Inhibitory effect of a cholecystokinin antagonist on the proliferative response of the pancreas to pancreateobiliary diversion

P Watanapa, E F Efa, K Beardshall, J Calam, C E Sarraf, M R Alison, R C N Williamson

Abstract

Since pancreateobiliary diversion probably stimulates pancreatic growth by increasing cholecystokinin secretion, the effect of the cholecystokinin antagonist CR-1409 on this adaptive response was tested. Male Wistar rats (n=108) weighing 220–250 g were randomised to receive either pancreateobiliary diversion (n=60) or sham diversion (n=48) and thereafter to receive either saline injections or CR-1409 (10 mg/kg/day, subcutaneously). Rats were killed at four, seven, and 14 days postoperatively, when blood was obtained for cholecystokinin assay and the pancreas was assessed for proliferative activity by three techniques: nucleic acid and protein assay, bromodeoxyuridine labelling, and metaphase arrest after vincristine administration (1 mg/kg, intraperitoneally). Pancreateobiliary diversion increased plasma cholecystokinin concentrations by 91% at seven days and 137% at 14 days, irrespective of CR-1409 treatment. Total pancreatic RNA content was doubled by pancreateobiliary diversion at four days (2.15 v 1.07 mg/100 g body weight: p<0.001) and at seven days (3.43 v 1.76 mg/100 g: p<0.001), and trebled at 14 days (4.27 v 1.32 mg/100 g: p<0.001). Pancreateobiliary diversion increased bromodeoxyuridine labelling index from 1.1 to 3.7% at seven days and the cell birth rate from 0.09 to 0.06%. CR-1409 completely abolished this proliferative response and partly prevented the rise in RNA. The results confirm pancreatic hypertrophy and increased acinar cell proliferation after pancreateobiliary diversion. CR-1409 prevents this adaptive growth, probably by blocking cholecystokinin receptors. Bromodeoxyuridine labelling and the metaphase arrest technique may be used to assess pancreatic cell kinetics.

Methods

EXPERIMENTAL DESIGN

Male Wistar rats (n=108) weighing 220–250 g were housed in groups of five in animal quarters with a 12 hour day/night cycle. Standard pelleted rat food (Paterson and the Christopher Hill Group, Porton-diet PRD) and water were freely available. After 72 hours of acclimatisation, animals were randomised to receive either pancreateobiliary diversion (n=60) or sham pancreateobiliary diversion, comprising triple small bowel transection and resuture (n=48). Pancreateobiliary diversion involved transposition of 50 cm jejunum to lie between the pylorus and duodenal papilla, while in shamms the small bowel was divided immediately distal to the pylorus, at the duodenojejunal junction and again at the level of the mid-small bowel. Operations were carried out under light ether anaesthesia. A continuous 6/0 silk suture was used for intestinal anastomoses.

Immediately after the operation, half the animals in each group were further randomised to receive either CR-1409 (10 mg/kg/day) or saline (2.5 ml/kg) by daily subcutaneous injection. CR-1409 was dissolved in distilled water.
pancreas was excised, trimmed free of adherent fat and lymph nodes, and weighed. Small samples were removed for protein and nucleic acid content measurement, bromodeoxyuridine labelling index, and metaphase arrest counting after vincristine injection (see below).

**BROMODEOXYURIDINE LABELLING INDEX**

The thymidine analogue bromodeoxyuridine is readily and specifically incorporated into DNA during the S-phase of the cell cycle. One hour before sacrifice, rats were given an intraperitoneal injection of 5-bromo-2'-deoxyuridine (50 mg/kg body weight). A small sample of the excised pancreas was placed in Carnoy’s solution for two to four hours and transferred to 75% ethanol. Tissue was processed in chloroform, wax embedded, and 4 μm sections were cut. Sections were dewaxed and dehydrated in 100% alcohol. Endogenous peroxidase activity was removed by subsequent immersion in methanol/hydrogen peroxide (98%-4%/1-6%). Dissociation of histones was achieved by using 1 mol/l HCl, which also partially denatured DNA. Each section was treated with a drop of primary antibody (mouse anti-human bromodeoxyuridine, Dako, High Wycombe; 1:20 dilution in phosphate buffered saline). After overnight incubation at 4°C, the section was incubated with a peroxidase conjugated second antibody (anti-mouse IgG, 1:50 dilution with phosphate buffered saline) for one hour at room temperature. The colour reagent diaminobenzidine plus H2O2 was used for staining, and each section was counterstained with haematoxylin. The number of S-phase cells in each section was counted and expressed as a percentage of 3500 exocrine pancreatic acinar cells (=labelling index).

