How well does stool form reflect colonic transit?

L P Degen, S F Phillips

Abstract

Background—Watery stools are equated with rapid and hard stools with slow intestinal transit; however, the relation between stool form and transit through specific regions of the gut is not clear cut. In addition, more information is needed on interindividual variability of these measurements.

Aim—To examine the relations between stool form and gastric emptying, small bowel and colonic transit.

Methods—Regional gut transit was assessed scintigraphically and segmental colonic transit was also quantified by radio-opaque markers. On two occasions, 32 healthy volunteers (12 men, 20 women) were studied, women during the follicular and luteal phases of menstruation, men twice within a similar four week period. Diets were standardised and stool form was recorded on a seven point scale.

Results—Women had significant harder stools; hard stools were correlated significantly with slow transit and loose stools with fast transit through the colon.

Conclusions—Stool form could not be related to gastric emptying or small bowel transit.

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Keywords: colonic transit, stool form.

Many persons with functional bowel disorders pass stools that vary greatly in form and consistency and, moreover, correlations among stool frequency, faecal form, and gastrointestinal transit have been suggested.1, 2 Most reports have examined stool form and total gut transit, assessed by the faecal excretion of radio-opaque markers; however, there are reasons for proposing that colonic transit in itself may be an important determinant of stool form.3, 4 We have described conditions under which colonic transit was manipulated experimentally; colonic transit and stool form were closely related.5 We wished to extend these observations by measuring regional colonic transit through the unprepared bowel of healthy people who maintained diaries of stool form. We also sought to describe interindividual variability in transit and stool form, by studying each person twice under standardised conditions.

Methods

Experimental subjects

Twelve women and 20 men, aged between 19 and 45 years, were recruited by public advertisement. All considered themselves healthy and, especially, none complained of gastrointestinal symptoms or had a history of gastrointestinal disease or abdominal surgery other than appendicectomy or herniorrhaphy. Functional bowel diseases were excluded specifically using the criteria of Manning.6 Any symptoms of acute infections or use of drugs thought to change gastrointestinal function were reasons for exclusion. Smoking habits, and alcohol and coffee consumption were assessed by standardised questions. After discussion of the procedure in detail, written consent was obtained for a protocol approved previously by the Institutional Review Board and the Radiation Control Committee of Mayo Clinic. All female volunteers had a negative plasma β-human chorionic gonadotropin pregnancy test no longer than 48 hours before each scintigraphic study.

Experimental procedures

Volunteers ingested a weight maintaining diet that was based on the Harris-Benedict equation,7 with adjustment for daily physical activity. All were asked to avoid unusually intensive physical activity. Meals were provided by the Mayo General Clinical Research Center and their composition was normalised to 53% carbohydrate, 17% protein, and 30% fat. Fibre intake was standardised to 15 gram per day, consisting of 60% (9 gram) water insoluble and 40% (6 gram) water soluble fibre.8

On each of three days, volunteers ingested at 9 am a capsule containing 24 radio-opaque markers (SITZMARKS, Lafayette Pharmacol, 4200 South Huluken Street, Fort Worth, TX 76109). At 9 am on the fourth day an abdominal x ray was taken to assess the location of radio-opaque markers.9 On day 4 also, at 7 am, after fasting since midnight, the scintigraphic transit study began.

Women had two studies corresponding to the menstrual cycle: one was on day 7–10 (=follicular phase) and one on day 21–24 (=luteal phase). Men had the study repeated at equal intervals as did the women; one assessment was followed by a second within 14–17 days. Immediately before each transit study, a blood sample was drawn in women to measure the concentration of progesterone and oestradiol. At the beginning of each scintigraphic study, all participants judged their physical activity in the past week according to the Harvard Alumni Activity Survey questionnaire10 and completed the self report inventory SCL-90-R11 to reflect their psychological status for the past week.

