Intestinal transit in anxiety and depression

D A Gorard, J E Gomborone, G W Libby, M J G Farthing

Abstract

Background—Patients with anxiety and depression often have bowel symptoms. Until now, studies investigating a link between altered bowel habit and psychological illness have focused on patients with disturbed defecation presenting to gastroenterologists.

Aims—To determine whether patients with anxiety and depression have objective evidence of abnormal intestinal transit irrespective of any bowel symptoms.

Methods—21 psychiatric outpatients fulfilling research criteria for generalised anxiety disorder and/or major depression, and 21 healthy volunteers were studied. Orocaecal transit time (OCTT) was measured by lactulose hydrogen breath test. Whole gut transit time (WGTT) was measured by abdominal radiography after ingestion of radio-opaque markers.

Results—Median (range) WGTT was shorter in patients with anxiety (14 (6–29) hours) than in patients with depression (49 (35–71) hours) (p<0.001), and controls (42 (10–68) hours) (p<0.001). In patients with anxiety, oroocaecal transit time was shorter (60 (10–70) minutes) than in patients with depression (110 (60–180) minutes) (p<0.01), and shorter than in controls (75 (50–140) minutes (p<0.05). The prolongation of transit times in depression compared with controls was not significant. However, WGTT correlated with both the Beck Depression Inventory score (r=0.59, p<0.001) and the depression score of the Hospital Anxiety and Depression scale (r=0.66, p<0.001).

Conclusions—These objective measurements of intestinal transit in affective disorders are consistent with clinical impressions that anxiety is associated with increased bowel frequency, and depressed patients tend to be constipated; mood has an effect on intestinal motor function.

(Gut 1996; 39: 551–555)

Keywords: intestinal transit, anxiety, depression, oroocaecal, constipation.
TABLE I  Psychiatric questionnaire scores in patients with DSM-III-R criteria for major depression or generalised anxiety disorder and in control subjects

<table>
<thead>
<tr>
<th></th>
<th>Patients (n=21)</th>
<th>Controls (n=21)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Anxiety (n=8)</td>
<td>Depression (n=9)</td>
</tr>
<tr>
<td>BDI</td>
<td>11.5 (7-19)</td>
<td>25 (10-41)</td>
</tr>
<tr>
<td></td>
<td>15 (11-17)</td>
<td>12 (7-18)</td>
</tr>
<tr>
<td>HAD-D</td>
<td>5.5 (2-10)</td>
<td>11 (9-18)</td>
</tr>
</tbody>
</table>

Values are median (range). BDI=Beck Depression Inventory; HAD-A=Hospital Anxiety and Depression score: anxiety subscale; HAD-D=Hospital Anxiety and Depression score: depression subscale.

WHOLE GUT TRANSIT TIME
The WGTT was measured by a radio-opaque marker technique using a single abdominal radiograph. Subjects swallowed 20 radio-opaque polyvinyl chloride markers on three consecutive mornings at 0900 hours. On the fourth morning, a plain abdominal radiograph was taken at 0900 hours. The WGTT was calculated by multiplying the total number of markers seen by 1.2 as previously described.12 Segmental colonic transit times were calculated from the numbers of markers seen within the left colon, right colon, and rectosigmoid regions.

OROCOEAL TRANSIT TIME
After an overnight fast and 20 ml mouth rinse with 0.2% w/v chlorhexidine gluconate, subjects ingested 20 ml (13-4 g) lactulose. End expiratory breath samples were collected before and at 10 minute intervals after lactulose ingestion. Hydrogen concentration in each breath sample was determined using an electrochemical detector (GMI Medical Ltd, Renfrew, UK). The OCTT was defined as the time elapsed between lactulose ingestion and a sustained (>10 ppm above baseline) rise in breath hydrogen.

BOWEL SYMPTOMS
Subjects completed an abridged version of a validated bowel symptom questionnaire.16 This asked them to describe their weekly frequency of defecation as ≤1, 2, 3-4, 5-8, 9-12, 13-16, 17-21, 22-26, or >26, and enquired about the presence of abdominal pain and Manning criteria15 for diagnosing irritable bowel syndrome.

