539: Validation of electronic (web-based & smartphone) administration of measures of pelvic floor dysfunction

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Objective: To validate responses on electronically administered (web-based and smartphone) validated instruments of pelvic floor dysfunction with responses on the same paper instruments **Methods:**

•Women presenting for evaluation of pelvic floor disorders at 5 FPMRS clinic sites invited to participate

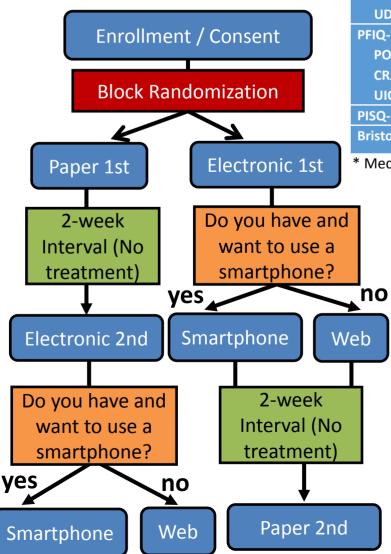
•Excluded if had intervention in the 2 weeks between initial and repeat questionnaire completion

•Randomized order of completion (electronic versus paper) of 4 validated questionnaires (PFDI-20, PFIQ-7, PISQ-12, Bristol Stool Scale) 2 weeks apart

•Power analysis: 54 subjects per arm (108 total) to detect effect size 0.5, 80% power, 2-tailed alpha 0.05.

•Analysis: Continuous variables compared with paired and student t-test; categorical values with chi-square; Pearson's & Spearman's coefficients used for correlation

Figure 1: Randomization scheme



Results:

•234 subjects enrolled; 132 (56%) completed study with no intervening treatment

•Mean age 58 (+-15) years-old, BMI 28 (+-6), parity 2, 77% white, 10% other, 7% Asian, 6% African-American

Complaints: 58% urinary, 37% prolapse, 5% bowel
No differences in demographics between those who completed data collection vs those who did not

•No age difference between those who completed electronic version on web vs smartphone

•No significant difference in responses for each total scale and individual scale between time points (first and second administration)

Table 1: Correlation of paper versus electronicresponses obtained 2 weeks apart

Questionnaire	Score Mean (std dev)	Correlation r, p-value
PFDI-20	77.5(47.6)	0.74, <0.001
POPDI	22.2 (18.8)	0.67, <0.001
CRADI	16.8 (15.5)	0.66, <0.001
UDI	32.5 (23.5)	0.77, <0.001
PFIQ-7	48.9 (51.7)	0.63, <0.001
POPIQ	11.7 (18.1)	0.50, <0.001
CRAIQ	11.3 (19.4)	0.68, <0.001
UIQ	25.9 (24.8)	0.72, <0.001
PISQ-12	18.1 (16.4)	0.86, <0.001
Bristol Stool Scale	4 (3,4)*	0.58, <0.001

* Median (25th %ile, 75th %ile)

Conclusion:

Scales of pelvic floor dysfunction have moderate to strong correlations when administered electronically versus on paper.
Our results support prior work demonstrating strong correlation between paper-based and electronic administration of PFDI-20 and PFIQ-7 and further demonstrates strong correlation for the PISQ-12 and moderate correlation for the Bristol Stool Scale.

•Our study is unique in demonstrating correlation between paper and web-based or smartphone administration.

•Electronic questionnaires, administered via the web or on a smartphone , provide an acceptable alternative to paper based forms for both research and clinical practice.

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