**METAPHASE ARREST INDEX**

The proliferative status of exocrine pancreatic cells was also assessed by counting the number of cells arrested in metaphase by the stathmokinetic agent vincristine. Rats were killed serially 30, 60, 90, and 120 minutes after an intraperitoneal injection of vincristine sulphate (1 mg/kg body weight). Pancreatic tissue was placed in fixative solution and was stained with haematoxylin and eosin. The number of arrested metaphase figures was counted per 2000 pancreatic acinar cells. The cell birth rate (Kb) was determined from the slope of the line that was fitted to the data by the method of least squares.

**PLASMA CHOLECYSTOKININ ASSAY**

Plasma cholecystokinin peptides were extracted from cardiac blood samples with C18 ‘SepPak’ cartridges (Waters, Harrow, UK) and eluates were dried by centrifugal evaporation (Savant, Farmingdale NY, USA). Cholecystokinin was measured by a specific radioimmunoassay based on antiserum A2, raised by immunising a rabbit with natural porcine cholecystokinin 33 (donated by Professors V Mutt and S R Bloom). Antiserum A2 (1:60 000) was incubated at 4°C for three days with standard cholecystokinin 8 or with plasma samples plus cholecystokinin 8
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Figure 3: Pancreatic nucleic acid and protein contents per 100 g body weight. (Values mean (SEM).) Sham = transection (control); PBD = pancreatobiliary diversion. Significance v corresponding sham group. *p<0.05, **p<0.002, ***p<0.001.

Results

MORTALITY AND BODY WEIGHT (Fig 1)
There were nine early deaths from anastomotic leakage (pancreatobiliary diversion 7, transection 2). There was no significant difference in body weight between the groups at four days. At seven days, rats with pancreatobiliary diversion plus CR-1409 weighed 17% less than controls (triple transection + saline). By 14 days, pancreatobiliary diversion rats weighed less than shams (by 16–25%) whether or not they received CR-1409. In sham rats CR-1409 did not affect body weight.

PANCREATIC WEIGHT (Fig 2)
Pancreatic growth occurred with time in every group. Differences in absolute pancreatic weight were generally minor and seemed to reflect differences in body weight. When this variable

Significance v corresponding sham group *p<0.05.

Plasma cholecystokinin concentration in rats with pancreatobiliary diversion (PBD) and sham PBD. (Values mean (SEM).)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Cholecystokinin concentration (pmol/l)</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>At 4 days</td>
</tr>
<tr>
<td>Sham + saline</td>
<td>4.88 (0.34)</td>
</tr>
<tr>
<td>PBD + saline</td>
<td>7.36 (1.91)</td>
</tr>
<tr>
<td>Sham + CR-1409</td>
<td>5.42 (1.15)</td>
</tr>
<tr>
<td>PBD + CR-1409</td>
<td>6.62 (0.87)</td>
</tr>
</tbody>
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tracer labelled with iodine-125 (1000 cpm, Amersham, UK) in 0.05 mol/l sodium phosphate buffer (pH 7.4) with 0.25% gelatin and 0.01 mol/l EDTA. Free and bound tracer were separated by the addition of 6% (weight/volume) charcoal (Norit PNS, BDH, Poole, UK) with 0.6% (weight/volume) dextran. The concentrations of pure peptides that produced half maximum inhibition of binding of tracer to A2 were 2.0 pmol/l for cholecystokinin 8, 2.4 pmol/l for cholecystokinin 33, and 2.2 pmol/l for gastrin 17. The coefficient of variation within assays was 8.2% and between assays was 12.8%. The sensitivity of the assay (defined as minimal amount of cholecystokinin 8 that could be distinguished from zero cholecystokinin with 95% confidence) was 0.2 pmol, and the recovery of cholecystokinin 8 and 33 through SepPak and assay procedure was 79%.

