Topical phenylephrine increases anal canal resting pressure in patients with faecal incontinence

M J Cheetham, M A Kamm, R K S Phillips

Abstract

Introduction—The internal anal sphincter receives a stimulatory alpha, adrenergic innervation. Use of an adrenergic agonist may therefore have a role in treating patients with faecal incontinence.

Methods—Ten patients (seven females, median age 66 years) with passive faecal incontinence related to weak internal anal sphincter were studied. All patients had intact anal sphincters as assessed by endoanal ultrasound. Phenylephrine gel was applied in a double blind manner in concentrations of 0%, 10%, 20%, 30%, and 40% (Slaco Pharma (UK) Ltd, Watford, UK) on separate days. Maximum resting anal pressure (MRP), anodermal blood flow, blood pressure, and pulse rate were measured before, and one and two hours after application.

Results—All concentrations of phenylephrine gel increased median MRP (43, 48, 54, 65, and 70 cm H2O, for placebo, 10% (p=0.122), 20% (p=0.170), 30% (p=0.002), and 40% (p=0.004), respectively at one hour; comparisons with placebo). This was sustained at two hours. There was a clear dose-response relationship at one hour. Higher concentrations raised median MRP to within the normal range (> 60 cm H2O). At two hours, all concentrations greater than 20% increased the pressure to a similar degree, suggesting that the exact concentration may be important for the initial effect but given a certain threshold is less important after a period of time. Toxicity was rare. Two patients experienced transient perianal burning which settled within a few minutes. There was no significant effect on anodermal blood flow, blood pressure, or pulse rate.

Conclusion—This study has demonstrated the feasibility of using topical phenylephrine to raise resting anal tone in patients with faecal incontinence. Randomised controlled trials are required to assess the efficacy of this agent.

Keywords: phenylephrine; faecal incontinence; anal tone

Faecal incontinence is often related to structural sphincter damage, such as obstetric trauma,1 accidental trauma, or following anorectal surgery. Incontinence can also occur when the anal sphincter muscles are structurally intact, as assessed by endoanal ultrasound. The commonest cause for this appears to be a degenerative disorder of the internal anal sphincter.7 In this condition the internal sphincter becomes fibrotic and weak, leading to a reduced maximum anal canal resting pressure (MRP) and episodes of passive faecal incontinence (loss of stool without the patient’s awareness).7

In minor degrees of faecal incontinence, the use of a pad or anal plug1 may ease symptoms. The use of loperamide and low fibre diet may also be appropriate. In patients with external anal sphincter disruption, surgical repair of the sphincter is effective.7 In severe cases, major reconstructive surgery, such as an artificial sphincter implant or graciloplasty,7 may merit consideration. Such procedures are however still in the development phase and carry a high rate of complications and reoperation. Many patients have relatively minor symptoms which do not merit such radical surgery.

The internal anal sphincter exists in a state of tonic contraction, and is the main factor responsible for the generation of anal canal resting pressure.8 Given that the internal anal sphincter receives an extrinsic innervation which modulates its tone, we hypothesised that it may be possible to mimic the effect of this innervation and pharmacologically increase internal anal sphincter tone.

Phenylephrine is a selective alpha, agonist which causes internal sphincter contraction in vitro10 and elevates MRP in animal studies.11 A recent pilot study from this unit has shown that topical phenylephrine can increase internal anal sphincter tone in healthy volunteers.12 This dose ranging study demonstrated that 10% phenylephrine gel was the optimum concentration, raising MRP by a mean of 33%. This elevation in MRP persisted for a median of seven hours following a single application. However, when the efficacy of 10% phenylephrine in treating faecal incontinence was assessed in a controlled trial, the clinical response was disappointing.13 Of 36 patients enrolled in the study, six reported a significant improvement in their symptoms. Patients with faecal incontinence are known to have an internal anal sphincter which is less responsive to alpha adrenoceptor agonists in vitro.14 The poor in vivo response may therefore have been related to an inadequate concentration of the applied gel.

This study therefore aimed to assess the effect of topical phenylephrine on MRP in patients with passive faecal incontinence and to determine its optimal concentration.

