A physiological study of the ano rectal reflex in patients with functional anorectal and defecation disorders

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Abstract

OBJECTIVE: The symptoms of fecal incontinence and constipation can arise from a variety of alterations of anorectal function. The aim of this study was to investigate components of the anorectal reflex in patients with these...
METHODS: 21 healthy volunteers (controls), 78 incontinent and 74 constipated patients were recruited and symptom burden determined using the Wexner incontinence and constipation questionnaires. All participants underwent standardised anorectal physiology including anorectal manometry, anorectal distension and electrosensitivity thresholds, rectal mucosal blood flow and rectoanal inhibitory reflex measurement.

RESULTS: Patients with passive incontinence had lower resting sphincter pressures than controls (38 vs 87cmH₂O, p<0.05), while those with urge incontinence had lower squeeze pressures than controls (37 vs 119cmH₂O, p<0.05). Patients with urge incontinence had lower maximal tolerable volumes (100 vs 166mL, p<0.05). Patients with slow transit constipation had elevated rectal electrosensitivity thresholds compared to controls (31.4 vs 20.2, p<0.05), and also showed lower mucosal blood flow than patients with evacuation difficult and controls (107 vs 162 (evacuation difficult) vs 169 (controls), p<0.05). Only patients with passive incontinence were associated with reflex abnormalities (prolonged recovery phase (1.2 vs 0.5msec, p<0.05) and total duration of reflex (6.3 vs 4.3msec, p<0.05)).

CONCLUSION: Anorectal motor, sensory and reflex abnormalities are seen in distinct patterns in patients with fecal incontinence and constipation. This would suggest distinct physiological differences which may predict the
potential for different treatment neuromodulation and behavioural modalities in these conditions.

**Keywords:** anorectal physiology, rectoanal inhibitory reflex, fecal incontinence, constipation
Introduction

Idiopathic fecal incontinence (FI) is a functional anorectal disorder that affects up to 8% of individuals,1 whilst functional defecation disorders affect up to 19% of the North American population.2 Both have a higher prevalence amongst females and the elderly.3 4 The significant negative impact between symptom burden and impaired quality of life is well established in patients suffering from FI or constipation.5

Anorectal physiology forms an important component of FI assessment, although its value remains controversial. Currently the most common recorded variables are: static and squeeze anal sphincter pressures, sensitivity to rectal distention and mucosal electro-stimulation.6 Studies of anal sphincter function do not reliably correlate with symptoms, as for example, not all patients with low sphincter pressures suffer from FI.7 8 Moreover, a sub-set of patients with FI and proven low anal sphincter pressures report an improvement in symptoms with sacral nerve stimulation, which does not alter anal sphincter function.9 This has led to the theory that the etiology of functional FI involves not just anal sphincter function, but also rectal function.

Extrinsic neural control of rectal and anal sphincter through the pudendal and autonomic nerves (ANS) is central to controlling continence. Pudendal nerve function can be measured by Pudendal Nerve Terminal Motor Latency (PNTML), however the results do not consistently correlate with anorectal function, symptoms or treatment outcomes.10 11 ANS function may be
assessed using rectal mucosal blood flow (RMBF) as a surrogate marker of gut specific function, and has been shown to correlate with severity of constipation in patients with multiple sclerosis (MS) and spinal cord injury.\textsuperscript{12}

The rectoanal inhibitory reflex (RAIR) is an intrinsic reflex influenced by descending spinal pathways. This dynamic reflex is characterised by relaxation and then a contraction of the anal sphincters in response to rectal distension, and is a major contributor to continence.\textsuperscript{13} Abnormalities of the RAIR have been implicated in both constipation and FI in patients with diabetes, scleroderma, multiple sclerosis and spinal cord injury.\textsuperscript{12,14,15}

The relationship between altered anorectal physiology and functional FI and constipation is poorly understood. The study of static and dynamic anorectal physiology may provide an important tool for understanding the pathology of functional anorectal disease and may reveal potential treatment options. Therefore, the aim of this study was to investigate the relationship between anorectal physiological parameters (including: PNTML, manometry, mucosal electro-stimulation, RAIR and RMBF) in patients with symptoms of functional anorectal and defecation disorders.