Volunteers recorded the date and time of each bowel movement and scored its consistency (Table I). Scoring was by a modified analogue table,5 as described originally by

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TABLE 1 Description of bowel movements

<table>
<thead>
<tr>
<th>Score</th>
<th>Description*</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Separate, hard lumps – like nuts</td>
</tr>
<tr>
<td>2</td>
<td>Sausage-shaped and lumpy</td>
</tr>
<tr>
<td>3</td>
<td>Sausage-shaped, cracked surface</td>
</tr>
<tr>
<td>4</td>
<td>Sausage or 'snaky', smooth, soft</td>
</tr>
<tr>
<td>5</td>
<td>Soft blobs, clear cut edges</td>
</tr>
<tr>
<td>6</td>
<td>Fluffy pieces, ragged edges, 'mushy'</td>
</tr>
<tr>
<td>7</td>
<td>Watery, no solids</td>
</tr>
</tbody>
</table>

*Modified from the descriptions of O'Donnell et al.1

O'Donnell et al.1 Women recorded these for one menstrual cycle, men for 28 days.

Procedure
Gastric, small bowel and colonic transit was measured by the non-invasive scintigraphic method developed in our laboratory.12–14 Briefly, polystyrene Amberlite 120-IR-Plus resin pellets (average diameter 1 mm; range 0.5–1.5 mm) were labelled with 100 μCi of 99mTcCl3.13,14 A capsule filled with approximately 0.5 g pellets, and coated with one layer of methacrylate, was given to the fasting volunteers. As shown already,3,4,12 this capsule could be expected to dissolve in the ileocaecal region. It was therefore used to quantify colonic transit. External radioactive markers were placed over both anterior superior iliac spines to estimate the location of the capsule. As soon as the radiolabelled capsule passed into the small bowel, a breakfast was eaten within five minutes. It consisted of two scrambled eggs, one slice of whole wheat bread, and skimmed milk (35% protein, 52% carbohydrate, 13% fat, 219 kcal). The scrambled eggs were mixed and cooked with 1 mCi of 99mTc-labelled Amberlite 410 resin pellets (average diameter 1 mm) to a firm consistency, to provide a solid medium.

Four hours after breakfast, a standardised non-radiolabelled lunch (chicken, potato, butter, tapioca pudding, and water; 535 kcal) and eight hours after breakfast a dinner (steak, salad, dessert; 21% protein, 49% carbohydrate, 30% fat; 561 kcal) was consumed. During the study volunteers were permitted normal physical activity.

Gammacamera imaging
Gammacamera imaging started immediately after the radiolabelled breakfast was eaten, and used with a large field of view gammacamera with a medium energy, parallel hole collimator (GE Starcam, General Electric, Milwaukee, WI). Anterior and posterior images were acquired with the subject erect. For the 99mTc counts a 140 keV and for the 111In counts a 245 keV energy window (each with ±20% window) was utilised. The estimated whole body dose equivalent was 130 mRem.

Using variable regions of interest, radioactivity was quantified in the stomach and ascending colon for 99mTc and in four regions of the colon (ascending, transverse, descending, rectosigmoid) for 111In.12 The geometric means of the counts obtained from anterior and posterior images were calculated for each region and then corrected for radionuclide decay. The downscatter of 111In into the 99mTc window was adjusted. For two days, stools were collected and the radioactivity for 111In counts was assessed and corrected for decay.

Colonic transit time measured by radio-opaque marker method
The localisation of the radio-opaque markers on the abdominal film taken 24 hours after ingestion of the last radio-opaque markers were related to bony landmarks and gaseous delineations.15 Markers located to the right of the vertebral spinous processes above a line from the fifth lumbar vertebrae to the pelvic outlet were assigned to the right colon. Markers to the left of the vertebral spinous process and above an imaginary line from the fifth lumbar vertebrae to the anterior superior iliac crest were allocated to the left colon. Markers inferior to a line from the pelvic brim on the right and the superior iliac crest on the left were assigned to the rectosigmoid and rectum.15 However, if bowel outlines clearly showed a pelvic caecum, an unusual transverse colon or a large sigmoid loop above the fifth lumbar vertebrae, markers were judged to be in the anatomic segment based on the bowel silhouette.