Abbreviations used in this paper: MRP, maximum anal resting pressure.
**Table 1** Effect of a single application of a gel containing different phenylephrine concentrations on maximum anal resting pressure (MRP)

<table>
<thead>
<tr>
<th>Phenylephrine</th>
<th>Baseline MRP</th>
<th>MRP at 1 h</th>
<th>p*</th>
<th>MRP at 2 h</th>
<th>p*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Placebo</td>
<td>35.5 (29–68)</td>
<td>42.5 (31–64)</td>
<td>N/A</td>
<td>48.0 (30–58)</td>
<td>N/A</td>
</tr>
<tr>
<td>10%</td>
<td>35.5 (19–77)</td>
<td>41.0 (28–82)</td>
<td>0.122</td>
<td>63.5 (32–76)</td>
<td>0.068</td>
</tr>
<tr>
<td>20%</td>
<td>38.0 (29–60)</td>
<td>54.0 (33–81)</td>
<td>0.170</td>
<td>65.0 (31–82)</td>
<td>0.006</td>
</tr>
<tr>
<td>30%</td>
<td>37.0 (28–68)</td>
<td>65.0 (32–77)</td>
<td>0.002</td>
<td>67.5 (29–80)</td>
<td>0.002</td>
</tr>
<tr>
<td>40%</td>
<td>43.0 (29–60)</td>
<td>69.5 (28–89)</td>
<td>0.004</td>
<td>63.5 (31–82)</td>
<td>0.006</td>
</tr>
</tbody>
</table>

Values are median (range).

Results from 10 patients with passive faecal incontinence. Values at one and two hours were compared with values pretreatment (baseline). Subjects were studied on five separate days.

*Comparison with placebo using Friedman multiple comparisons (Bonferroni corrected value).

**Patients and methods**

We studied patients with exclusively passive faecal incontinence—that is, loss of stool without the patient’s awareness. All complained of episodes of faecal incontinence to liquid or solid stool.

All patients were assessed by anorectal physiological testing and endoanal ultrasound scanning. Patients were included only if they had circumferentially intact sphincters demonstrable on endoanal ultrasound scanning, and a low MRP indicative of impaired internal anal sphincter function.

Patients were excluded if they were pregnant, had ischaemic heart disease, aortic aneurysm, uncontrolled hypertension, inflammatory bowel disease, or other disorders known to cause secondary faecal incontinence.

Patients attended for five visits a minimum of 48 hours apart. At the first visit, incontinence scores were assessed for each patient. Continence was rated from 0 (perfect continence) to 24 (daily episodes of incontinence to solid stool with major lifestyle restriction) using a recently published faecal incontinence grading scale.

On each of the five study days, measurements were made of MRP, anodermal blood flow, blood pressure, and pulse rate at baseline (pre-gel application), and at one and two hours after application. When baseline measurements had been obtained, a 2.5 cm strip of the test gel was applied to the distal anal canal using the gloved index finger of the investigator.

MRP was measured in each subject using an eight channel water perfused anorectal manometry system (Muir Scientific, Ontario, Canada). A station pull through technique was used at 1 cm intervals to identify the point of MRP. The catheter was then left in situ to establish a stable resting pressure. MRP was determined by taking the mean of the eight circumferential pressure measurements. Anodermal blood flow was measured using the DRT4 laser Doppler flowmeter (Moor Instruments, Devon, UK). The skin probe was applied to the skin of the anus and held until a steady reading was obtained. Anodermal blood flow was calculated by taking the mean of four quadrant readings around the anus.

Gels containing no active ingredient (placebo), and 10%, 20%, 30%, and 40% phenylephrine were used. All gels were supplied in identical coded foil tubes by Slaco Pharma (UK) Ltd (Watford, UK). Both the investigator and patients were unaware of the nature of each gel. Gels were applied in a random order defined before the start of the study.

**Results**

**Patients**

Ten patients (three males) with a median duration of incontinence of two years (range 1–7) were studied. Median incontinence score was 13.5 (range 7–20 out of 24).

Endoanal ultrasound demonstrated the integrity of both anal sphincters in all patients. The mean thickness of the internal anal sphincter was 1.9 mm (range 1.1–2.7). At the time of patient selection, anorectal physiological testing demonstrated a median MRP of 38.5 cm H2O (range 13–60, normal in our laboratory >60 cm H2O), with a median squeeze pressure of 100 cm H2O (range 51–288, normal >60 cm H2O). Median left pudendal nerve terminal motor latency was 2.1 (range 1.3–2.2). Median right pudendal nerve terminal motor latency was 2.2 (range 1.6–2.2).

**Effect on maximum resting pressure**

There was no significant difference between baseline MRPs for the five study days when the placebo and four different concentrations were tested. Application of all concentrations of phenylephrine gels resulted in an increase in median MRP at both time points (Table 1). This effect was statistically significant (p<0.05 compared with placebo) at both one and two hours for phenylephrine gels in concentrations of 30% and 40% only. Application of 30% and 40% phenylephrine resulted in an increase in median MRPs to within the normal range for this unit (60–120 cm H2O). Individual patient data are shown in figs 1–3. Three patients had minimal response to all concentrations of phenylephrine studied (defined as an elevation less than 10 cm H2O in maximum resting pressure at either time point) (figs 1–3).