RESULTS
Twenty one patients (five men, median age 27, range 17-45 years) with DSM-III-R criteria for major depression, or generalised anxiety disorder, or both were recruited. Eight had anxiety, nine had depression, and four had research criteria of both generalised anxiety and depression. Twenty one healthy volunteers (six men, median age 24, range 19-45 years) served as the control group.

PSYCHOLOGICAL SCORES
Table I shows the psychological scores on the BDI, HAD-A, and HAD-D. On the BDI, a score of 0-10 is rated as normal, 11-16 is mild depression, 17-20 is borderline clinical depression, 21-30 is moderate depression, and greater than 30 is severe depression. With either of the HAD subscales, a score of 0-7 is considered normal, 8-10 is mild anxiety or depression, and a score above 10 is definite anxiety or depression, up to a maximum of 21.

TRANSIT TIMES
Median (range) WGTT was 42 (10-68) hours in the control group, 14 (6-29) hours in patients with generalised anxiety disorder, 49 (35-71) hours in patients with depression, and 34 (18-66) hours in patients fulfilling DSM-III-R criteria of both anxiety and depression (Fig 1). The WGTT was shorter in patients with anxiety than in patients with depression (p<0.001), and shorter than in controls (p<0.001). The difference in WGTT between depressed patients and control subjects did not reach statistical significance (p=0.09). Table II shows segmental colonic transit times, demonstrating that the faster WGTT in anxiety occurred in all colonic regions.

The OCTT was 75 (50-140) minutes in the control group, 60 (10-70) minutes in patients with anxiety, 110 (60-180) minutes in patients with depression, and 70 (60-90) minutes in patients with both anxiety and depression (Fig 2). In patients with anxiety, OCTT was shorter than in patients with depression (p=0.01), and shorter than in controls (p=0.05). The prolongation of OCTT in depressed patients compared with controls did not reach significance (p=0.08).

In the whole group of psychiatric patients, there was a correlation between WGTT and

TABLE II  Segmental colonic transit times in controls and in patients with generalised anxiety or major depression

<table>
<thead>
<tr>
<th></th>
<th>Right colon transit time (h)</th>
<th>Left colon transit time (h)</th>
<th>Rectosigmoid transit time (h)</th>
<th>Whole gut transit time (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Controls (n=21)</td>
<td>11.2 (2-27)</td>
<td>14 (0-37)</td>
<td>15 (4-32)</td>
<td>42 (10-68)</td>
</tr>
<tr>
<td>Anxiety (n=8)</td>
<td>4 (0-13)</td>
<td>2 (0-11)</td>
<td>4 (0-10)</td>
<td>14 (6-29)</td>
</tr>
<tr>
<td>Depression (n=9)</td>
<td>16 (1-29)</td>
<td>14 (0-39)</td>
<td>11 (2-29)</td>
<td>49 (35-71)</td>
</tr>
</tbody>
</table>

Values are median (range).

Figure 1: Whole gut transit times in patients with DSM-III-R criteria of generalised anxiety disorder, or major depression, or both, and in controls.
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Figure scores on p<0.001).

Bowel symptoms

In the patients with anxiety, the median number of bowel movements per week was 13 to 16, and ranged from 5–8 to 22–26. In patients with depression the median frequency of defecation per week was lower at 3–4, and ranged from 1 to 17–21 (p<0.001). In the control group, the median number of weekly bowel movements was 5–8, with a range from 2 to 17–21. Using a definition of recurrent abdominal pain and three or more Manning criteria, three of the eight patients with anxiety qualified for a diagnosis of irritable bowel syndrome. Six of the nine patients with depression and three of the four patients with both anxiety and depression qualified for a diagnosis of irritable bowel syndrome. Thus within the entire group of psychiatric patients, 12 of 21 (57%) had irritable bowel syndrome. By contrast, two of the 21 controls (10%) fulfilled criteria for diagnosing irritable bowel syndrome (p<0.05).

