The vagus, the duodenal brake, and gastric emptying

M. SHAHIDULLAH, T. L. KENNEDY, AND T. G. PARKS

From the Department of Surgery, Queen’s University, and Royal Victoria Hospital, Belfast, N. Ireland

SUMMARY  It has been suggested that an intact vagal supply is essential for the normal function of the receptors in the duodenum and proximal small bowel, which influence the rate of gastric emptying. This paper reports the effect of vagal denervation on gastric emptying and also examines the site and mode of action of receptors in the proximal small bowel.

It has been demonstrated in the dog that most, if not all, the receptors controlling gastric emptying lie in the proximal 50 cm of the small bowel. Following truncal vagotomy the emptying time of each instillation increased significantly and the differential rate of emptying of different instillations remained unchanged. The proximal 50 cm of small bowel was capable of differentiating between different instillates even after selective extragastric vagotomy, in which the duodenum was vagally denervated and, therefore, duodenal braking receptors function independently of vagal innervation.

It is well established that meals of different composition empty from the stomach at different rates and that this is due to regulation by receptors in the duodenum (Hunt, 1957; Hunt and Knox, 1968a). Differences of opinion exist as to the effects of vagotomy and drainage on the rate of gastric emptying; some workers report that the rate of gastric emptying is increased (Goodall, 1966; George, 1968; Aylett, Wastell, and Wise, 1969; McKelvey, 1970) but others find that it is delayed (Isaac, Ottoman, and Weinberg, 1950; Dragstedt and Woodward, 1951; Buckler, 1967; Tinker, Kocak, Jones, Glass, and Cox, 1970).

In this study we have tried to identify the site of the braking receptors and also to determine the role of the vagus in the regulation of gastric emptying when the pylorus is intact.

Materials and Methods

Two different experimental models were prepared.

1  Gastric and duodenal fistulae were produced by inserting two Gregory cannulae, one into the stomach and one into the duodenum (fig 1).

2  In another group of animals the duodenum was divided immediately beyond the intact pylorus which was then implanted end to side 50 cm further down the small bowel. The duodenal stump was closed. At a second operation two cannulae—one into the stomach and one into the duodenum—were inserted in order to produce gastric and duodenal fistulae (fig 2).

Five dogs belonging to model 1 underwent truncal vagotomy, i.e., both right and left trunks of the vagus were divided transthoracically. Two dogs belonging to model 2 underwent extragastric selective vagotomy: hepatic and coeliac branches were divided but all other gastric branches, including the nerves of Latarjet, were preserved (fig 3).
M. Shahidullah, T. L. Kennedy, and T. G. Parks

Fig 2 Model 2: Bypass experimental model in which the intact pylorus is implanted 50 cm down the small intestine.

The gastric fistula facilitated the carrying out of measurements of the rate of gastric emptying. The duodenal fistula was used to introduce different substances into the duodenum and proximal small bowel during the study, eg, deionized water, 250 mM hydrochloric acid, 10% dextrose, and 5% fat. The double-sampling dye dilution technique of George (1968) was used throughout to measure the rate of gastric emptying. In the case of the meal containing fat, a special procedure was adopted before optical density was read. It was ensured that specimens were very thoroughly mixed by means of vigorous shaking immediately before removing 4 ml to be made up with 10 ml buffer and distilled water to 100 ml. The fat was extracted from this final solution by mixing equal volumes of the solution and diethyl ether. This mixture was very thoroughly shaken, allowed to sit at 4°C for 30 minutes, then it was centrifuged at 4°C. The ether layer and the meniscus between the ether and aqueous layers were removed and the optical density of the residue was read. Calculations were carried out in the usual manner with these results.

Emptying studies were carried out between six and 10 weeks after gastric and duodenal fistulae had been prepared.

All the studies were performed on healthy, conscious dogs. The dogs were kept fasting for about 24 hours before any emptying study was carried out. During the tests, dogs were placed in a Pavlov stand on an elevated table. These dogs had been trained for some days previously to stand in the frame before actual emptying tests were carried out. The cap of the gastric cannula was removed and the stomach allowed to drain on its own. A rubber teat with a small hole at its centre for the passage of the Levin

Fig 3 Diagrammatic representation of selective extragastric vagotomy.

