Relationship between undissociated acidity of gastric juice and gastric protein secreted in response to graded doses of pentagastrin in duodenal ulcer patients

T. POPIELA,¹ Z. SZAFRAN, HALINA SZAFRAN, AND M. KOMOROWSKA

From the Division of Surgical Gastroenterology, Department of Surgery I, and the Department of Clinical Biochemistry, Institute of Paediatrics, Nicolaus Copernicus Medical Academy, Krakow, Poland

SUMMARY
Concentrations of free and total hydrogen ions, total protein and pepsin were measured in gastric juice fractions collected during basal secretion and upon stimulation by graded doses of pentagastrin administered intravenously. Undissociated hydrogen ion and non-pepsin protein concentrations were calculated as derived quantities. The studies were carried out in nine patients with duodenal ulcer both before and after truncal vagotomy. It was found that after vagotomy the undissociated hydrogen ion concentration was significantly lower and non-pepsin protein higher than before the operation. No correlation was found between the two quantities both before and after vagotomy. It was concluded that in duodenal ulcer patients either not all non-pepsin protein takes part in buffering of hydrogen ions secreted by parietal cells, or that non-protein buffers play a more important role.

Human gastric juice contains variable amounts of soluble protein, of which a considerable part has no well-defined physiological role. Makhlouf et al. (1970) have suggested that one of the possible functions of non-pepsin protein may be the buffering of hydrogen ions secreted by parietal cells. It is obvious on theoretical grounds that any protein with an isoelectric point higher than the actual pH of gastric juice will have some buffering capacity. This means that some hydrogen ions will be bound by these proteins when, in the process of secretion, they come into contact with parietal fluid and pH changes from the relatively high value of the cell interior to the low value of gastric juice. The buffering capacity of such protein will depend on the number of proton-releasing groups with pKₐ values contained within the range covered by the above-mentioned pH extremes.

Makhlouf et al. (1970) found that, in a healthy human subject, undissociated hydrogen ion concentration was correlated with non-pepsin protein secreted upon stimulation with gastrin II or histamine. Our previous studies showed (Szafran et al., 1976) that, in duodenal and gastric ulcer patients, the concentration of undissociated hydrogen ion was positively correlated with total protein in the course of the basal secretion. No such correlation was observed in these patients for gastric juice secreted upon stimulation with a single dose of pentagastrin or histamine.

The aim of the present study was to evaluate the relationship between the secretion of the total, non-pepsin and pepsin protein and undissociated hydrogen ion concentration in the course of intravenous pentagastrin tests with graded doses of the stimulant carried out in duodenal ulcer patients both before and after surgical vagotomy.

Methods
A group of nine men treated for duodenal ulcer was studied. The mean age of the patients was 36 years (range from 27 to 52 years). Truncal vagotomy with pyloroplasty was applied in all these patients as a basic treatment. Vagotomy was complete in all cases
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as confirmed by the negative results of the insulin test (Popiela et al., 1972). Gastric secretory tests were carried out twice in each patient, first before the operation, and then one to 17 months after vagotomy.

Gastric juice was collected through a gastric tube by applying reduced pressure. The stomach was first emptied from the residuum, whereupon three 10 minute fractions of basal secretion were collected. This was followed by the intravenous infusion of pentagastrin solution (Peptavlon, ICI, England) lasting for three hours. The following doses of pentagastrin were applied consecutively: 0.008, 0.064, 0.256, 1.024, 2.048, and 8.192 μg/kg h. Each dose was infused during 30 minutes. Eighteen 10 minute fractions were collected during the pentagastrin infusion. Any fraction contaminated with bile was rejected. Dental cotton pledges inserted into the mouth and frequently changed were used to prevent contamination of gastric juice with saliva. The volume of each fraction was measured, and the fractions were then centrifuged for 20 minutes at 12,000 × g in a refrigerated centrifuge.

