**STOMACH**

*Helicobacter pylori* infection modifies gastric and plasma ghrelin dynamics in Mongolian gerbils

H Suzuki, T Masaoka, T Hosoda, T Ota, Y Minegishi, S Nomura, K Kangawa, H Ishii

Background and aim: Although ghrelin, a novel growth hormone releasing peptide localised mainly in the gastric fundus, is reported not only to accelerate food passage and gastrointestinal motility but also to affect appetite and weight control, regulation of gastric ghrelin secretion under the conditions of gastric *Helicobacter pylori* infection is unknown. The present study was designed to investigate plasma and gastric ghrelin levels in Mongolian gerbils with *H pylori* colonisation of the gastric mucosa.

Methods: Gerbils orally inoculated with *H pylori* were examined after inoculation. To examine preproghrelin mRNA expression in the gastric mucosa, cDNA encoding the gerbil preproghrelin and glyceraldehyde-3-phosphate dehydrogenase homologue was isolated and a quantitative reverse transcription-polymerase chain reaction system was established.

Results: In gerbils showing *H pylori* colonisation (*H pylori* group), expression of preproghrelin mRNA and total ghrelin levels were significantly decreased 17 and 23 weeks later (p<0.01). Although the number of ghrelin immunoreactive cells decreased as the stomach weight increased, the gastric contents of total and active ghrelin in this group were the same as those in controls. Gastric myeloperoxidase activity showed a positive correlation with plasma ghrelin levels. On the other hand, at 17 weeks, plasma ghrelin levels were significantly increased in the *H pylori* group (p<0.05), suggesting a compensatory increase in secretion of the peptide at this time point.

Conclusion: The present experimental study demonstrated that gastric and plasma ghrelin dynamics are altered in response to *H pylori* infection.

METHODS

Cloning of gerbil ghrelin and GAPDH cDNA

Total RNA was obtained from the Mongolian gerbil (*Meriones unguiculatus*) stomach using the RNasy Mini-kit (Qiagen, Hilden, Germany). In order to clone a gerbil ghrelin homologue, degenerative primers were designed from human and mouse ghrelin cDNA (forward primer M01: 5'-CCGGAATTCTCAGCATGCTCTGGATGGACATGGCCATGGCAGG-3' and reverse primer M02: 5'-CCGGAATTCTCAGCGTGCCGCCTCTTGTGACCCTCTTCCCA-3'). Approximately 2 μg of total RNA were used as the template and one step reverse transcription-polymerase chain reaction (RT-PCR) was performed with the M01 and M02 primers (50°C for 30 minutes, 95°C for 15 minutes, and 35 cycles of 55°C for 30 seconds, 72°C for one minute, and 94°C for 20 seconds). The size of the PCR product was close to the expected size (332 bp); the product was subcloned into the pBluescript SK + vector (Stratagene, La Jolla, California, USA), and the DNA sequences were determined using the 310 Genetic Analyzer (PE Applied Biosystems, Norwalk, Connecticut, USA).

Abbreviations: GAPDH, glyceraldehyde-3-phosphate dehydrogenase; RT-PCR, reverse transcription-polymerase chain reaction; RIA, radioimmunoassay; DAB, 3,3’-diaminobenzidine tetrahydrochloride; MPO, myeloperoxidase

To obtain the full length cDNA of ghrelin, we performed 5′-RACE and 3′-RACE with primers designed based on the sequences obtained. For 5′-RACE, two gene specific antisense primers, M06 (5′-TCAAGCTGCGGCGCTTTGACGCT-3′) and M07 (5′-GCTCGTACAGCTTTGACGCTAAGGTC-3′), were designed, and 5′-RACE was performed using the 5′ RACE system for rapid amplification of the cDNA ends (Invitrogen Corp., Carlsbad, California, USA). For 3′-RACE, a gene specific sense primer, M05 (5′-TGCGGATCCGTACAGCTTTGACGCTAAGGTC-3′) and reverse primer M11 (5′-CGGAAATATACCTCTGAGGCGGCTTCTT-3′) were used. The sizes of the PCR products were close to the expected size (990 bp). For 5′-RACE, a gene specific antisense primer, M17 (5′-CTCTGCTGACGCTAAGGTC-3′), was designed, and for 3′-RACE, a gene specific sense primer, M18 (5′-TCTTCTCCACCATGGAAGGGCGGTAAG-3′), was used. Each amplified fragment was directly sequenced using the 310 Genetic Analyzer (PE Applied Biosystems).

