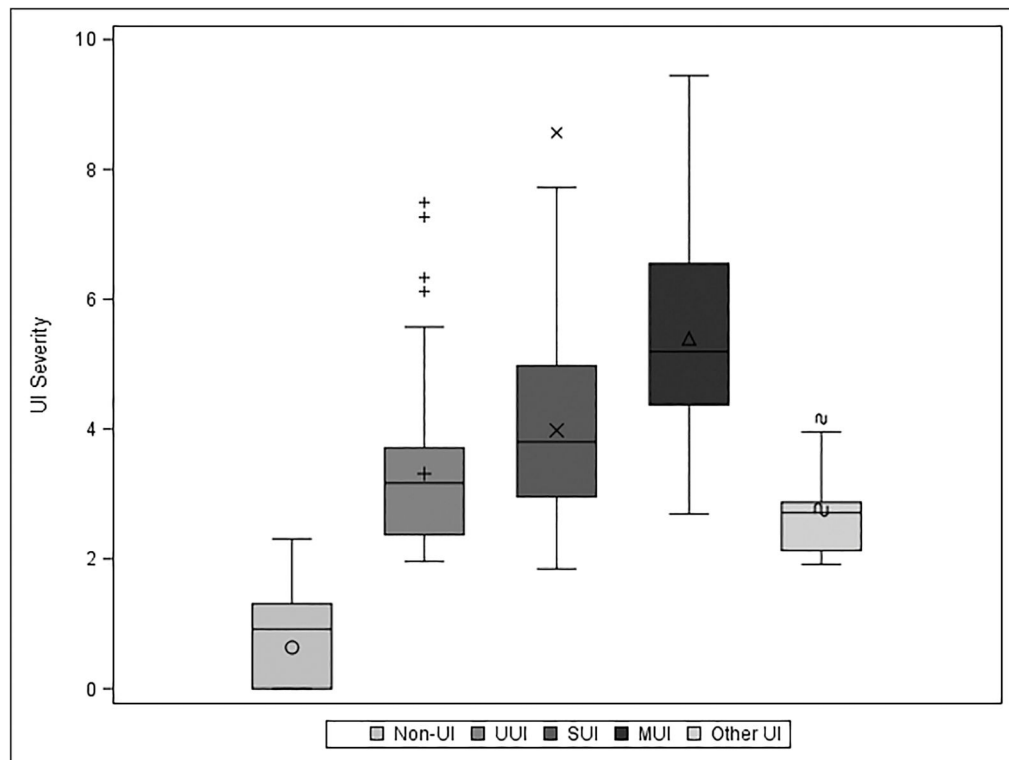
Figure 1.
UI Severity by subtype. UI severity was calculated as the weighted Euclidean distance (square root of sum of squared responses) of 7 LUTS Tool incontinence questions. Weights were calculated using the ratio of average correlation of a given question to the average total correlation of all 7 questions in order to account for potential redundancy in questions.