**Methods**

This single centre prospective case-control study was undertaken at the Physiology Unit at University College Hospital, London, UK. Full ethical approval was granted by St Mark’s & Northwick Park Ethics Committee.
Patients & controls

Adult patients referred to the Physiology Unit at University College Hospital were recruited. Patients under 18 years old, a previous history of colorectal or anal surgery (including surgical or orifice treatment for haemorrhoids), symptoms suggestive of irritable bowel syndrome (as defined by the Rome III criteria) or any other systemic disease affecting gastrointestinal function (including: diabetes, multiple sclerosis, spinal cord injury and connective tissue disorders were excluded.

All patients underwent a full clinical assessment including a detailed history of symptom burden. Patients with a medication history that may influence bowel function were excluded. Organic disease was further excluded through comprehensive laboratory tests and lower gastrointestinal endoscopy. The Rome III criteria was applied to differentiate functional anorectal disorders (namely FI predominant symptoms) from functional defecation disorders. This enabled classification into one of four groups: 1) urge FI, 2) passive FI, 3) slow transit and 4) evacuation difficulty. Following group allocation, symptom load was then assessed using the validated Wexner questionnaires for constipation and FI. [16, 17]

Healthy volunteers (controls) were recruited by hospital advertisements and no financial incentives were offered. The same exclusion criteria used for recruitment of patients from the physiology unit were also applied to healthy controls.
All patients and controls underwent a series of standardised physiological investigations including: anal manometry, evaluation of rectal sensory thresholds, evaluation of pudendal nerve function and rectal mucosal blood flow.

**Anorectal physiological investigations**

**Whole gut transit study**

Patients allocated to the constipation predominant groups underwent whole gut transit time assessment through ingestion of radio opaque markers (Dunn Clinical Nutrition, Cambridge, UK) over a three day period and timed plain abdominal films using the validated method described by Evans et al (1992). Healthy volunteers should retain approximately 20% of markers within 12 hours and less than 80% after 120 hours. This enabled classification of patients into two groups: slow transit constipation or rectal evacuation disorder with normal transit time.

**Anorectal manometry**

Anorectal manometry was performed using an eight-channel water perfused catheter linked to a pneumohydraulic water perfusion system (Medial Measurements Systems, Enschede, Netherlands). Functional anal canal length and maximum voluntary and involuntary squeeze pressures were determined using a stationary pull through method.

**Rectal distension sensitivity thresholds**
Rectal sensation was assessed by slow inflation of a latex rectal balloon (Ardmore Healthcare, Bognor Regis, UK) with air at 1 mL/s to elicit the following variables: threshold volume for first constant sensation, defecatory urge volume (DDV), and maximum tolerable volume (MTV). Based on previous studies in this unit, normal ranges were set as: threshold volume of 20-110mL, DDV of 60-170mL and MTV of 110-320mL.12

Pudendal nerve terminal motor latency

Pudendal nerve terminal motor latency (PNTML) was assessed with the St Mark’s Pudendal Stimulating Electrode (Dantec Medical, Skovlunde, Denmark). Age adjusted PNTML values were used in order to assess for the presence of pudendal neuropathy (PNTML >2.3ms below 40-years-old, and >2.5ms in those 40-years or older).19

Anorectal electrosensitivity thresholds

A bipolar electrode ring catheter (Gaeltec Devices Ltd, Strathclyde, UK) and the Dantec Keypoint EMG/NCS/EP Workstation (Natus Medical Incorporated, Pleasanton, CA, USA) was used to assess anal and rectal mucosal electrosensitivity thresholds. Briefly, the protocol involved initial electrical stimulation of the anal canal at 5Hz with a pulse width of 0.1ms, followed by an incremental increase in the current to a maximum of 20mA or until the patient reported a change in sensation. Values for the rectum were an initial stimulation of 10Hz, pulse width 0.5ms and increased to a maximum of 50mA.20
Rectoanal inhibitory reflex

The rectoanal inhibitory reflex (RAIR) was assessed by insertion of a latex rectal balloon (Ardmore Healthcare, Bognor Regis, UK) into the lower rectum and rapid inflation and deflation with 50mL of sterile water. Changes in pressure were recorded using the eight-channel manometry catheter. Persistent failure to elicit the RAIR despite repositioning the catheter prompted an increase in the inflation volume to 100mL. The following parameters were measured: excitation latency, the percentage reduction in amplitude of the anal pressure compared to resting anal pressure, recovery time and total duration of the reflex (further details provided in figure 1). If the RAIR was observed in multiple catheter channels the one demonstrating the largest change in amplitude was selected for analysis.