For cloning of the gerbil homologue of GAPDH, the same method as that described above for ghrelin was used. In brief, to generate a fragment of GAPDH, forward primer M08 (5′-CGGGGATCCGTATGGGGCGCTTTGACGCG-3′) and reverse primer M11 (5′-CGGAAATATACCTCTGAGGCGGCTTCTT-3′) were used. The sizes of the PCR products were close to the expected size (990 bp). For 5′-RACE, a gene specific antisense primer, M17 (5′-CTCTGCTGACGCTAAGGTC-3′), was designed, and for 3′-RACE, a gene specific sense primer, M18 (5′-TCTTCTCCACCATGGAAGGGCGGTAAG-3′), was used. Each amplified fragment was directly sequenced using the 310 Genetic Analyzer (PE Applied Biosystems).

Quantitative RT-PCR
Total RNA was extracted from the stomach of the Mongolian gerbils using the RNeasy Mini Kit (Qiagen). A TaqMan quantitative real time RT-PCR was performed to detect preproghrelin mRNA and GAPDH mRNA with the ABI PRISM 7700 sequence detection system (PE Applied Biosystems). The following primers were used to amplify preproghrelin mRNA: ghrelin-F (5′-GGA ATC CAA GAA GCC ACC AGC-3′), ghrelin-R (5′-GCT CCT GAC AGC TTG ATG CCA-3′), and ghrelin-Taq (5′-FAM-AAC TGC AGC CAG CAC GTC TGG AAG GC-GAMRAA-3′); to amplify GAPDH mRNA as the internal control, the following primers were used: GAPDH-F (5′-TTC AAC GGC ACA GTC AAG GC-3′), GAPDH-R (5′-GCC TTC TCC ATG GTG GTG AAG-3′), and GAPDH-Taq (5′-FAM-CCC ATC ACC TTC CGC GAG CGA TAMRA-3′).

The PCR fragments of the Mongolian gerbil preproghrelin and GAPDH were amplified and cloned into the pDrive cloning vector (Qiagen) and respectively used as standards. Preproghrelin mRNA expression levels were normalised using GAPDH mRNA expression levels.

Radioimmunoassay (RIA) for ghrelin

Stomach, duodenum, or jejunum sampling
Fresh whole anterior wall specimens of the glandular stomach, duodenum (5–15 mm from the pyloric ring), or jejunum (120–130 mm from the pyloric ring) were frozen immediately after collection and stored at −80°C. Each sample was boiled for five minutes in a 10-fold volume of water to inactivate the intrinsic proteases. The solution was adjusted to 1 M acetic acid after cooling, and the tissue was homogenised since acidification of sample solution prevented the loss of extraction recovery. The supernatant was lyophilised and then subjected to ghrelin RIA. The extraction efficiency of tissue ghrelin was more than 95%.

Plasma sampling
Whole blood samples (1 ml) were obtained from the right ventricle in tubes containing EDTA-2Na (1 mg/ml) and aprotinin (500 kIU/ml). Each plasma sample (500 μl) was treated with a C18 Sep-Pak cartridge (Waters, Milford, Massachusetts, USA) for peptide extraction. The cartridge was washed and eluted; the eluate (50 μl equivalent of plasma) was lyophilised and subjected to ghrelin RIA. The extraction efficiency of plasma ghrelin was approximately 90%.

RIA for ghrelin
The two RIA techniques were used for measuring ghrelin, as described previously. Briefly, ghrelin levels were measured using two polyclonal rabbit antibodies raised against the N terminal (1–11) (Gly1-Lys11) or C terminal (13–28) (Gln13-Arg28) fragment of rat ghrelin. Two tracer ligands were synthesised: [Tyr29]-rat ghrelin for antirat ghrelin (1–11) antiserum and [Tyr12]-rat ghrelin (13–28) for antirat ghrelin (13–28) antiserum. RIA incubation mixtures, containing 100 μl of either standard ghrelin or unknown sample with 200 μl of antiserum diluted in RIA buffer containing 0.5% normal rabbit serum, were initially incubated for 12 hours. Then, 100 μl of 125I labelled tracer (15 000 cpm) were added and the mixture was incubated for 36 hours. Antirabbit IgG goat serum (100 μl) was added prior to an additional 24 hour incubation period. Free and bound tracers were then separated by centrifugation at 3000 rpm for 30 minutes. Following aspiration of the supernatant, radioactivity in the pellet was quantitated using a gamma counter (ARC-600; Aloka, Tokyo, Japan). All assays were performed at 4°C. The antirat ghrelin (1–11) antiserum specifically recognised the n-octanoylated form of rat ghrelin but not the des-acyl form. The antirat ghrelin (13–28) antiserum recognised both the acylated and des-acyl forms of rat ghrelin equally. Both antisera were equally cross reactive with human and gerbil ghrelin and did not recognise the other enteric peptides. The respective intra- and interassay coefficients of variation for the N terminal RIA were 3% and 6%, and for the C terminal RIA, 6% and 9%.