Data analysis
Gastric emptying was assessed by the gastric lag time, post-lag emptying rate, and the half emptying time (T1/2). The gastric lag time (min) was the time taken for 10% of radioactivity to empty from the stomach.16 The gastric post-lag emptying rate (%/min) was calculated by the slope of linear regression of data points immediately beyond the lag time until 90% of the radiolabel had emptied from the stomach.14

Small bowel transit time (min) was calculated by subtracting the time for 10% of the radio-labelled breakfast to empty from the stomach from the time taken for 10% to enter the colon.13

Colonic transit was evaluated by the geometric centre of counts in the colonic regions of interest (ROI). The geometric centre was the weighted average of proportions of counts in four designed ROI of the colon.3 The regions, designated by numbers 1–4 as weighting factors were the ascending, transverse, descending, and rectosigmoid colons. The stool was designated as the fifth ROI. The proportion in each region was multiplied by the weighting factor and the sum calculated. A low geometric centre indicated that most radio-label was closer to the caecum, whereas a high value indicated that the major part of the radiolabel was closer to the stool.

Colonic transit time measured by radio-opaque marker method
The total number of all markers for each colonic segment was taken as the mean colonic transit time for that segment. The mean total
colonic transit time was the sum of the mean segmental transit times.  

Stool form (Table 1) was expressed as the integer median consistency of all stools passed during each of the two standardised study periods. A median was taken to eliminate the effects of day to day variability and the possibility of a skewed distribution. Subsequently, we grouped the stool consistency into loose, intermediate, and hard. Scores 5–7, describing more watery motions, we defined as loose; scores below 4 were considered hard stools.

Serum concentrations of progesterone and oestradiol were assessed with enhanced luminiscence and radioimmunoassays, respectively. A concentration of progesterone above 2 ng/ml was regarded as consistent with the luteal phase, values below 0.7 ng/ml with the follicular phase.

Physical activity score, SCL-90-R score, smoking habits, alcohol, and coffee consumption

Energy expenditure was expressed as total kilocalories/week, and the psychological symptom scores for the primary symptom dimensions and global indices of distress were rated. Smoking habits were quantified by pack years of actual consumption, alcohol consumption by units a week (1 unit beer, wine or spirits=10 g alcohol), and coffee by cups a day.

Statistical analysis

The colonic transit data were expressed as box and whisker plots showing the median, the interquartile interval, and the total range.

Sex distribution was examined with the \( \chi^2 \) test. Correlations between stool form and gut transit were by linear regression analysis. Data in different stool form groups were analysed by analysis of variance, and for pairwise comparisons, with the Newman-Keuls procedure. To detect any possible dependent variables that may have influenced stool form, a multiple analysis of variance was used. Significance was declared at \( \alpha<0.05 \).

Results

Characteristics of the groups

The mean age in both groups was 29 years (range, women: 19–44; men: 21–45). Although the body mass index did not differ significantly (mean (SEM) women: 23.97 (0.95); men: 25.48 (0.63)), weight and height were significantly different (p<0.001). Men were heavier (84 (2) v 68 (3 kg)) and taller (182 (2) v 168 (2 cm)). Although men consumed slightly more calories, the values were not significantly different between the sexes or between the two studies. Women: study 1: 2845 (420), study 2: 2562 (288) kcal/day, men: 3384 (570) and 3296 (479) kcal/day. Estimated smoking, coffee or alcohol consumption were not significantly different.

Fluctuations of hormonal concentrations between the follicular and luteal phases were significant for progesterone and oestradiol (p<0.01 and p=0.018, respectively). Progesterone measured in the follicular phase was 0.57 (0.07), in the luteal phase 6.07 (1.11) ng/ml. Oestradiol concentrations were 79-67 (11.09) and 137-33 (16.43) pg/ml, respectively.

Stool form

Characteristics of the stools varied widely among subjects and transit of radio-opaque markers was highly correlated with stool form (y=4.7 - 0.04x, p<0.001; Fig 1). Stool form also varied considerably between the two study periods in some subjects. Although the median difference for duplicate studies was essentially zero in both sexes (Fig 2) there was wide variability within subjects, with women and men having the same degree of scatter.

Stool form was subsequently grouped into three major categories, hard (scores 1–3), intermediate (score 4), and loose stools (scores 5–7). Using these groupings, women had significantly harder stools in at least one observation period than did men: hard=8 men, 7 women; intermediate=4 men, 5 women,
loose = 8 men, 0 women; p = 0.04, χ² = 6.59). Women’s stool form did not change significantly during the menstrual cycle.

Colonic transit, when expressed for both sets of studies as the geometric centre at 6 hours (data not shown) and 24 hours (Fig 3A), was significantly different when persons with hard and loose stools were compared (p = 0.01). By the radio-opaque marker method, persons with hard stools had significantly slower total colonic transit (Fig 3B, p < 0.01). Segmental transits (data not shown) in the right colon and rectosigmoid region were also slower (p = 0.005, p = 0.02, respectively).