Rectal mucosal blood flow

Rectal mucosal blood flow (RMBF) was assessed using a Doppler probe (DP6A Moor Instruments, Axminster, UK) at 10cm from the anal verge at four circumferential points after allowing stabilisation of the trace for at least 30 seconds.

Statistical analysis

Five study groups were defined based on clinical and physiological assessment: 1) controls, 2) urge FI, 3) passive FI, 4) slow transit and 5) evacuation difficult constipation. All controls were matched for age and sex. Normally distributed variables were expressed as mean (+/- SD) otherwise they were recorded as median and ranges. Comparisons between control and
patient groups were performed using one-way analysis unpaired t-test for parametric data, or Mann-Witney U test for non-parametric data. SPSS 20 (IBM, Armonk, New York, USA) was used for all statistical tests.

**Results**

Twenty one healthy volunteers (controls, 14 female; mean age 35) and 152 patients (age range 18–72) were included. Seventy eight patients (63 female; mean age 45) had FI-predominant symptoms (44 urge FI and 34 passive FI), and 74 patients (59 female; mean age 32) had constipation-predominant symptoms (47 slow transit, defined by whole gut transit studies and 27 with evacuation difficulty.

**Questionnaires**

Mean (SD) scores for Wexner FI and constipation scores are shown in table 1. The symptom questionnaires demonstrated that controls had low symptom scores, consistent with previous studies. Patients with urge or passive FI had significantly higher Wexner FI scores than controls (p<0.001). Similarly, patients with slow transit (p<0.03) or evacuation difficulty (p<0.01) had higher Wexner constipation scores than controls.

**Anorectal physiology**

**Anal manometry**

Mean (SD) resting and squeeze sphincter pressures are shown in table 2. Patients with FI-predominant symptoms had lower resting and squeeze pressures than controls (p<0.05). Patients with passive FI had lower resting
sphincter pressures than controls (38 vs 87 cmH2O, p<0.05) and patients with urge FI (38 vs 53 cmH2O, p<0.05). Patients with urge FI had lower squeeze pressures than controls (37 vs 119 cmH2O, p<0.05) and patients with passive FI (37 vs 84 cmH2O, p<0.05). Patients with constipation-predominant symptoms had no disturbance of anal sphincter pressures.

**Rectal sensory function to distension (table 2)**

Patients with urge FI required lower volumes to elicit first sensation, DDV and MTV as compared to all other groups. However, only the reduced MTV was statistically significant when compared to controls (100 Vs 166 mL, p<0.05). Patients with slow transit and evacuation difficulty-constipation reported elevated volumes to elicit these variables, although only DDV was statistically significant volume when compared to controls and both urge and passive FI (94 (slow transit), 90 (evacuation difficulty) vs 69 mL (controls), p<0.05).

**Pudendal nerve terminal motor latency & anorectal electrosensitivity thresholds (figure 2)**

All controls and patients had age appropriate PNTMLs. All controls and patients with urge or passive FI and evacuation difficulty had normal anorectal electrosensitivity thresholds. Patients with slow transit constipation had elevated rectal electrosensitivity thresholds compared to controls (31.4 vs 20.2, p<0.05).

**Rectal mucosal blood flow**
RMBF in patients with incontinence-predominant symptoms were similar to those observed in controls. Patients with slow transit constipation had decreased RMBF when compared to both patients with evacuation difficult and controls (107 vs 162 (evacuation difficult), 169 (controls), p<0.05).

