Human migrating myoelectric complex in relation to gastrointestinal transit of a meal

J L Madsen, K Dahl

Abstract
Feeding interrupts the migrating myoelectric complex in most mammals. This study aimed to assess whether resumption of the migrating myoelectric complex in the human duodenum after eating was related to the gastrointestinal transit of the meal. Five healthy subjects participated in the study. After eating a radiolabelled test meal consisting of mixed liquid and solids, duodenal myoelectric activity and gastrointestinal transit of the meal were determined simultaneously. In spite of considerable variation in entire gastric emptying time between subjects (range 2.5-5.0 hours), significant correlation was found between the completion of gastric emptying and the resumption of duodenal phase III activity within subjects (p<0.01). A new technique for recording the duodenal myoelectric activity was used.

The migrating myoelectric complex is a specific pattern of electrical response activity that occurs cyclically in the small intestine of most mammals in the fasted state.1-4 Migrating myoelectric complexes generally involve the duodenum and propagate distally.4 After eating, the cycle is interrupted for a considerable period, depending on the composition of the meal.5-7 We examined the possible association between gastric emptying and resumption of the duodenal myoelectric complex cycle activity, which has not been formally studied before.

Materials and methods

RADIOLABELLED TEST MEAL
The test meal consisted of 400 g of mixed solids and liquid (80 g bread, 30 g cheese, 10 g butter, 50 g yoghurt, 230 g water) that had an energy value of 1600 kJ. Two hundred mg of cellulose fibre labelled with 40 MBq of technetium-99m was added to the yoghurt. The use of this radiolabelled marker in solid foods has been validated in a recent study.4

DUODENAL MYOELECTRIC RECORDINGS
A new electrode technique was developed to record the duodenal myoelectric activity. The myoelectric signal was recorded with a single bipolar surface electrode (length 21 mm, diameter 3 mm, weight 0.6 g) consisting of two stainless steel leading-off rings (width 7 mm) which were mounted on the ends of a fitted teflon cylinder and connected to two recording wires (polyfile stainless steel, teflon coated, highly flexible, diameter 0.3 mm) (Fig 1).

On the evening before the start of the investi-
Figure 2: Representative tracing of duodenal phase III activity.

obtained using a 15% energy window over the 140-keV $^{99m}$Tc photopeak. Each imaging period lasted two minutes. Imaging was repeated at 30 minute intervals until no activity could be detected in the small intestine. Subjects were sitting between imaging. Data were stored on a computer for later analysis.

CALCULATIONS

Regions of interest for integration of activity were delineated manually around the stomach, the small intestine, and colon on each image. The obtained count rates were corrected for physical decay. The geometric means of corresponding anterior and posterior count rates were used as attenuation corrections. The corrected count rates were then converted into fractions of the maximum count rate. For each gastrointestinal segment, these fractions were plotted against time. The moment that the first phase III migrating myoelectric complex occurred was marked out on these curves. The corresponding amount of activity in each segment was then read by interpolation from the curves. In addition, the entire gastric emptying time and the corresponding phase III interruption time were compared by linear regression. The obtained correlation coefficient was tested against 0.

The percentage distribution of radiolabelled cellulose fibre in the various gastrointestinal segments of five subjects at resumption of duodenal phase III activity after a meal. (For comparison with Fig 3, subjects are ranked according to phase III interruption time. Rank 1 means shortest interruption time.)

<table>
<thead>
<tr>
<th>Subject</th>
<th>Stomach</th>
<th>Small intestine</th>
<th>Colon</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>2</td>
<td>88</td>
<td>10</td>
</tr>
<tr>
<td>2</td>
<td>0</td>
<td>100</td>
<td>0</td>
</tr>
<tr>
<td>3</td>
<td>0</td>
<td>41</td>
<td>59</td>
</tr>
<tr>
<td>4</td>
<td>0</td>
<td>79</td>
<td>14</td>
</tr>
<tr>
<td>5</td>
<td>0</td>
<td>45</td>
<td>55</td>
</tr>
</tbody>
</table>

Figure 3: The entire gastric emptying time of five subjects plotted against the corresponding phase III interruption time after a meal. The regression line is drawn.

Results

Results of the gastrointestinal transit measurements and the myoelectric recordings are shown in the Table and in Figure 3.

Linear regression showed a significant correlation between the entire gastric emptying time and the corresponding phase III interruption time (correlation coefficient 0.96, p<0.01, slope of regression line 1.04, intercept on y-axis 0.04). Obviously, a correlation did not exist between resumption of phase III activity and the corresponding content of radiolabelled marker in the small intestine or colon.

Discussion

The migrating myoelectric complex cycle or its motor correlate in the human small intestine is interrupted by a meal. The duration of this interruption is proportional to the caloricific value of the meal and the composition of the food ingested. Lipids interrupt the migrating myoelectric complex cycle for longer than carbohydrates and proteins of equal caloricific value. As gastric emptying of larger meals is delayed compared with that of smaller meals and lipids are emptied more slowly than non-lipids, the interruption of the migrating myoelectric complex cycle may be related to the gastrointestinal transit of the meals. Such a relation has not yet, however, been described precisely.

This study showed a very close temporal association between the end gastric emptying and the resumption of the duodenal migrating myoelectric complex cycle. The cause of this association is not clear as the control mechanisms involved in the migrating myoelectric complex cycle have not been fully explained. Our results, however, can probably be explained by the hypothesis that when duodenal chemoreceptors are no longer stimulated, cycle activity controlled by the enteric nervous system is unblocked.

The considerable variation in the time that the first phase III activity occurs after a meal reported earlier may, as in this report, reflect the variation in gastric emptying kinetics.
The physiological role of the migrating myo-electric complex and the corresponding motor activity is still unknown. Our results possibly support the thesis that the cycle is important in propelling residual food along the small intestine since this segment contained considerable amounts of the food marker when the migrating myoelectric complex cycle recurred after feeding.

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