Fig 4 A Levin tube in situ, introduced via a rubber teat and gastric cannula.
tube was placed at the open end of the cannula to prevent fluid leaking from the stomach (fig 4). A Levin tube size 14 was then introduced into the stomach through the gastric cannula, making sure that just the tip of the tube was lying in the stomach. Another Levin tube was introduced in a similar fashion through the duodenal cannula.

The stomach was washed out with 200 ml of deionized water. It was then ensured that the stomach was emptied as completely as possible by means of careful aspiration with a syringe, moving the tube around the stomach. At the end of this operation the Levin tube and the rubber test were removed and the stomach was allowed to drain on its own, the cannula being at the most dependent part of the stomach and the dog kept in the erect position.

**Results**

In one dog with an intact gastroduodenal junction five emptying studies using 600 ml of water showed the high degree of reproducibility of the test (table I).

<table>
<thead>
<tr>
<th>Rate of Emptying (min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
</tr>
<tr>
<td>40</td>
</tr>
</tbody>
</table>

**Table I  Rate of emptying of water from the stomach**

*Daily variation in a single dog

In seven dogs with an intact gastroduodenal junction emptying studies were carried out using 600 ml of fluids. It was shown that the rate of emptying of different instillates varied greatly. The mean emptying times in minutes were as follows: deionized water 44, hydrochloric acid 85, dextrose 126, and fat 144. There were highly significant differences in the rates of emptying between different pairs of instillates in the intact stomach. Detailed results are shown in table II.

<table>
<thead>
<tr>
<th>Fluids in Stomach</th>
<th>Before Vagotomy</th>
<th>After Vagotomy</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Number of Studies</td>
<td>Mean Time (min) ± SEM</td>
</tr>
<tr>
<td>Water</td>
<td>26</td>
<td>44 (± 1.78)</td>
</tr>
<tr>
<td>250 mM HCl</td>
<td>8</td>
<td>85 (± 4.35)</td>
</tr>
<tr>
<td>10% dextrose</td>
<td>18</td>
<td>126 (± 3.18)</td>
</tr>
<tr>
<td>5% fat</td>
<td>15</td>
<td>144 (± 5.93)</td>
</tr>
</tbody>
</table>

**Table II Emptying time before and after truncal vagotomy**

Truncal vagotomy was performed in five dogs belonging to model 1; six weeks after vagotomy emptying was then studied using the same instillates. During insulin hypoglycaemia no animal had an acid output of more than 20-equiv/l or 10 m-equiv if there was no basal acidity, i.e., all tests were negative according to Hollander's criteria.

The results of emptying tests are shown in table II and figure 5. It can be seen that the emptying time was significantly altered by interruption of the vagi. After truncal vagotomy the emptying time for each substance became significantly longer, yet the proportional differences between the effects of different substances were maintained.

In eight dogs belonging to model 2, i.e., with the pylorus implanted into the small bowel, gastric emptying studies were carried out putting 500 ml of different substances into the stomach with 100 ml of deionized water introduced through the duodenal cannula (fig 6). With water in the duodenum the
different types of instillates all emptied from the stomach at similar rates: for example, water took 54 minutes and dextrose 53 minutes (table III).

<table>
<thead>
<tr>
<th>Fluid in Stomach</th>
<th>Agent in Small Bowel</th>
<th>No. of Studies</th>
<th>Mean Time (minutes)</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal saline</td>
<td>Water</td>
<td>5</td>
<td>54</td>
<td>3.27</td>
</tr>
<tr>
<td>Water</td>
<td>Water</td>
<td>12</td>
<td>54</td>
<td>1.86</td>
</tr>
<tr>
<td>250 mM HCl</td>
<td>Water</td>
<td>12</td>
<td>53</td>
<td>2.33</td>
</tr>
<tr>
<td>10% dextrose</td>
<td>Water</td>
<td>12</td>
<td>53</td>
<td>2.57</td>
</tr>
<tr>
<td>5% fat</td>
<td>Water</td>
<td>11</td>
<td>54</td>
<td>1.48</td>
</tr>
</tbody>
</table>