Table 1

Demographics and medical history of LURN female cohort

	Total (n=510)	Non-UI (n=90)	UI (n=420)	p-value*	SUI (n=70)	UUI (n=85)	Mixed UI (n=240)	p-value**
Age	56.4 (14.4)	55.8 (17.1)	56.6 (13.8)	0.953	56.8 (15.8)	53.0 (13.4)	57.6 (13.1)	0.022
Race				0.448				0.178
American Indian/Alaskan Native	5 (1%)	2 (2%)	3 (1%)		1 (1%)	0 (0%)	1 (0%)	
Asian	14 (3%)	2 (2%)	12 (3%)		3 (4%)	4 (6%)	5 (2%)	
African-American	59 (12%)	8 (9%)	51 (12%)		14 (16%)	6 (9%)	30 (13%)	
Native Hawaiian/Pacific Islander	1 (0%)	0 (0%)	1 (0%)		1 (1%)	0 (0%)	0 (0%)	
White	418 (82%)	74 (82%)	344 (82%)		62 (73%)	59 (84%)	200 (84%)	
Multi-racial/Other	12 (2%)	4 (4%)	8 (2%)		4 (5%)	1 (1%)	3 (1%)	
Education				0.159				0.204
< HS Diploma/GED	11 (2%)	1 (1%)	10 (2%)		1 (1%)	2 (3%)	7 (3%)	
HS Diploma/GED	45 (9%)	4 (5%)	41 (10%)		6 (7%)	5 (7%)	29 (12%)	
Some college/tech school - no degree	117 (23%)	17 (20%)	100 (24%)		19 (22%)	10 (14%)	64 (27%)	
Associate's degree	57 (11%)	7 (8%)	50 (12%)		9 (11%)	11 (16%)	26 (11%)	
Bachelor's degree	153 (30%)	29 (34%)	124 (30%)		27 (32%)	21 (30%)	68 (29%)	
Graduate degree	119 (24%)	28 (33%)	91 (22%)		23 (27%)	21 (30%)	42 (18%)	
Employment				0.145				0.054
Employed part-time	72 (14%)	12 (14%)	60 (14%)		14 (16%)	10 (14%)	35 (15%)	
Employed full-time	197 (39%)	30 (34%)	167 (40%)		32 (38%)	38 (54%)	81 (34%)	
Unemployed (looking for work)	14 (3%)	0 (0%)	14 (3%)		1 (1%)	3 (4%)	10 (4%)	
Not employed (not looking for work)	221 (44%)	46 (52%)	175 (42%)		38 (45%)	19 (27%)	110 (47%)	
BMI	30.6 (7.8)	28.2 (5.9)	31.1 (8.1)	0.002	30.6 (7.9)	27.9 (6.4)	32.4 (8.3)	<.001
< 25	137 (27%)	28 (32%)	109 (26%)	0.015	25 (29%)	30 (43%)	47 (20%)	0.006
25–30	131 (26%)	31 (36%)	100 (24%)		19 (22%)	17 (24%)	55 (23%)	
30–35	108 (22%)	16 (18%)	92 (22%)		18 (21%)	13 (19%)	60 (26%)	.
>35	126 (25%)	12 (14%)	114 (27%)		23 (27%)	10 (14%)	73 (31%)	
Current or Former Smoker	174 (35%)	22 (26%)	152 (37%)	0.050	22 (26%)	23 (33%)	96 (41%)	0.040
Diabetes	74 (15%)	13 (15%)	61 (15%)	0.972	12 (14%)	6 (9%)	39 (16%)	0.257
>2 UTIs in the past year	236 (48%)	32 (37%)	204 (50%)	0.029	34 (41%)	32 (46%)	128 (55%)	0.057

	Total (n=510)	Non-UI (n=90)	UI (n=420)	p-value*	SUI (n=70)	UUI (n=85)	Mixed UI (n=240)	p-value**
Sleep Apnea	90 (18%)	10 (11%)	80 (19%)	0.087	12 (14%)	4 (6%)	59 (25%)	<.001
History of psychiatric diagnosis	218 (43%)	32 (36%)	186 (45%)	0.156	33 (39%)	26 (37%)	116 (49%)	0.104
Previous brain or spinal surgery	37 (7%)	5 (6%)	32 (8%)	0.511	6 (7%)	1 (1%)	23 (10%)	0.071
Number of Vaginal Births	1.7 (1.4)	1.5 (1.5)	1.8 (1.4)	0.148	1.7 (1.4)	1.7 (1.2)	1.8 (1.5)	0.606
Post-menopausal	323 (64%)	52 (59%)	271 (66%)	0.245	59 (69%)	39 (56%)	155 (66%)	0.170
Hormone Use	55 (17%)	11 (21%)	44 (16%)	0.430	8 (14%)	12 (32%)	23 (15%)	0.034
Anticholinergic medication use	11 (2%)	3 (3%)	8 (2%)	0.397	1 (1%)	1 (1%)	5 (2%)	0.838
Anti-Constipation medication use	32 (6%)	8 (9%)	24 (6%)	0.260	5 (6%)	4 (6%)	13 (5%)	0.985
Previous Surgeries (multiple possible)								
Urgency incontinence	6 (1%)	0 (0%)	6 (1%)	0.254	1 (1%)	2 (3%)	3 (1%)	0.600
SUI/Prolapse	66 (13%)	10 (11%)	56 (13%)	0.569	11 (13%)	3 (4%)	38 (16%)	0.042
Hysterectomy	154 (30%)	23 (26%)	131 (31%)	0.328	27 (32%)	21 (30%)	76 (32%)	0.948
Urethral Dilation	3 (1%)	0 (0%)	3 (1%)	0.421	0 (0%)	0 (0%)	2 (1%)	0.523
Other	1 (0%)	0 (0%)	1 (0%)	0.643	0 (0%)	0 (0%)	1 (0%)	0.723
Functional Comorbidity Index	2.4 (2.2)	1.9 (1.8)	2.5 (2.2)	0.012	2.2 (2.0)	1.9 (1.8)	2.8 (2.3)	0.002
Prolapse				<.001				0.397
Stage 0	134 (31%)	34 (44%)	100 (28%)		25 (34%)	16 (27%)	50 (25%)	
Stage 1	140 (32%)	17 (22%)	123 (35%)		27 (37%)	22 (37%)	69 (34%)	
Stage 2	131 (30%)	15 (19%)	116 (33%)		20 (27%)	17 (28%)	74 (37%)	
Stage 3	26 (6%)	11 (14%)	15 (4%)		1 (1%)	5 (8%)	8 (4%)	
Stage 4	1 (0%)	0 (0%)	1 (0%)		0 (0%)	0 (0%)	1 (0%)	
Anterior vaginal wall (Aa, cm)	−1.7 (1.3)	−1.7 (1.5)	−1.7 (1.3)	0.520	−1.8 (1.3)	−1.7 (1.2)	−1.6 (1.3)	0.448
Posterior vaginal wall (Ba, cm)	−1.6 (1.9)	−1.4 (2.3)	−1.7 (1.8)	0.764	−1.8 (1.3)	−1.3 (1.5)	−1.7 (2.0)	0.214
Pelvic Floor Muscle Strength (Oxford Scale)				0.501				0.082
Grade 0	39 (11%)	6 (9%)	33 (11%)		8 (13%)	4 (8%)	18 (11%)	
Grade 1	81 (22%)	15 (22%)	66 (22%)		15 (25%)	12 (23%)	36 (21%)	
Grade 2	84 (23%)	13 (19%)	71 (24%)		13 (21%)	8 (15%)	47 (28%)	
Grade 2	80 (22%)	16 (23%)	64 (21%)		10 (16%)	12 (23%)	38 (22%)	
Grade 4	61 (17%)	11 (16%)	50 (17%)		8 (13%)	13 (25%)	27 (16%)	
Grade 5	24 (7%)	8 (12%)	16 (5%)		7 (11%)	4 (8%)	3 (2%)	
PVR (ml)	44.8 (58.7)	53.3 (68.4)	42.9 (56.2)	0.319	46.2 (63.8)	31.0 (56.1)	43.3 (52.7)	0.064