Discussion

These objective measurements of intestinal transit in patients with affective disorders show that mood disturbances are associated with altered OCTT and WGTT. The WGTT consists predominantly of colonic transit time. Although OCTT comprises gastric emptying as well as small intestinal transit time, small intestinal transit is the same as OCTT when using lactulose alone. Therefore differences of OCTT in this study are likely to be due to differences in transit through the small intestine.

Patients with anxiety and/or depression often have somatic symptoms including disturbed defecation. Thus constipation is found in almost a third of depressed patients, but there have been no previous objective studies to support altered intestinal transit in psychiatric patients. Tucker et al, however, did show that personality factors influence stool output in healthy men. Positive outgoing personality traits were associated with increased stool output. Other studies linking psychological factors and bowel habit have focused on patients presenting with disturbed defecation. There is increased psychiatric disturbance in patients complaining of constipation.

However, increased psychopathology is not universally reported in patients attending hospital with constipation, and in some studies constipated patients with normal transit had increased psychomorbidity, but constipated patients with slow transit did not. A link between disturbed bowel habit and psychological factors is also strengthened by the increased psychomorbidity seen in patients with irritable bowel syndrome attending hospital.

Our study differs from those above because patients with psychiatric illness were studied.
rather than patients with bowel symptoms seen by gastroenterologists or surgeons. In the only other study investigating transit in patients with psychiatric disturbance, patients had presented with gastrointestinal symptoms. Whole gut transit times were shortened in anxiety and prolonged in depression, in keeping with the findings of the current study, but no correlations were found between transit times and psychiatric morbidity scores. By contrast, in our study, patients were all psychiatric outpatients and were not seeing gastroenterologists.

Intestinal transit (both WGTT and OCTT) was faster in anxious patients than in the depressed and control groups. Our finding of shortened OCTT in anxiety is in keeping with the finding that short periods of laboratory induced stress can shorten small intestinal transit in healthy volunteers. We also showed that patients with anxiety have accelerated WGTT. This is consistent with the description of some patients with functional bowel symptoms as having ‘nervous diarrhoea’. Depression, on the other hand, tended to slow transit, but the prolongation of OCTT and WGTT in depressed patients did not reach significance. A significant difference may have been reached if larger numbers of depressed patients were studied. However, there are practical and ethical difficulties in recruiting psychiatrically ill patients who are not taking any drugs, into a study of this nature. Patients requiring urgent drug treatment or admission to hospital were not eligible. Patients had to be outpatients because relative immobilisation in hospital might slow transit compared with the control group – however, the effects of exercise and immobilisation on bowel transit are disputed. Although the depressed patients did not have significantly greater transit times, there were significant positive correlations between WGTT and scores on both the BDI and the HAD-D. These findings provide, for the first time, evidence for an association between severity of depression and colonic inertia. Mechanisms (autonomic neural connections or hormonal) by which mood can alter colonic motor function remain unknown.

Assessment of bowel symptoms was a minor part of this study, yet, interestingly, it showed that the faster intestinal transit of anxious patients was associated with more frequent defecation, whereas the slower transit of the depressed group was associated with less frequent bowel actions. Other studies have shown that frequency of defecation is not correlated with colonic transit time. Symptoms compatible with a diagnosis of irritable bowel syndrome were present in more than half of the psychiatric patients, and were most prevalent in patients with depression. This contrasts with the prevalence of irritable bowel syndrome in the general population of about 20%, but is consistent with studies reporting a prevalence of irritable bowel syndrome in patients with anxiety and depression of about 30–40%.

Our findings are consistent with clinical impressions that anxiety is associated with increased bowel frequency, and depressed patients tend to be constipated. The transit differences between anxious and depressed patients and the correlation of depression scores with prolonged transit show an effect of mood on bowel motility. Just as acute perturbations of the central nervous system by stress and emotion acutely alter intestinal motility, so more chronic perturbation of the central nervous system by affective disorder also influences intestinal motor function.

This work was presented to the 1995 autumn meeting of the British Society of Gastroenterology and published in abstract form (Gut 1995; 37 (suppl 2): A47).

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