Trophic effect of gastrin on the enterochromaffin like cells of the rat stomach: Establishment of a dose response relationship

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Abstract
Gastrin was given to rats by continuous subcutaneous infusion through implanted osmotic minipumps in doses covering a wide range of the dose response relationship for gastrin with regard to the trophic effect on the enterochromaffin like cells of the oxyntic mucosa. Thirty five rats were divided into five groups (each of seven rats), one group receiving a control solution of 1% albumin, the others receiving gastrin in 1% albumin at doses of 2·5, 5, 10, and 15 μg/kg/h, respectively. The plasma gastrin concentrations in the various groups increased in the same order of magnitude as expected from the gastrin doses given. Gastrin induced a dose dependent increase in enterochromaffin like cell density, oxyntic mucosal histamine concentration and histidine decarboxylase activity up to the dose of 5 μg/kg/h, where the increase levelled off. Hence, the dose response relationship for the trophic effect of gastrin on the enterochromaffin like cells seems to follow a polynomial rather than a linear function. These findings may also contribute to the understanding of the trophic effect of gastrin on enterochromaffin like cells in man with conditions associated with hypergastrinaemia.

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It is generally accepted that gastrin controls the growth of the enterochromaffin like cells of the oxyntic mucosa. Thus, profound inhibition of acid secretion by omeprazole or histamine 2 receptor blockers in the rat led to endogenous hypergastrinaemia and enterochromaffin like cell hyperplasia. The hypergastrinaemia and enterochromaffin like cell hyperplasia induced by these agents were abolished by antrectomy. Furthermore, hypergastrinaemia after antrectomy or partial gastric corpectomy and exogenous hypergastrinaemia induced by continuous infusion of gastrin evoked proliferation of the enterochromaffin like cells in the rat stomach. Therefore acid inhibition is not a prerequisite for the hyperplasia. It has been suggested that there is a linear correlation between plasma gastrin concentration and the enterochromaffin like cell density in the oxyntic mucosa of the rat. No report has fully established the relationship between the gastrin concentration and the trophic effect on the enterochromaffin like cell, however, as neither the minimal nor the maximal effective gastrin concentration with regard to the trophic effect on the enterochromaffin like cell has been assessed. From the study of Larsson et al, it could appear that the number of enterochromaffin like cells would increase infinitely with the gastrin concentration. In the rat, the enterochromaffin like cells produce and store most of the histamine in the oxyntic mucosa. From our laboratory, it has previously been reported that the acid stimulatory effect of gastrin can be fully explained by the stimulation of histamine release. Gastrin induced histamine release in a concentration-dependent fashion with threshold concentration of 2 pmol/l and reaching a maximum effect at a gastrin concentration of about 260–520 pmol/l. Furthermore, the histamine synthetising enzyme, histidine decarboxylase, and the histidine decarboxylase mRNA abundance are regulated by gastrin. The present study was done to establish a dose response relationship for the trophic effect of gastrin on the enterochromaffin like cells by giving exogenous gastrin in graded doses.

Methods

ANIMALS
The procedure described by Ryberg et al was followed. Thirty five female Sprague-Dawley rats each weighing approximately 200 g, were used. The animals were divided into five different groups, each group consisting of seven rats. One group was given a control solution (1% albumin) and the other four groups were given [Leu15]-gastrin-17 (L-G-17) in 1% albumin in the doses shown in the Table. Human L-G-17 and albumin were purchased from Sigma (St Louis, Mo, USA). L-G-17 or the control solution was administered subcutaneously through osmotic minipumps (ALZET 2 ML1, ALZA Corp, Palo Alto, Calif, USA) implanted on the back of the rats under general anaesthesia (0·2 ml 100 g/body weight of a solution containing fluanisone 2·5 mg/ml, phenytoin 0·05 mg/ml, and midazolam 1·25 mg/ml). The animals were treated for 28 days and the minipumps were changed every seventh day.

At day 28, the rats were bled by heart puncture under general anaesthesia (as described above). The stomach was removed, opened along the
greater curvature, rinsed in ice cold 0.9% saline and weighed. Subsequently the stomach was laid out on a glass plate and a tissue specimen, 4 mm in diameter, was taken by a punch press from the anterior wall on the greater curvature. The antrum was discarded and the oxyntic mucosa scraped off, weighed, diluted to 100 mg/ml in ice cold 0.01 M sodium phosphate buffer, pH 7.4, and homogenised. Aliquots of the homogenates were stored at −70°C for later determination of the histidine decarboxylase activity. One portion of the homogenates was boiled for 10 minutes, centrifuged at 5000 g for 10 minutes and the supernatant stored at −20°C for the later determination of histamine. The tissue specimens were fixed overnight in freshly prepared Stefanini’s solution with 1% (weight/volume) 1-ethyl-3(3-dimethylaminopropyl)-carbodiimide (Sigma), dehydrated in ethanol, and embedded in paraffin.

The study was approved by the animal welfare committee at the University Hospital of Trondheim.

PLASMA GASTRIN

Plasma gastrin concentration was determined in 200 μl plasma using a double antibody liquid phase 125I-Radioimmunoassay (Diagnostic Products Corporation, Los Angeles, Calif, USA). These antibodies have the same immunoreactivity towards L-G-17 and gastrin-17 (unpublished data). Results are expressed as pmol equivalents of synthetic human gastrin 17/1 plasma.

HISTAMINE IN OXYNTIC MUCOSA

The determination of histamine in the oxyntic mucosal homogenates was performed using a commercially available radioimmunoassay kit (Immunotech, Marseilles, France) with high specificity and sensitivity. Results are expressed as nmol histamine/g wet weight of oxyntic mucosa.

