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SENSITIVITY OF THE PFDI-20 TO THE PRESENCE OR ABSENCE OF ANAL INCONTINENCE IN THE GENERAL POPULATION.

Hypothesis / aims of study

To establish whether PFDI-20 scores for women in the general population differ based on the presence and type of anal incontinence (AI) and to determine whether scores correspond with symptoms and degree of bother.

Study design, materials and methods

Subjects were recruited for this cross-sectional study during Twins Day Festivals in Twinsburg from 2004-2009. An anonymous survey on pelvic floor dysfunction, including questions assessing for anal incontinence (AI), was completed. In 2004, the PFDI was included; the PFDI-20 was used in subsequent years. For 2004 data, PFDI items constituting the PFDI-20 were analyzed. The PFDI-20 and its subscales (Colorectal-anal Distress Inventory (CRADI-8), Pelvic Organ Prolapse Distress Inventory (POPDI-6), and Urinary Distress Inventory (UDI-6)) were scored as previously described [1]. Scores were compared between continent and incontinent women and between incontinent subtypes by Wilcoxon rank-sum tests.

Results

PFDI-20 and all subscale scores differed significantly between subjects with AI of flatus only (n=265), AI of liquid stool (n=79), or AI of solid stool (n=44) and those denying any AI (n=1162) (Fig 1). CRADI-8, UDI-6, and PFDI-20 scores differed significantly between subjects socially bothered by AI of flatus and those unbothered. CRADI-8 and PFDI-20 scores differed significantly between subjects socially bothered by AI of liquid stool and those unbothered. PFDI-20 and all subscale scores differed significantly between subjects socially bothered by AI of solid stool and those unbothered (Fig 2).

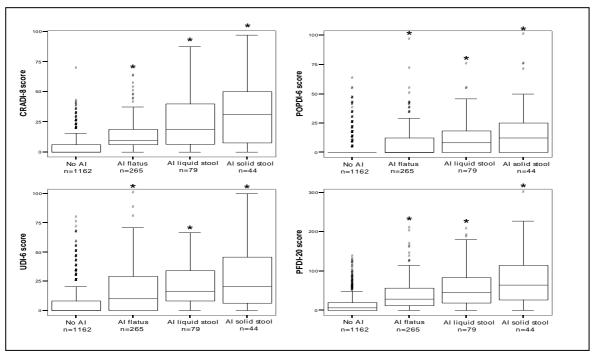
Interpretation of results

Women in the general population reporting AI of flatus, liquid stool, and solid stool segregate themselves from those without these conditions based on CRADI-8 and PFDI-20 scores; furthermore, scores for symptomatic subjects differ depending on whether their quality of life (social bother) is affected by their AI.

Concluding message

The PFDI-20 appears sensitive to patient symptomatology and degree of bother among women in the general population with

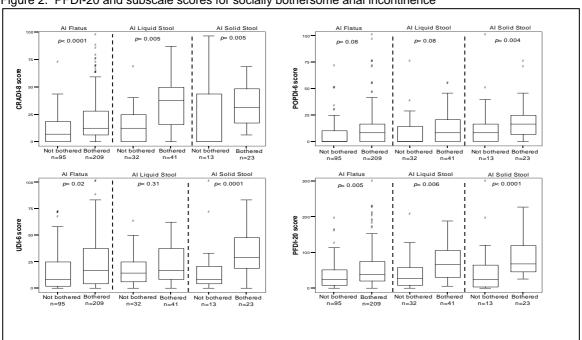
Figure 1. PFDI-20 and subscale scores for women with and without anal incontinence



CRADI-8, Colorectal-anal Distress Inventory; POPDI-6, Pelvic Organ Prolapse Distress Inventory; UDI-6, Urinary Distress Inventory; PFDI-20, short-form of the Pelvic Floor Distress Inventory

p<0.0001 when comparing incontinence subtype to no UI by Wilcoxon rank-sum test

Figure 2. PFDI-20 and subscale scores for socially bothersome anal incontinence



CRADI-8, Colorectal-anal Distress Inventory; POPDI-6, Pelvic Organ Prolapse Distress Inventory; UDI-6, Urinary Distress Inventory; PFDI-20, short-form of the Pelvic Floor Distress Inventory p-value determined by Wilcoxon rank-sum test

References

1. Am J Obstet Gynecol 2005;193:103-113.

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Is this a clinical trial?	No
What were the subjects in the study?	HUMAN
Was this study approved by an ethics committee?	Yes
Specify Name of Ethics Committee	NorthShore University HealthSystem Institutional Review Board Project#: EH03-260 (exempt research)
Was the Declaration of Helsinki followed?	Yes
Was informed consent obtained from the patients?	No