BIOCHEMICAL STUDIES
Pancreatic tissue was homogenised in phosphate buffered saline (136.9 mmol/l NaCl, 2.7 mmol/l KCl, 8.1 mmol/l Na2HPO4, 1.5 mmol/l KH2PO4, 0.9 mmol/l MgCl2, 0.5 mmol/l CaCl2, 6H2O, adjusted to pH 7.5 with NaOH), and the concentration of the homogenate was adjusted to 1 mg tissue per ml. Pancreatic contents of DNA and RNA were assayed by the fluorimetric method of Boer et al. but using an excitation wavelength of 536 nm and an emission wavelength of 602 nm. Protein content was determined by the method of Lowry et al. Since body weight has a profound effect on pancreatic weight, protein and nucleic acid contents were expressed as mg per 100 g body weight to correct for this variable, in line with other workers. The coefficient of variation within assays was 8.2% and between assays was 12.8%. The sensitivity of the assay (defined as minimal amount of cholecystokinin 8 that could be distinguished from zero cholecystokinin with 95% confidence) was 0.2 pmol, and the recovery of cholecystokinin 8 and 33 through SepPak and assay procedure was 79%.

Statistical analysis
Student's t test for unpaired data was used for the group analysis. Results were given as mean (SEM). The Pearson product moment correlation was used to assess correlations between the results of the bromodeoxyuridine labelling index and metaphase arrest studies.

Total DNA/100 g body weight

<table>
<thead>
<tr>
<th>Group</th>
<th>Days after operation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham + saline</td>
<td>0.64 (0.21)</td>
</tr>
<tr>
<td>PBD + saline</td>
<td>0.85 (0.27)*</td>
</tr>
<tr>
<td>Sham + CR-1409</td>
<td>0.92 (0.28)</td>
</tr>
<tr>
<td>PBD + CR-1409</td>
<td>0.89 (0.26)</td>
</tr>
</tbody>
</table>

Total RNA/100 g body weight

<table>
<thead>
<tr>
<th>Group</th>
<th>Days after operation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham + saline</td>
<td>4.44 (0.24)</td>
</tr>
<tr>
<td>PBD + saline</td>
<td>4.28 (0.26)*</td>
</tr>
<tr>
<td>Sham + CR-1409</td>
<td>4.27 (0.25)</td>
</tr>
<tr>
<td>PBD + CR-1409</td>
<td>4.33 (0.26)</td>
</tr>
</tbody>
</table>

Total protein/100 g body weight

<table>
<thead>
<tr>
<th>Group</th>
<th>Days after operation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham + saline</td>
<td>2.25 (0.21)</td>
</tr>
<tr>
<td>PBD + saline</td>
<td>2.36 (0.29)</td>
</tr>
<tr>
<td>Sham + CR-1409</td>
<td>2.36 (0.28)</td>
</tr>
<tr>
<td>PBD + CR-1409</td>
<td>2.38 (0.27)</td>
</tr>
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</table>
NUCLEIC ACID AND PROTEIN CONTENTS (Fig 3)
Pancreatic DNA content steadily increased after pancreatobiliary diversion. Initially, values were lower than in shams but by 14 days there was a (non-significant) 29-34% increase. Likewise, alterations in protein content were slight, with a trend towards greater values 14 days after pancreatobiliary diversion. By contrast, pancreatobiliary diversion increased RNA content substantially at every time point – by 101% at four days, 95% at seven days, and 224% at 14 days. These increments were partially inhibited by the hormone antagonist CR-1409, being 33% at seven days (p<0.05 v pancreatobiliary diversion + saline) and 139% at 14 days.

BROMODEOXYURIDINE LABELLING INDEX (Fig 4)
Pancreatobiliary diversion sharply increased the bromodeoxyuridine labelling index at seven days (3-74 v 1.1%). Thereafter, the proliferative response declined, and at 14 days the difference was much smaller (1.77 v 0.34%). CR-1409 abolished the effect of pancreatobiliary diversion, so that there was no significant difference from sham values at any time point. CR-1409 had no effect in rats with sham pancreatobiliary diversion.

METAPHASE ARREST INDEX (Fig 5)
Broadly similar results were obtained by the stathmokinetic technique. Four days after pancreatobiliary diversion, cell birth rate (K₀) was already higher than in shams (0.32 v 0.20%) and there was a dramatic difference at seven days (0.60 v 0.09%) but no residual effect at 14 days. Again, CR-1409 largely abolished the increased birth rate seven days after pancreatobiliary diversion (0.14% v 0.09% in shams) but had no consistent effect within the sham groups. There was a close correlation between the bromodeoxyuridine labelling and the cell birth rates with a Pearson correlation coefficient of 0.863 (Fig 6).