Stool frequency was not significantly correlated with stool form. Rates of gastric emptying and small bowel transit did not influence stool form significantly (Table II), though there was a trend for faster small bowel transit to be associated with looser stools. Psychological symptom scores did not change between studies and had no relation to transit times or stool form.

During the entire period of scintigraphy (48 hours), hard stools were significantly associated with lower geometric centres than were those for loose stool (Fig 4). The areas under the curve, which express the geometric centre progression over time, were significantly different (p < 0.001). The intermediate group showed considerable overlap (data not shown).

**Discussion**

Descriptors of stool form have been used to estimate gastrointestinal transit and applied to epidemiological studies of transit in health and patients with gastrointestinal dysfunction.

Others have reported significant correlations between stool consistency and objective measurements by a penetrometer, viscometer or by analysis of water content. Reproducible results require that a single investigator do the scoring and that volunteers themselves record consistency of their stools. As to how precisely descriptor scales of stool form reflect specific transit functions is not clear. Thus, although the correlation of stool form with whole gut transit times was significant, considerable overlap was evident. On the other hand, we reported that colonic transit, when changed experimentally, correlated well with stool form. The present protocol featured standardised conditions in healthy volunteers and used the scintigraphic method developed in our laboratory to assess separately gastric emptying, small bowel transit, and segmental colonic transit. Moreover, intraindividual variations were assessed by repeating the tests in all persons.

Correlations between the seven point descriptor scale and transit were similar to our earlier findings. However, an overriding influence of one colonic segment, specifically the rectosigmoid colon was not confirmed. All parts of the colon exhibited similar qualitative relations with hard and loose stools. Stool consistency did not correlate with any of the indices of upper gastrointestinal transit in these healthy persons. However, in patients with

**TABLE II Stool form and upper gut transit**

<table>
<thead>
<tr>
<th>Stool form</th>
<th>Lag phase (min)</th>
<th>Post-lag emptying rate (%/min)</th>
<th>Transit (min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hard</td>
<td>56 (17)</td>
<td>0.302 (0.177)</td>
<td>211 (82)</td>
</tr>
<tr>
<td>Intermediate</td>
<td>58 (25)</td>
<td>0.355 (0.200)</td>
<td>178 (65)</td>
</tr>
<tr>
<td>Loose</td>
<td>54 (14)</td>
<td>0.419 (0.163)</td>
<td>136 (30)</td>
</tr>
</tbody>
</table>

*p = 0.08

*p = 0.17

*p = 0.11

*p = 0.09

*By analysis of variance. Data shown as mean (SEM).

![Figure 3: (A) Medians, interquartile distributions, and ranges of geometric centres of colonic transit for healthy subjects with hard, intermediate or soft stools. (B) Medians, interquartile distributions, and ranges of total colonic transit by the radio-opaque marker method.](http://gut.bmj.com/)

![Figure 4: Continuous plot of geometric centres of healthy subjects with hard or loose stools. The higher geometric centre values for loose stools represents more distal passage of the isotopic marker, an effect that becomes more pronounced with time.](http://gut.bmj.com/)
functional bowel disease, a significant influence of orocecal transit time, assessed by the breath hydrogen method, on stool consistency has been reported.21 In addition, we have reported an overrepresentation of rapid gastric emptying in patients with non-organic diarrhoea.22

We confirmed that women had significantly harder stools. The mechanism may entail more pronounced prolongation of colonic transit in the distal segments, as suggested by additional findings reported elsewhere.23 In agreement with an earlier report we did not find any systematic variation of stool form or colonic transit23 during the menstrual cycle. Results suggesting an influence of female sex hormones25 may have been influenced by different diets2 or underlying functional bowel disorders in the women studied.26 No other factor in these non-obese, healthy volunteers, such as body mass index, smoking, coffee consumption, physical activity, caloric intake or psychological profile significantly influenced stool form.

In summary, the extremes of the stool form, hard and loose textures, discriminate usefully between slow and fast colonic transit. We could not show any predominant influence of a specific colonic segment. Intrinsic variability of colonic transit is expressed by different stool forms.

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