Treatment of Puborectalis Syndrome with Progressive Anal Dilation

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PURPOSE: The aim of this study is to assess the ability of progressive anal dilations to improve frequency of spontaneous bowel movements in patients with puborectalis syndrome (PRS). METHOD: Thirteen patients (9 females and 4 males; mean age, 37 years) with severe, chronic constipation caused by PRS were treated with daily, progressive anal dilation for a three-month period. Three dilators of 20, 23, and 27 mm in diameter were used. Dilators were inserted every day for 30 minutes (10 minutes each dilator). Patients were evaluated with anorectal manometry and defecography halfway through treatment, at the end of treatment, and six months after the end of treatment. At six months, patients also underwent physical examination. RESULTS: There was a significant improvement of weekly mean spontaneous bowel movements from zero to six (P < 0.0001), and the need for laxatives decreased from 12 patients with a weekly mean of 4.6 to 2 patients once per week (P <0.001). Enemas used before treatment by eight patients who had a weekly mean of 2.3 were, after treatment, needed only by three patients once per week (P < 0.01). During straining, tone measured with anorectal manometry decreased from 93 to 62 mmHg after six months of the end of therapy (F = 6.97; P < 0.01), and anorectal angle measured with defecography during the strain increased from 95° to 110° (P = not significant). CONCLUSIONS: Daily progressive anal dilation should be considered as the first and most simple therapeutic approach in patients with PRS. [Key words: Puborectalis syndrome; Anal dilators; Anorectal manometry; Defecography; Chronic constipation]

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 ${\bf F}$ unctional chronic constipation is divided into two groups: slow transit constipation as a result of colonic inertia and outlet obstruction,¹⁻¹⁰ which is mainly caused by puborectalis syndrome (PRS). Physical examination, anorectal manometry (ARM), electromyography (EMG), balloon evacuation proctography, and defecography are key examinations for diagnosis.⁸ Other useful investigations are digital scan defecography, defecography-computerized tomography scan, long-lasting rectosigmoid manometry with mechanical or drug-induced stimulation, and computerized anorectal manometry with three-dimensional study of the sphincter. At physical examination, patients with PRS show a paradoxical puborectalis contraction. During straining, failure of pelvic floor muscles to relax and an increased intraluminal pressure were detected. With balloon evacuation proctography, anorectal angle profile and its modifications during straining are studied. Defecography shows, besides anorectal angle modifications, the presence of rectocele and/or internal mucosal prolapse. Electromyography is useful during straining in showing a pathologic involvement of puborectalis function.

PRS, although simple to diagnose using the previously described methods, has always been difficult to treat. The aim of this study is to assess the ability of progressive anal dilations in improving the frequency of spontaneous bowel movements in patients with PRS.

PATIENTS AND METHODS

Thirteen patients (9 females and 4 males) with a mean age of 37 (range, 17–73) years who had severe chronic constipation ("outlet" type) were studied in our colorectal unit from 1993 to 1994. Diagnosis was made with physical examination, barium enema, colonoscopy, colonic transit time study, anorectal manometry, and defecography. Only three patients underwent electromyography.

All patients reported a history of incomplete, prolonged, and difficult evacuation with the need for constant use of enemas, laxatives, and digital defecation. In eight patients, physical examination during straining showed failure to relax the puborectalis muscle and an increased contraction of muscles in five patients. Defecography in all 13 patients showed increased activity of the puborectalis muscle and failure to expel rectal contents during defecation. All patients underwent ARM, which revealed high pressure levels during straining.

We treated the 13 patients with progressive anal dilation by daily insertion of three dilators (20, 23, and 27 mm in diameter) for a period of ten minutes each from the smallest to largest for a three-month period.

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Patients were followed up with ARM and defecography halfway through treatment, at the end of treatment, and six months after the end of treatment. Clinical assessment was performed before and six months after the end of therapy. Wilcoxon's signedrank test was used as a test of statistical significance to correlate clinical findings, and functional examinations were statistically correlated using analysis of variance.

RESULTS

Six months after the end of treatment, all patients reported good clinical outcome. All were satisfied with this therapy; and none complained of incontinence. Natural bowel movements (*i.e.*, no laxatives, bowel stimulants, or suppositories) increased for a mean of 0 to 6 per week (P < 0.0001; Table 1). Laxative need before treatment was reported by 12 patients as being a mean of 4.6 times per week; after treatment, laxatives were needed by only two patients and in both cases only once per week (P < 0.001; Table 1).

Similarly, before treatment eight patients required the use of enemas a mean of 2.3 times per week, whereas after treatment enema use was required by only three patients once per week (P < 0.001; Table 1). During straining, tone, measured with ARM, decreased from a mean of 93 mmHg before treatment to a mean of 77 mmHg halfway through treatment and 57 mmHg at end of treatment. At six months after treatment, a mean pressure of 62 mmHg was revealed

 Table 1.