	Total (n=510)	Non-UI (n=90)	UI (n=420)	p-value[*]	SUI (n=70)	UUI (n=85)	Mixed UI (n=240)	p-value[**]
Alcoholic Beverages Consumed Per Week				0.624				0.580
0–3	334 (68%)	62 (74%)	272 (66%)		58 (69%)	47 (67%)	150 (65%)	
4–7	60 (12%)	8 (10%)	52 (13%)		12 (14%)	11 (16%)	24 (10%)	
8–14	14 (3%)	2 (2%)	12 (3%)		3 (4%)	2 (3%)	7 (3%)	
>14	3 (1%)	1 (1%)	2 (0%)		1 (1%)	0 (0%)	1 (0%)	
Never consumed alcohol	83 (17%)	11 (13%)	72 (18%)		10 (12%)	10 (14%)	49 (21%)	

[*] P-value for wet vs. dry from chi-square test or Wilcoxon 2-sample test

[**] P-value for SUI vs. UUI vs. Mixed UI from chi-square test or Kruskal-Wallis test

Table 2

Pelvic Floor Measures among Women with and without Urinary Incontinence

Indices	N Non-UI	Mean (SD) Non-UI	N UI	Mean (SD) UI	p-value[*]	Adjusted p-value[**]
PISQ NSA-Condition Specific	29	21.9 (21.3)	187	29.1 (30.0)	0.168	
PISQ NSA-Partner Related	31	60.7 (26.7)	192	59.8 (29.7)	0.912	
PISQ NSA-Global Quality	29	38.1 (32.8)	188	48.4 (33.1)	0.168	
PISQ NSA-Condition Impact	30	12.2 (22.3)	188	24.0 (31.7)	0.031	0.11
PISQ SA-Arousal Orgasm	50	66.4 (19.0)	198	61.8 (19.7)	0.189	
PISQ SA-Partner Related	48	75.8 (24.6)	184	78.4 (24.5)	0.596	
PISQ SA-Condition Specific	49	96.7 (8.9)	195	89.8 (15.8)	<.001	0.05
PISQ SA-Global Quality	49	62.9 (26.9)	197	62.6 (30.4)	0.961	
PISQ SA-Condition Impact	49	92.5 (14.8)	198	79.8 (28.0)	<.001	0.01
PISQ SA-Desire	50	55.5 (21.9)	197	54.1 (21.9)	0.767	
GUPI Pain Subscale	65	4.1 (4.2)	300	4.7 (5.1)	0.461	
PFDI POPDI6	87	13.8 (14.2)	406	17.1 (20.0)	0.104	
PFDI CRADI8	87	14.8 (18.2)	404	21.0 (20.4)	0.022	0.15
PROMIS GI Constipation T-score	86	48.4 (8.8)	404	51.9 (8.9)	0.003	0.02
PROMIS GI Diarrhea T-score	88	46.2 (8.6)	410	49.5 (9.6)	0.008	0.02
PROMIS GI Bowel Incontinence Raw Score	87	4.8 (2.2)	389	5.4 (2.5)	0.058	

[*]
P-value from t-test.