Rectoanal inhibitory reflex (table 3)
Urge FI was not associated with any significant abnormalities of the RAIR when compared to controls. In contrast, passive FI was associated with a prolonged recovery phase (1.2 vs 0.5msec, p<0.05) and total duration of reflex (6.3 vs 4.3msec, p<0.05) when compared to controls. Patients with constipation predominant symptoms were not associated with any abnormalities in the RAIR reflex parameters compared with controls or incontinence-predominant patients.

Discussion
This was a single centre study investigating the physiological parameters of the anorectal reflex in patients with functional anorectal and defecation disorders. This heterogeneous cohort of patients often have limited treatment options, which are compounded by unpredictable treatment outcomes. These challenges even more complex in the absence of known organic causes, meaning that patients with functional (idiopathic) anorectal and defecation disorders represent a particularly challenging cohort of patients to manage.

The relationship between anorectal physiology and functional FI and constipation has previously been investigated by Zbar et al (1998).21
However, their study failed to address the heterogeneity of symptoms within each patient group (i.e. urge FI vs passive FI and slow transit vs evacuation difficult constipation). This represents an important source of bias, making interpretation of results difficult and hinders application to real life clinical practice. In contrast, our study overcame this limitation by analysing patients according to the exact nature of their FI and constipation in order to demonstrate physiological differences underpinning symptom burden. This may enable more effective targeting to treatment options.

Patients with functional (idiopathic) FI or constipation had relatively disparate symptom profiles. Unsurprisingly, those with FI had low constipation scores whilst those with predominant symptoms of constipation had low FI scores. Of note, this is in contrast to patients with central neuropathy syndromes who describe a mixed symptom burden. Abnormalities in rectal sensation are considered to be important in the development of anorectal dysfunction. In this study, rectal balloon distension demonstrated reduced rectal sensory thresholds in patients with urge FI, but not passive FI. Chan et al (2005) demonstrated similar findings in a subset of patient with urge FI, suggesting that symptom burden within this cohort of patients is secondary to rectal hypersensitivity. Moreover, patients with urge FI in this study also reported decreased voluntary anal squeeze pressures (external anal sphincter function) suggesting a pathophysiological association between rectal sensation and voluntary sphincter action within patient group. It is unclear if lower squeeze pressures leads to rectal conditioning to lower rectal volumes in an effort to avoid urge FI. Alternatively, rectal hypersensitivity may lead to
impaired tolerance of rectal distention to relatively low rectal volumes and in turn to urge FI.

This study did not show any variation in PNTML and symptom burden across all patient groups. The relevance of this is unclear, and may reflect the findings of previous studies that suggest measurement of PNTML is an insensitive test for underlying neuropathy.\(^{25}\) For examples, Hill et al (2002) found that 31% of patients with bilaterally prolonged PNTML had normal squeeze pressures, and 49% with normal PNTML had abnormal squeeze pressures.\(^8\) It is possible that these results reflect the underlying limitations of this method of measuring extrinsic neuronal control, whereby the presence of a few nerve fibres (insufficient to provide reliable control of continence) result in false positives. Furthermore, the test is highly operator dependant and poorly tolerated by patients.

Patients with known functional (idiopathic) constipation have been shown to have diminished rectal sensation to distention, resulting in rectal mucosal sensation to electrical stimulation.\(^{26}\) This study revealed that this finding is unique to patients with slow transit, but not evacuation difficult constipation. In contrast to FI, this indicates an element of rectal hyposensitivity, and similar findings have been reported in patients with diabetes and MS.\(^{12, 14}\) Sensation to electrical stimulation is detected by non-myelinated c-fibres within the rectal mucosa, and is transmitted to higher centres via parasympathetic fibres within the nervi erigentes. Abnormal myenteric plexus transmission and altered
neurotransmitter concentrations have been observed in patients with slow transit constipation indicating visceral neuropathy etiology.\textsuperscript{27}

Consistent with previous studies,\textsuperscript{28} we found that RMBF is reduced in patients constipation-predominant symptoms, although this study revealed that this finding is unique to patients with slow transit constipation only. RMBF is used as a surrogate measurement of colonic and rectal autonomic tone. Reduced RMBF is associated with psychological stress,\textsuperscript{29} muscarinic antagonists\textsuperscript{30} and constipation. Indeed, RMBF has previously been shown to correlate with severity of constipation.\textsuperscript{31} As functional (idiopathic) anorectal and defecation disorders are often multifactorial, these findings provide evidence that autonomic tone is affected in patients suffering from constipation-predominant symptoms.