The pH of gastric juice fractions was measured by using the Beckman Expandomatic pH-meter supplied with No. 39042 stomach electrode and No. 40249 Ag/AgCl reference electrode. Instrument calibration was carried out with 0.1 mol/l hydrochloric acid solution (pH 1.1) and pH 2.6 citrate buffer. Free hydrogen ion concentration was calculated from pH values by using appropriate factors as given by Moore (1968). Total hydrogen ion concentration was measured by the titration with 0.01 mol/l NaOH solution to pH 7.4 using the Beckman automatic endpoint titration system. Undissociated hydrogen ion concentration was calculated by subtracting free hydrogen ion concentration from total hydrogen ion concentration.

Protein concentration was determined by the method of Lowry et al. (1951). Bovine albumin (Cohn fraction V, Koch-Light, England) was used as a standard. Pepsin activity was measured by a modification of the Anson method (Rick, 1963). Crystalline pepsin (2 × cryst. from pig gastric mucosa, Koch-Light, England) was used as a standard. Non-pepsin protein concentration was calculated by subtracting pepsin concentration (as expressed in grams per litre) from total protein concentration.

Results

The changes in the concentration of total and free hydrogen ions during intravenous infusion of graded doses of pentagastrin in duodenal ulcer patients both before and after vagotomy are shown in Fig. 1. It can be seen that the changes of free acidity are similar to those of total acidity. As might be expected, the values after vagotomy were significantly lower than those before the operation. In patients before vagotomy there was already a definite increase in hydrogen ion concentration in response to the lowest dose of pentagastrin. The peak value was attained in fraction 9, after which there was no further significant change. After vagotomy the response was slower, but the rise of acid concentration lasted longer (up to fraction 15). As the Table

![Fig. 1 Changes in free and total hydrogen ion concentration in the course of intravenous pentagastrin test performed in nine patients with duodenal ulcer before and after vagotomy. Fractions —2, —1, and 0: basal secretion. Fractions 1 to 18: secretion during pentagastrin infusion. ○ Free hydrogen ions before vagotomy; ▲ free hydrogen ions after vagotomy; O total hydrogen ions before vagotomy; △ total hydrogen ions after vagotomy. Mean values ± SEM are given.](http://gut.bmj.com/)
shows, the dose response to pentagastrin, as expressed by $D_{50}$ value, was 0·62 μg/kg h before the operation and 1·28 μg/kg h after vagotomy. The corresponding $V_{\text{max}}$ values for the total hydrogen ions were 0·65 and 0·19 mmol/min respectively.

The changes of undissociated acidity, representing the difference between total and free hydrogen ion concentrations, are shown in Fig. 2. There was no clear dependence on pentagastrin dose, inasmuch as the changes from fraction to fraction exhibited no definite pattern. On the other hand, there was a clearly visible difference between the values obtained before and after vagotomy. In the latter case the values of undissociated acidity were definitely lower. The differences were statistically significant ($p < 0·05$) for all fractions of basal secretion and for fraction nos. 1, 2, 3, 5, 6, 8, 9, 11, 12, and 18 secreted during pentagastrin infusion.

Figures 3 and 4 show the changes in protein concentration, separate curves being displayed for pepsin, non-pepsin, and total protein. Only pepsin protein increases in response to pentagastrin stimulation. Neither total nor non-pepsin protein changed appreciably with the increase in pentagastrin dose. Only a slight tendency to rise was observed after vagotomy. Nevertheless, the values obtained in patients after vagotomy were higher than before vagotomy, both for total and non-pepsin protein. For non-pepsin protein the difference was statistically significant for fraction nos 5, 6, 8, 9-11, and 13-18, and for total protein for fraction nos. 9, 11, and all the following numbers. This result was rather unexpected in view of the markedly lower values of undissociated acidity observed after vagotomy.

Another striking feature was found when the correlation between undissociated acidity and non-pepsin protein was tested for all fractions of gastric juice obtained during basal secretion and pentagastrin-induced secretion. There was no significant correlation between the two parameters, as reflected by the very low values of correlation coefficient: for basal secretion—0·21 before vagotomy and 0·19 after vagotomy, and for stimulated secretion 0·14 and 0·13 respectively.

**Discussion**

The results of this study revealed that, in duodenal ulcer patients, vagotomy brings about marked changes in the undissociated acidity of gastric juice. The observed values were significantly lower after vagotomy than before the operation. At the same time, there was no parallel change in gastric protein concentration. Mean values of total and non-pepsin protein concentrations were higher after vagotomy. It seems logical that, in patients with duodenal ulcer, the parietal component of the
secretion should dominate, and thus the concentration of undissociated hydrogen ions should be lower before vagotomy than after it. It is, therefore, difficult to find a satisfactory explanation for the fact that the opposite phenomenon occurs.