HISTIDINE DECARBOXYLASE ACTIVITY

The microprocedure described by Beaven et al with some modifications was used. We incubated 80 μl oxyntic mucosal homogenate with [1-14C]L-histidine (24 nCi, 0.48 nmol; New England Nuclear, Boston, Mass, USA), 5·10⁻⁴ M L-histidine, and 10⁻⁵ M pyridoxal 5-phosphate in a total reaction volume of 160 μl at 37°C for 60 minutes. The reaction was stopped by adding 80 μl 2 M perchloric acid followed by an incubation at 37°C for 30 minutes. The expelled 14CO2 was trapped in 50 μl Protosol (New England Nuclear) and counted after adding scintillation fluid. Results are expressed as pmol 14CO2/mg wet weight per hour.

MORPHOMETRIC ANALYSES

Sections of 5 μm thickness were cut perpendicularly to the mucosal surface. For the determination of enterochromaffin like cells, specimens were stained with histamine antibodies no. 8531 (Milab, Malmö, Sweden, kindly provided by Prof R Håkanson, University of Lund, Sweden) and visualised using the peroxidas reaction. In the rat this method can be regarded as specific for the enterochromaffin like cells, because the mast cells are few in the mucosa and restricted to the superficial layer. The enterochromaffin like cell density was determined in randomly selected fields. An eye piece ×10 with an ocular grid inserted and an objective ×40 were used. Epithelial cells located between the lines of the grid (0.25 mm mucosal length) in at least three visual fields in two different sections from each specimen were counted. Only enterochromaffin like cells with a visible nucleus were counted. The enterochromaffin like cell density is expressed as number of cells per mm mucosal length. The specimens were coded and the examiner was unaware of the group to which they belonged.

STATISTICAL ANALYSIS

A non-parametric one way analysis of variance, the Kruskal-Wallis H-test, was used for examination of a global difference between all groups. When significance was indicated, the two sample Mann-Whitney U-test was applied. All values are given as mean (SE). The level of statistical significance was set at 0.05.

Results

The results are summarised in the Table and Figure. The plasma gastrin concentrations increased from group to group as expected from the dose given. The rat weights did not differ between the groups. There was a significant increase in the weight of the whole stomach and oxyntic mucosa (p=0.030 and p=0.018, respectively). These differences were mostly caused by the increase in the weights in the treated groups compared with the control group.

The enterochromaffin like cell density increased from 66 (7) in the control group to 78 (12) (NS) in the group receiving 2.5 μg/kg/h and to 127 (12) (p=0.002) in the group receiving 5.0 μg/kg/h. The mucosal histamine content increased from 264·1 (20·1) nmol/g to 379·0 (New England Nuclear) and counted after adding scintillation fluid. Results are expressed as pmol 14CO2/mg wet weight per hour.
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Discussion

For the first time a wide range dose response relationship for the trophic effect of gastrin on the enterochromaffin like cells of the rat stomach has been established. The dose response curve for the trophic effect of gastrin is best described by a quadratic function with a maximal effect of gastrin at a plasma concentration in the range 250–400 pmol/l. Thus, the enterochromaffin like cell density, histamine content and histidine decarboxylase activity of the oxyntic mucosa all increased with increasing gastrin dose up to a maximal level at a gastrin concentration in the same range as found for the maximal histamine releasing and acid stimulatory effect of gastrin. Hence, the histamine releasing and trophic effect of gastrin on the enterochromaffin like cells show about the same concentration dependence with regard to the minimal concentration of gastrin causing maximal effect. This may suggest that the histamine releasing and the trophic effects of gastrin on the enterochromaffin like cells are mediated by interaction with the same receptor.

Ryberg et al using the same radioimmunoassay for determination of plasma gastrin, found higher concentrations for the corresponding gastrin doses than we did. The basal gastrin concentration, however, was also proportionately lower in the present study. Furthermore, Ryberg et al showed that continuous infusion of L-G-17 through osmotic minipumps gave somewhat lower plasma gastrin concentrations just before the pumps were changed. Therefore, the gastrin concentrations in the present study are probably measured at the lowest level. The linear relationship between plasma gastrin concentration and the enterochromaffin like cell density previously reported may only reflect a more prolonged endogenous hypergastrinaemia in those with the highest gastrin concentration two hours after administration of the drug. In the study by Ryberg et al gastrin was probably given in such a dose that the gastrin concentration just reached the maximal effective level. In another study by Ryberg et al the enterochromaffin like cell labelling index was shown to increase linearly with the plasma gastrin concentration. From their figure, however, it can be extrapolated that the labelling index tended to level off, at a gastrin concentration corresponding to the lowest concentration giving maximal effect in the present study. In the study by Mattsson et al describing development of enterochromaffin like carcinoids in rats after long term hypergastrinaemia caused by partial corpectomy, the plasma gastrin concentration might have been above the minimal concentration necessary to give maximal trophic effect (supramaximal concentration).

Extrapolation of data from rat to man is difficult, but it is tempting to speculate that the findings in the present study may have its parallel in man and explain some findings reported in patients with diseases accompanied by hypergastrinemia. Blair et al reported that circulating gastrin accounted for approximately 90% of acid secretion in response to eating. They found that on average, a gastrin concentration of 1000 pg/ml (=500 pmol/l) gave maximal gastric acid

![Graph](image)

Density of enterochromaffin like (ECL) cells (A), histamine content (B), and histidine decarboxylase activity (C) in the oxyntic mucosa of the rat after continuous, subcutaneous treatment with various doses of [Leu 8]-gastrin-17 for 28 days. All values are mean (1 SE).