Table III Emptying time in a bypass model

Further studies were carried out in four dogs belonging to model 2. Five hundred ml of water was put into the stomach and at the same time 100 ml of different instillates, ie, water, 250 mM hydrochloric acid, 10% dextrose, or 5% fat solution were introduced on different days through the duodenal cannula (fig 7). In these dogs the water emptied from the stomach at different rates depending on the nature of the substance introduced through the duodenal cannula. When water was introduced into the small bowel, the water emptied in 54 minutes, but when 250 mM HCl or 10% dextrose or 5% fat was introduced through the duodenal cannula the mean emptying time became 80, 137, and 160 minutes respectively (table IV).

<table>
<thead>
<tr>
<th>Fluid in Stomach</th>
<th>Agents in Small Bowel</th>
<th>No. of Studies</th>
<th>Mean Time (minutes)</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>Water</td>
<td>12</td>
<td>54</td>
<td>1.86</td>
</tr>
<tr>
<td>Water</td>
<td>250 mM HCl</td>
<td>10</td>
<td>100</td>
<td>4.15</td>
</tr>
<tr>
<td>Water</td>
<td>10% dextrose</td>
<td>13</td>
<td>137</td>
<td>4.85</td>
</tr>
<tr>
<td>Water</td>
<td>5% fat</td>
<td>11</td>
<td>160</td>
<td>4.84</td>
</tr>
</tbody>
</table>

Table IV Emptying time in a bypass model

In two dogs belonging to model 2, selective extragastric vagotomy (division of hepatic and coeliac vagal branches) was performed and six weeks later emptying studies were carried out using water, 10% dextrose, and 5% fat. When 500 ml of each of these substances was introduced on different days into the stomach and 100 ml of deionized water was put into the duodenum, the rate of emptying of each instillate from the stomach was similar (table V).

<table>
<thead>
<tr>
<th>Fluids in Stomach</th>
<th>Agents in Small Bowel</th>
<th>No. of Studies</th>
<th>Mean Time (minutes)</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>Water</td>
<td>10</td>
<td>51</td>
<td>2.43</td>
</tr>
<tr>
<td>10% dextrose</td>
<td>Water</td>
<td>7</td>
<td>53</td>
<td>3.21</td>
</tr>
<tr>
<td>5% fat</td>
<td>Water</td>
<td>7</td>
<td>52</td>
<td>3.34</td>
</tr>
</tbody>
</table>

Table V Emptying time following selective extragastric vagotomy in a bypass model

Discussion

As one would expect that the passage of the Levin tube into the stomach through the mouth of healthy conscious dogs would disturb the physiological state of the animals, it was considered preferable to introduce the Levin tube into the stomach through a gastric cannula. Knowing the length of the cannula it was possible to introduce the Levin tube so that just the tip lay in the stomach. Radiological screening was not necessary in order to find out if
The vagus, the duodenal brake, and gastric emptying

the tube was lying in the proper place or if the tube was curled up inside the stomach. The double-sampling dye dilution technique of George (1968) was found to be satisfactory though a modification was needed in cases of meals containing fat. Fluid meals empty in an exponential pattern. The main disadvantage of the test is that it is only applicable to fluid meals, and it can be argued that the evacuation of fluid from the stomach does not reflect how the stomach deals with a normal mixed meal. Using a mixed meal, Griffith, Owen, Kirkman, and Shields (1966) demonstrated an exponential pattern.

Hunt and Spurrell (1951) have shown that there is very little daily variation in the emptying rates of an individual stomach and the present study has confirmed this. This emptying pattern was found to be exponential in the intact dogs corresponding with the findings of Hunt and Spurrell (1951). On the other hand, Tamarit Torres, Enriquez de Salamanca, and Castro-Rial Canosa (1954) found a hyperbolic relationship between the volume of meal remaining in the stomach and time. In the present studies, different substances were used at different concentrations and it was found that normal saline emptied fastest whereas 5% fat solution took longest to empty, confirming previous work which showed that the rate of emptying varies according to the nature of the meal (Hunt, 1957, 1960; Hunt and Knox, 1968b).