No clear effect of pentagastrin dose was observed either on gastric protein or on the undissociated acidity of gastric juice. It is only the concentration of total and free acid, as well as pepsin, that exhibits definite response to pentagastrin infusion. Our results concerning the changes in total hydrogen ion concentration closely resemble those obtained by other authors. The values of $V_{max}$ and $D_{50}$ (pentagastrin dose response) found for our patients were similar to those obtained by Petersen and Myren (1975) for duodenal ulcer patients. After vagotomy the values of $V_{max}$ were significantly lower, and $D_{50}$ correspondingly higher than before the operation.
Makhloff et al. (1970) found a significant positive correlation between undissociated acidity of gastric juice and non-pepsin protein. These authors suggested that non-pepsin protein is the chief constituent of gastric juice taking part in the buffering of hydrogen ions. Our results are not consistent with this view. There was no significant correlation between undissociated acidity and non-pepsin protein of gastric juice in duodenal ulcer patients both before and after vagotomy in the course of pentagastrin stimulated and basal secretion.

There are many factors which may account for the differing results of the two studies. Makhloff et al. (1970) carried out repeated tests in one healthy subject, whereas we have studied a group of duodenal ulcer patients, performing two pentagastrin tests (one before and the other after vagotomy) in each patient. The end-point pH value of gastric juice titration was also different: 7-4 in our experiments and 8-3 in those of Makhloff et al. (1970). In addition, these authors used the difference between titratable acidity to pH 8-3 and pH 7-0 as an index of the true undissociated acidity. In fact, Makhloff et al. (1970) calculated the correlation between the values of this index and protein concentration.

The main argument of Makhloff et al. (1970) for using pH 8-3 as an end point for gastric juice titration was that only at such a pH is there a balance between the main cations of gastric juice and Cl\(^{-}\) ions. Evidently, it is only at so high a value that all the protons attached to the buffering groups are liberatered. In physiological conditions, however, it seems improbable that so high a pH value obtains at any time during the process of gastric secretion. Evidence is still lacking that the protons back-titrated to pH 8-3 are really 'parietal' protons buffered by gastric proteins or their degradation products in the course of the secretory process.

It is also the mode of stimulant administration that may influence the relation between gastric protein and undissociated acidity. The types of secretory tests were different in the both studies. Inasmuch as protein is secreted by at least two types of gastric mucosal cells (Makhloff et al., 1968), the different cells may not give the same response to different stimulating agents, and even to different modes of stimulant administration.

The results of our study seem to indicate that in patients with duodenal ulcer either not all non-pepsin protein takes part in buffering of hydrogen ions or non-protein buffers play a more important role. Unfortunately, the individual constituents of non-pepsin protein have not been well characterised. We also have no information on isoelectric points of gastric juice proteins, except for pepsin and intrinsic factor (Fruton, 1971; Marcoullis and Gråsbeck, 1975). Among non-protein buffers, organic acids are present in too low a concentration to be of any importance (Piper et al., 1967), but phosphate with its concentration in gastric juice ranging from 0-1 to 4-0 mmol/l (Burhol et al., 1966) may account for a significant part of buffering capacity of gastric juice constituents. It is also possible that the buffering capacity of the constituents of non-pepsin protein fraction may increase upon pepsin degradation. This view may be supported by the studies of Fordtran and Walsh (1973), who found that the buffer capacity of protein food increased by 33% after incubation with gastric juice. Whereas Roland et al. (1974) observed a lower concentration of pepsin in the gastric juice of vagotomised patients, our results showed no significant change in pepsin concentration after vagotomy. Although we have shown that undissociated acidity and non-pepsin protein are not correlated in patients with duodenal ulcer, we cannot definitely ascribe the buffering to any other specific component of gastric juice.

References


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T Popiela, Z Szafran, H Szafran and M Komorowska

Gut 1977 18: 208-213
doi: 10.1136/gut.18.3.208

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