**Effect on skin flux**

There was wide variation in values obtained for anodermal skin flux, both within and between patients. No significant effect on skin flux was...
noted with any of the concentrations of phenylephrine studied.

LOCAL SIDE EFFECTS
One patient reported a stinging sensation immediately on application of 30% phenylephrine gel. Another patient reported transient burning on application of 20% and 40% phenylephrine, but not with other concentrations. In all cases, local discomfort settled within 20 minutes. The remaining eight patients did not report any local side effects on direct questioning. No patient reported a rash.

Three patients who had no response or only a minimal response to phenylephrine were identified (non-responders). There were no clear differences between responders and non-responders in relation to clinical or investigational parameters. The numbers were too small to compare statistically, given that there were three non-responders and seven responders.

Discussion
This study has demonstrated that it is possible to increase the resting tone of the internal anal sphincter in patients with faecal incontinence. This has not been demonstrated previously. We have also shown that this is possible using a topical preparation, minimising the potential for systemic side effects. A dose-response relationship was demonstrated, with higher doses causing an increase in resting pressure. With higher doses the increase in resting pressure was of a magnitude which increased the pressure to within the normal range, suggesting that this effect has therapeutic potential.

Minimal toxicity was noted. Phenylephrine has been used topically in ophthalmology with reports of local irritation and cardiovascular side effects. However, the cornea may offer greater access to the systemic circulation than the anoderm. In the current study, even with higher concentrations of gel, only minor skin irritation was observed by two patients and this resolved after a few minutes. Longer term studies will need to explore local side effects in greater detail. Anodermal blood flow was not affected, suggesting that the vasoconstrictive effect of phenylephrine was balanced by other local factors. No effect on blood pressure was observed.

Previous work from our unit demonstrated that 10% topical phenylephrine significantly increased the MRP in healthy volunteers, and that higher concentrations did not achieve a greater resting pressure. This concentration was then used in a double blind study of patients with an ileoanal pouch and faecal incontinence. These patients had a structurally normal anal sphincter and a normal resting pressure, with faecal incontinence thought to relate to lack of normal neural sphincter control. Half of the patients achieved clinical benefit, and one third had complete resolution of their incontinence.

However, when this concentration of phenylephrine was used in a double blind trial of patients with faecal incontinence due to a weak but intact sphincter, benefit was experienced by only a minority of patients, with no significant effect for the group as a whole. This led us to reconsider the nature of the pathology in these patients, and consequently pharmaco-
Topical phenylephrine increases anal canal resting pressure

were slightly greater than 60 cm H2O. This on several separate occasions. On some of these range for this unit). Each patient was studied than 60 cm H2O (the lower limit of the normal

Although concentrations of 10% and 20% phenylephrine resulted in a modest increase in median resting pressure, this was not statistically significant. In contrast, application of 30% and 40% phenylephrine resulted in a significant increase in resting pressure at both time points compared with placebo.

One hour after application there was a dose-response relationship for different concentrations. The pressure response was maintained at two hours. Given this two hour response, and data from volunteer studies suggesting that the effect of a single application of topical phenylephrine is sustained for a median of seven hours, we believe there is the potential for a sustained clinical response.

At inclusion, all patients had an MRP of less than 60 cm H2O (the lower limit of the normal range for this unit). Each patient was studied on several separate occasions. On some of these subsequent occasions, baseline values for MRP were slightly greater than 60 cm H2O. This reflects natural variation in measurement. These patients did however have passive faecal incontinence and had a low resting pressure measured on more than one occasion.

This dose ranging study was restricted to patients with a weak sphincter and passive faecal incontinence. This group is likely to need the greatest concentration of phenylephrine to improve sphincter function. It is possible that this type of topical therapy will affect sphincter function even in the presence of structural damage, or when the main complaint is urgency and the sphincters are intact and functioning normally. Phenylephrine causes relaxation of rectal circular muscle in vitro, and this effect may also contribute to improved continence. An open study in which phenylephrine was administered systemically to volunteers demonstrated no alteration in colonic motility; these studies suggest that accidental rectal administration of topical phenylephrine will not adversely affect the rectoanal pressure gradient. The aim of the present study was restricted to exploring the feasibility of using a topical alpha, agonist to increase anal resting pressure. Further studies may assess the effect of topical phenylephrine on rectal motility.

Clinical studies are now required in patients with faecal incontinence to assess the efficacy of this treatment.

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