Normal RAIR is mediated by both sacral and myenteric neurones, and abnormal RAIR morphology has previously been shown to correlate with FI.\textsuperscript{32} Within this study, abnormal RAIR morphology was found only in patients with passive FI. Previous investigations of healthy volunteers have shown that resting pressures are largely governed by the internal anal sphincter, which relaxes in response to rectal distention, whilst the external anal sphincter contracts involuntarily (during the so-called rectal sampling reflex).\textsuperscript{33} Patients with passive incontinence had prolonged relaxation and recovery phases of the RAIR. The combination of lower resting pressures, any prolongation of the RAIR is likely to reduce sphincter pressures further, and therefore predispose
to passive FI. Similar reflex patterns amongst patients with idiopathic passive incontinence have been demonstrated.\textsuperscript{21}

This group has previously reported on disturbances in anorectal physiology in patients with neurogenic (MS and SCI) and diabetic-associated bowel disorders\textsuperscript{12,14}. Severity of constipation in MS and SCI correlates with reduced RMBF, a finding that was mirrored for patients in this study with slow transit functional constipation. Severity of FI symptoms in patients with MS, SCI and diabetes correlates with abnormalities of the RAIR, namely prolonged duration and recovery. In this study, similar findings were shown for patients with passive FI but not urge FI. While patients with neurogenic bowel disorders and constipation predominant symptoms are known to have impaired relaxation of the anal sphincters during the RAIR, patients with functional constipation consistently demonstrate an intact reflex. Further research should now be undertaken to evaluate the role of these investigations in the management of functional anorectal and defecation disorders. The observations reported in this study may be clinically useful in selecting appropriate treatment options for patients with functional anorectal and defecation disorders, whereby patients with preserved anorectal sensation may be suitable candidates for biofeedback, whereas trans-anal irrigation may be more appropriate in patients with intact anorectal reflexes.

It could be argued that the predominance of females included in the study may limit generalisability of our results. Alternatively, it may simply reflect the increased prevalence of FI and constipation among female patients.\textsuperscript{3} Indeed,
FI has been shown to be up to eight times more prevalent in women, which is largely attributed to obstetric factors.\textsuperscript{34} Similarly, the prevalence of functional (idiopathic) constipation has been reported as affecting up to 15% of women compared to 5.2% of men.\textsuperscript{35} All of the physiological measurements were performed in all subjects (controls and patients). While the order of investigations was standardised, it is possible that this order influenced subsequent measurements.

In summary, the findings of this novel study provide new insights into the pathophysiology of the different symptom profiles observed in patients with functional (idiopathic) anorectal and defecation disorders. Furthermore, alterations in anorectal physiology appear to reflect the changes seen in other non-related bowel disorders. Therefore, consideration of these physiological differences may aid management decisions within the complex patient group.
References


The rectoanal inhibitory reflex (RAIR), taken from Thiruppathy et al. (2015): the point of maximal stimulation is the starting point of the RAIR and the end-point is when the pressure has recovered to two thirds of the initial resting pressure. The time between these points is the duration of the reflex. Time taken for the pressure to return to resting from maximal amplitude is the excitation latency. Recovery time begins as soon as the RAIR reaches maximal relaxation.

![Diagram of RAIR](image-url)
Figure 2

Mean current (mA) required to elicit anorectal electrosensitivity thresholds across each of the five study groups.