The potential doubling time of rat pancreatic acinar cells (Tpd) was calculated as the reciprocal of the cell birth rate (K₀), assuming a rectangular age distribution. Tpd in shams was 21 days (four days postoperatively) and 46 days (seven days postoperatively). Pancreatobiliary diversion shortened the doubling time considerably, reducing Tpd to 13 days (at four days) and 6-9 days (at seven days). The cholecystokinin antagonist abrogated this effect, producing a Tpd of 30 days (at seven days after pancreatobiliary diversion plus CR-1409).

Discussion
Our data support a tropic role for cholecystokinin in pancreatic growth after pancreatobiliary diversion, since the operation doubled circulatory concentrations of the hormone. There are two possible explanations for this hypercholecystokininemia. Diversion of pancreatobiliary secretions from the transposed jejunum should suppress the negative feedback mechanism on cholecystokinin secretion, and the jejunal hyperplasia that follows the procedure might well involve the enteroendocrine cells and lead
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Figure 5: Cell birth rates (Kb) per hour, derived from linear regression of metaphase indices. Pancreatic acinar cell doubling time (Tpd) = 1/Kb:
Thus, Tpd (PBD+saline, at 4 days) = 1/0.0032 = 312 hours = 13 days
Sham = transection (control); PBD = pancreaticobiliary diversion

Figure 6: Linear relation between labelling index and cell birth rates (Kb) per hour. Pearson correlation coefficient = 0.963.

We thank the Royal Postgraduate Medical School and the Hammersmith and Queen Charlotte's Special Health Authority for supporting this research, Rotta Research Laboratories (Milan, Italy) for kindly providing the CCK antagonist CR-1409 and W K Man, S K Li for nucleic acid and protein assay.

Assays of nucleic acid and protein content can give only an indirect assessment of the proliferative activity of exocrine pancreatic cells. Moreover, ductular, endocrine, and connective tissue elements will contribute to the RNA, DNA, and protein concentrations in the pancreas. These shortcomings limit the value of looking at indices of pancreatic mass to determine short term changes in exocrine pancreatic growth. We have therefore adapted two direct measurements of cell proliferation to the exocrine pancreas. Bromodeoxyuridine labelling has been widely used in the gastrointestinal mucosa, thyroid, and endocrine pancreas but not previously in the exocrine pancreas. Likewise, the metaphase arrest technique is well established for intestinal mucosa and has now been applied to the pancreas. Although baseline numbers of cycling cells are low within pancreatic acini, each technique showed a clearcut burst of proliferative activity seven days after pancreaticobiliary diversion with values returning to normal by 14 days. There was an excellent correlation between the bromodeoxyuridine labelling index and cell birth rate but not between these values and nucleic acid or protein content. These techniques are labour intensive, however; they require two to three thousand cells to be counted because of the low level of mitotic activity in the exocrine pancreas.

CR-1409 abolished the hyperplastic response of the pancreas to pancreaticobiliary diversion and partially inhibited the hypertrophic response without reducing the high circulating cholecystokinin concentrations. These data confirm that cholecystokinin is indeed involved in the tropic response. The intensity of pancreatic adaptation, however, did not completely accord with circulating cholecystokinin concentrations. Mitotic activity was maximal at seven days and then gradually subsided despite persistent hypercholecystokininaemia. Others have found a similar pattern, which may reflect down regulation of the acinar cell. Nevertheless, pancreaticobiliary diversion causes a sustained increase in pancreatic mass and, in our experience, it enhances azaserine induced carcinogenesis.

to increased cholecystokinin production. Our finding that the specific cholecystokinin antagonist CR-1409 largely inhibits the proliferation of the pancreatic acinar cells in response to pancreaticobiliary diversion underlines its importance as an intermediary. Since relative pancreatic weight increased after diversion despite the cholecystokinin antagonist, however, the hypertrophic response was only partly blocked. CR-1409 is recognised as one of the most potent antagonists of cholecystokinin and is known to prevent the effects of the hormone and its analogue on pancreatic growth.38
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