 Individual Improvement with Progressive Anal Dilation

	Before	After
Spontaneous BMS*	0.15 ± 0.37	6.23 ± 4.10†
Laxative-assisted BMS*	4.61 ± 2.63	$0.15 \pm 0.37 \ddagger$
Enema-assisted BMS*	$\textbf{2.30} \pm \textbf{2.32}$	0.23 ± 0.43 §
* BMS = bowel movem	ents per week	

* BMS = bowel movements per week. † P < 0.0001. (F = 6.97; P < 0.01; Table 2). Anorectal angle measured during straining increased from a mean of 95° before treatment to a mean of 105° halfway through treatment and 114° by the end of treatment. Six months after treatment, it was approximately 110° (P = not significant; Table 2).

DISCUSSION

PRS was first described by Wasserman¹¹ in 1964 and is caused by the failure of the puborectalis muscle to relax during evacuation. More complex alterations of all pelvic floor musculature⁶ might coexist, but the deficit of puborectalis muscle is the most typical alteration directly connected with the disease.

The advent of many anorectal physiologic testings, such as ARM, electromyography, and defecography, has enhanced our ability to diagnose PRS. Yoshioka and Keighley¹² reported that defecography or EMG can be equally used to identify inappropriate puborectalis contraction. In accord with Corman,² we believe that defecography is preferable to that of EMG for making this diagnosis, because EMG is so uncomfortable and invasive. All 13 patients studied underwent defecography, and all showed increased activity of the puborectalis muscle and failure to expel rectal contents during defecation.

The role of ARM is still not clear.¹ We performed ARM in all patients and detected high pressure during straining.

Patients with PRS are usually first managed with a high-residue food diet to elicit rectal voiding. The next step is use of increasing doses of laxatives and enemas.¹³

Neither procedure is able to solve the problem. A wide variety of surgical and pharmacologic therapeutic approaches has been proposed to elicit puborectalis muscle relaxation.¹³

Anorectal myectomy, attempted with good results in Hirschsprung's disease, is not recommended for this syndrome. It does not show any lasting benefit and produces incontinence as a complication in 10 percent of patients.^{12, 14} More disappointing results

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ARM (During Strail	ning, mmHg) and Anored	tal Angle Grades at Defecogra	lphy			

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_	Before	Mid Point	End	6 Months After	
ARM Defecography	93.07 ± 28.90 95.38 ± 5.93	77.69 ± 20.06 105.76 ± 5.34	56.92 ± 14.36 114.23 ± 8.37	62.30 ± 15.35* 109.61 ± 10.60†	
ABM = anorecta	al manometry		<u></u>		

ARM = anorectal manometry.

* *F* = 6.97; *P* < 0.01.

† P = not significant.

[‡]*P* < 0.001.

[§] P < 0.01.

have been obtained with the controlled anal dilation with the Park's retractor.¹²

Local injection of *Clostridium botulinum* Type A toxin has been reported as another possible form of therapy.⁴ Botulinum toxin seems to be a promising treatment, but it is overshadowed by reports of a high morbidity rate. However, patients who have undergone treatment thus far are too few to conclude this with any degree of certainty. In our colorectal unit, work is now in progress to better test its efficacy.

Surgical division of the puborectalis muscle is now abandoned because of a high rate of postoperative incontinence.¹⁵ Several recent reports suggest that biofeedback training is the most effective treatment in PRS and has the highest success rate and lowest morbidity.¹³ Use of this procedure for treatment of PRS was first reported by van Baal and colleagues¹⁶ (one case treated) in 1984 and later by Weber *et al.*¹⁷ (six cases) in 1987. In 1990, Loening-Baucke⁷ reported results of biofeedback training in chronically constipated children with encopresis. His report supported the efficacy of this procedure. In the work of Wexner and colleagues,¹⁸ biofeedback training also appeared to be effective for treating adult patients affected with PRS with an 89 percent success rate.

In the present study, we report use of daily progressive anal dilations to treat PRS. We believe that using anal dilators affects not only the internal sphincter muscle but also the external sphincter and puborectalis muscles. Dilators oppose the resting physiologic contraction of the external sphincter and puborectalis muscles, decreasing the puborectalis paradoxical contraction that characterizes PRS. Anal dilators have already been successfully used in therapy of anal fissures,¹⁹ and we now think they can be of great help in treatment of PRS. All 13 patients treated with daily progressive anal dilation showed good clinical outcome. The most striking evidence is the appraisal of natural evacuations and the great decrease in use of laxatives and enemas. None of our patients was incontinent for formed stool, and none experienced mild mucus or urgency incontinence.

These objective results are correlated with an improvement of functional examinations. Moreover, this treatment is inexpensive, is easy to perform, can be done at home, can be repeated as many time as needed, and can be added to biofeedback training. We are now enrolling more patients to verify the promising results and following up the study patients to test how long the benefits last.

CONCLUSIONS

We believe that daily, progressive anal dilation should be considered first line therapy for patients with PRS and might help biofeedback training.

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