[**]
P-value from multivariable linear regression models. Models were built using best subset selection with potential adjustment variables listed in Table 1. Full model results can be found in Supplementary Table 1.

PISQ=Pelvic Organ Prolapse/Urinary Incontinence Sexual Questionnaire; NSA = Not Sexually Active; SA = Sexually Active; GUPI = Genitourinary Pain Index; PFDI = Pelvic Floor Distress Inventory; POPDI = Pelvic Organ Prolapse Distress Inventory; CRADI = Colo-rectal-anal Distress Inventory; PROMIS = Patient-Reported Outcomes Measurement Information System; GI=Gastrointestinal

Table 3

Pelvic Floor Measures among Women by Urinary Incontinence Subtype

Indices	UUI		SUI		Mixed UI		Overall ANOVA p-value[*]	SUI vs. Mixed p-value[*]	UUI vs. Mixed p-value[*]	Adjusted Overall p-value[**]	Adjusted SUI vs. Mixed p-value[**]	Adjusted UUI vs. Mixed p-value[**]
	N	Mean (SD)	N	Mean (SD)	N	Mean (SD)						
PISQ NSA-Condition Specific	40	22.0 (30.3)	20	28.5 (23.8)	115	31.4 (30.2)	0.270	0.715	0.129			
PISQ NSA-Partner Related	42	69.4 (29.0)	20	43.3 (21.3)	118	58.8 (29.5)	0.010	0.049	0.074	0.004	0.04	0.03
PISQ NSA-Global Quality	40	40.4 (32.4)	20	59.1 (37.5)	116	49.5 (32.1)	0.143	0.271	0.172			
PISQ NSA-Condition Impact	41	18.7 (30.2)	20	25.6 (31.9)	115	26.5 (32.3)	0.434	0.901	0.221			
PISQ SA-Arousal Orgasm	41	65.0 (18.8)	46	66.9 (16.2)	98	57.7 (21.3)	0.034	0.021	0.082	0.05	0.05	0.06
PISQ SA-Partner Related	37	79.7 (27.8)	44	84.7 (18.3)	92	74.1 (26.2)	0.099	0.042	0.288			
PISQ SA-Condition Specific	41	93.2 (14.4)	45	91.3 (12.4)	96	87.1 (18.1)	0.136	0.198	0.078			
PISQ SA-Global Quality	41	65.8 (31.8)	45	68.2 (24.8)	98	57.9 (32.4)	0.169	0.099	0.213			
PISQ SA-Condition Impact	42	87.4 (24.4)	45	86.3 (22.6)	98	73.3 (29.9)	0.010	0.019	0.013	0.01	0.05	0.01
PISQ SA-Desire	41	56.1 (23.9)	45	52.9 (20.0)	98	53.8 (22.8)	0.811	0.836	0.616			
GUPI Pain Subscale	57	4.0 (4.6)	55	4.2 (4.7)	170	5.0 (5.4)	0.388	0.346	0.258			
PFDI POPDI6	84	10.7 (14.7)	68	12.4 (15.0)	229	20.6 (22.4)	<.001	0.008	<.001	0.004	0.01	0.004
PFDI CRADI8	84	18.3 (19.0)	68	12.8 (13.7)	227	24.3 (21.4)	<.001	<.001	0.036	0.02	0.01	0.20
PROMIS GI Constipation T-score	85	50.9 (8.3)	66	50.3 (8.6)	229	52.6 (9.0)	0.139	0.099	0.169			
PROMIS GI Diarrhea T-score	85	47.9 (9.1)	68	46.0 (8.7)	232	51.3 (9.8)	<.001	<.001	0.013	0.01	0.01	0.07
PROMIS GI Bowel Incontinence Raw Score	82	5.2 (2.3)	63	4.5 (1.2)	220	5.8 (2.9)	0.004	0.002	0.099	0.01	0.004	0.10

[*]
P-values from one-way ANOVA with pairwise p-values for SUI vs. Mixed and UUI vs. Mixed.

[**]
P-values from multivariable linear regression models (overall and pairwise p-values for SUI vs. Mixed and UUI vs. Mixed). Models were build using best subset selection with potential adjustment variables listed in Table 1. Full model results can be found in Supplementary Table 2.

PISQ=Pelvic Organ Prolapse/Urinary Incontinence Sexual Questionnaire; NSA = Not Sexually Active; SA = Sexually Active; GUPI = Genitourinary Pain Index; PFDI = Pelvic Floor Distress Inventory; POPDI = Pelvic Organ Prolapse Distress Inventory; CRADI = Colo-rectal-anal Distress Inventory; PROMIS = Patient-Reported Outcomes Measurement Information System; GI=Gastrointestinal