Though the function of the pylorus in model 2 may perhaps have been temporarily interfered with due to the operative procedure its nerve supply and vascular supply were preserved very carefully. The anastomosis was carried out in one layer in order not to cause narrowing of the lumen. Functionally the pylorus behaved normally as shown from the figures in table IV, when water instilled into the stomach emptied at different rates depending on the nature of the instillate in the proximal small bowel. It would appear that the pylorus in this animal experimental model was in no way incontinent. Our studies seem to have identified the proximal 50 cm of the small bowel as the main site for receptors controlling gastric emptying. If there was a significant number of receptors more than 50 cm distal to the pylorus, then different instillations into the stomach might be expected to empty at different rates. That this was not the case might be due to the very much slower rate of passage of fluid from the stomach into the jejunum compared with the rate of infusion into the duodenum. This is probably not the explanation, however, as our findings in model 1 dogs with no bypass and an intact pylorus show that different instillations into the stomach do empty at different rates.

Emptying time following vagotomy and drainage is disputed (Buckler, 1967). Tinker et al (1970), who used meals of a more solid nature, showed delayed emptying after vagotomy and drainage. In our series the emptying time for each instillate following vagotomy was significantly prolonged. The ability to differentiate between different instillates was maintained.

Other investigators who have used fluid meals have reported an overall increase in the rate of gastric emptying (Goodall, 1966; George, 1968; Aylett et al, 1969; McKelvey, 1970) which is in conflict with our findings. Apart from species differences the main point of variance is the absence of pyloroplasty or any drainage procedure in our animals. McKelvey (1970) thought that abnormal emptying following vagotomy and drainage was due to loss of duodenal regulation of emptying but could not say whether this was due to the vagotomy or the drainage procedure. It may be that the abnormality of gastric emptying is mainly due to the drainage operation. The superior results following highly selective vagotomy without drainage are probably due to preservation of the pylorus rather than preservation of the vagal supply to the distal part of the stomach and small bowel.

Hunt and his colleagues (1966) have shown that there are duodenal receptors involved in the control of gastric emptying but they have not demonstrated precisely how far down the gut these receptors extend. From our studies on dogs it would appear that most, if not all, the braking receptors lie in the proximal 50 cm of the small intestine. It seems unlikely that there are a significant number of such receptors further distally.

It is concluded that vagal innervation of the stomach is important in the mechanism of gastric emptying but that duodenal innervation is not essential for the function of the braking receptors which appear to work through a hormonal mechanism. It should be stressed that these conclusions relate to canine experiments and are not necessarily applicable to human physiology.

We wish to record our thanks to the Royal Victoria

---

**Table VI Emptying time following selective extragastric vagotomy in new bypass model**

<table>
<thead>
<tr>
<th>Fluid in Stomach</th>
<th>Agents in Small Bowl</th>
<th>No. of Studies</th>
<th>Mean Time (minutes)</th>
<th>SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>Water</td>
<td>10</td>
<td>51</td>
<td>2.43</td>
</tr>
<tr>
<td>Water</td>
<td>10% Dextrose</td>
<td>9</td>
<td>136</td>
<td>3.34</td>
</tr>
<tr>
<td>Water</td>
<td>5% Fat</td>
<td>9</td>
<td>163</td>
<td>4.79</td>
</tr>
</tbody>
</table>

1 Different agents in proximal small bowel
2 Data as in table V
Hospital Group Research Sub-Committee for a grant to one of us (M.S.) from their endowment funds which has made it possible to undertake this work. Professor A. D. Roy has kindly provided facilities for the study in the Department of Surgery, Queen's University, Belfast. We also wish to thank Mr H. O. Nevin, Mr J. White, and Sister E. Crawford for technical assistance, and Mrs S. Leonard and Mrs J. Walker for secretarial help.

References


M. Shahidullah, T. L. Kennedy, and T. G. Parks

*Lancet*, 1, 1244-1245.