STC = slow transit constipation; ED = evacuation difficulty constipation

* p<0.05 compared to controls
Table 1

<table>
<thead>
<tr>
<th>Group</th>
<th>Wexner FI Score (0-24)</th>
<th>Wexner Constipation Score (0-30)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>1.1 (0.2)</td>
<td>3.1 (1.3)</td>
</tr>
<tr>
<td>Urge FI</td>
<td>15.4 (7.4)</td>
<td>4.1 (2.8)</td>
</tr>
<tr>
<td>Passive FI</td>
<td>18.1 (7.5)</td>
<td>4.2 (4.3)</td>
</tr>
<tr>
<td>Slow transit</td>
<td>5.7 (3.0)</td>
<td>14.9 (8.9)</td>
</tr>
<tr>
<td>Evacuation difficulty</td>
<td>7.5 (2.9)</td>
<td>20.6 (12.7)</td>
</tr>
</tbody>
</table>

Mean (SD) Wexner FI and constipation scores across each of the five study groups. The Wexner FI score ranges from 0-20 with a score of ≥9 shown to correlate with moderate-severe symptoms and the Wexner constipation score ranges from 0-30 with a score of ≥10 shown to correlate with moderate-severe symptoms.
Table 2

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Controls</th>
<th>Urge FI</th>
<th>Passive FI</th>
<th>Slow transit</th>
<th>Evacuation difficulty</th>
</tr>
</thead>
<tbody>
<tr>
<td>Resting pressure (cmH$_2$O)</td>
<td>87 (35)</td>
<td>53 (28)$^\Delta$</td>
<td>38 (25)$^*$</td>
<td>78 (21)$^\Delta$</td>
<td>75 (28)$^\Delta$</td>
</tr>
<tr>
<td>Squeeze pressure (cmH$_2$O)</td>
<td>119 (52)</td>
<td>37 (21)$^*$</td>
<td>84 (38)$^{\dagger}$</td>
<td>109 (36)$^{\dagger\Delta}$</td>
<td>102 (41)$^{\dagger\Delta}$</td>
</tr>
<tr>
<td>Threshold volume (mL)</td>
<td>34 (14)</td>
<td>26 (17)</td>
<td>39 (18)</td>
<td>49 (20)</td>
<td>43 (23)</td>
</tr>
<tr>
<td>DDV (mL)</td>
<td>69 (44)</td>
<td>58 (24)</td>
<td>68 (34)</td>
<td>94 (28)$^{\dagger}$</td>
<td>90 (30)$^{\dagger}$</td>
</tr>
<tr>
<td>MTV (mL)</td>
<td>166 (78)</td>
<td>100 (29)$^*$</td>
<td>126 (42)</td>
<td>158 (39)</td>
<td>173 (42)</td>
</tr>
</tbody>
</table>

Mean (SD) parameters for anal manometry and rectal sensory function to distention across each study group.

DDV = defecatory urge volume; MTV = maximum tolerated volume

* p<0.05 compared to controls; † p<0.05 compared to urge FI; $\Delta$ p<0.05 compared to passive FI
Table 3

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Controls</th>
<th>Urge FI</th>
<th>Passive FI</th>
<th>Slow transit</th>
<th>Evacuation difficulty</th>
</tr>
</thead>
<tbody>
<tr>
<td>Excitation latency (msec)</td>
<td>1.6 (0.9)</td>
<td>2.1 (0.9)</td>
<td>2.0 (0.8)</td>
<td>1.9 (1.0)</td>
<td>2.3 (0.9)</td>
</tr>
<tr>
<td>Recovery time (msec)</td>
<td>0.5 (0.4)</td>
<td>0.8 (0.4)</td>
<td>1.2 (0.4)*</td>
<td>0.3 (0.3)</td>
<td>0.6 (0.2)</td>
</tr>
<tr>
<td>Duration of reflex (msec)</td>
<td>4.3 (1.7)</td>
<td>4.8 (1.9)</td>
<td>6.3 (2.1)*</td>
<td>4.2 (1.8)</td>
<td>6.6 (1.6)</td>
</tr>
<tr>
<td>RAIR amplitude (%) relaxation</td>
<td>77 (36)</td>
<td>71 (30)</td>
<td>54 (33)</td>
<td>64 (37)</td>
<td>70 (32)</td>
</tr>
</tbody>
</table>

Mean (SD) parameters of the RAIR.

* p<0.05 compared to controls