Mongolian gerbil model of H pylori infection
All experiments and procedures carried out on animals were approved by the Keio University Animal Research Committee (No 023009). Thirty one specific pathogen free male Mongolian gerbils (MGS/Sea, five week old; Seac Yoshitomi, Fukuoka, Japan) were administered H pylori suspensions (ATCC43504: 106 colony forming units (CFU)/ml, 15 ml/kg) while 27 control gerbils were administered buffer solution alone after overnight deprivation of food. All animals were allowed free access to water and a standard pellet diet (CE-2; Clea Japan, Tokyo, Japan). Four, 17, and 23 weeks after inoculation, the gerbils were examined under ether anaesthesia after 16 hours of food deprivation, and sacrificed by an overdose of ether.

H pylori infection at each time point was examined by determining the number of CFU in a microaerobic bacterial culture. Briefly, the diluted homogenates of the stomachs were plated onto Brucella agar plates containing 10% horse blood, 2.5 μg/ml amphotericin B, 9 μg/ml vancomycin, 0.32 μg/ml polymyxin B, 5 μg/ml trimethoprim, and 50 μg/ml 2, 3, 5-triphenyl-tetrazolium chloride. The plates were then incubated at 37°C in a microaerobic atmosphere for seven days. The number of colonies was counted, and the amount of viable H pylori was expressed as the number of CFU/g of tissue.

Immunohistochemistry
Stomach tissue specimens of the gerbils were fixed in 10% neutralised formalin and embedded in paraffin. Paraffin sections were placed on slides pretreated with a 0.01% aqueous solution of poly-L-lysine. Deparaffinisation and hydration were conducted. Then, the antigens were retrieved by heating for 15 minutes at 121°C in citrate buffer (10 mM,
pH 6.0). After cooling, endogenous peroxidase was quenched by 0.3% hydrogen peroxide. After washing, non-specific binding was blocked by a blocking reagent (BlockAce; Dainippon Pharm, Osaka, Japan). All sections were incubated overnight at 4°C with antighrelin (13–28) antiserum (1:10 000). After washing with TBS-T, slides were incubated with peroxidase labelled dextran polymer conjugated goat antimouse IgG in Tris-HCl (EnVision/HRP; Dako Japan, Kyoto, Japan) for 30 minutes at room temperature and then visualised after colour development using 3,3′-diaminobenzidine tetrahydrochloride (DAB) solution for three minutes. Countereasting was performed with haematoxylin.

The stained sections were observed under a light microscope equipped with a 3CCD digital camera (C7700; Hamamatsu Photonics, Hamamatsu, Japan).21 and each image was stored as a PSD file (Adobe Photoshop 7.0). Haematoxylin and/or DAB stained nuclei were counted using a particle analysis program (Ultimage Pro. 2.6.4; Alliance Vision, France). The density of ghrelin immunoreactive cells (D:ghrelin) was computed using the following equation:

\[
D_{\text{ghrelin}} = (\frac{N_{\text{g}}}{N_{\text{t}}}) \times 100(\%)
\]

where \(N_g\) and \(N_t\) represent the number of ghrelin immunoreactive cells and the total cell number, respectively, in the region of interest.

To investigate colocalisation of parietal cells with ghrelin immunoreactive cells, some specimens were also stained for \(\text{H}^+\text{, K}^+\text{-ATPase}\). After ghrelin immunostaining with DAB, specimens were incubated with a mouse anti-\(\text{H}^+\text{, K}^+\text{-ATPase}\) subunit monoclonal antibody (RDI, Flanders, New Jersey, USA; 1:300) overnight at 4°C. The slides were then incubated with goat antimouse IgG conjugated to alkaline phosphatase labelled dextran polymer in Tris-HCl buffer (EnVision/AP) for 30 minutes at room temperature. Sections were then visualised after exposure to the Dako Fuchsin substrate chromogen system for seven minutes.

**MPO activity**

Tissue samples of gastric mucosa were collected in tubes containing phosphate buffered saline with protease inhibitors (100 μM phenylmethylsulphonyl fluoride, 10 μg/ml aprotinin) and sonicated over ice in 30 consecutive 0.5 second bursts at 0.5 second intervals at a power setting of 150 W (VCX 750; Sonics and Materials, Inc., Newton, Connecticut, USA). Total protein in the homogenates was measured using the modified Lowry method,22 as described by Smith and colleagues.23

Myeloperoxidase (MPO) activity, an index of polymorphonuclear cell accumulation, was determined using a previously described method, with some modifications.24 Samples of mucosal homogenates were centrifuged at 8000 g for 15 minutes at 4°C to pellet the insoluble cellular debris. The pellet was then rehomogenised in an equal volume of 0.05 M potassium phosphate buffer (pH 5.4) containing 0.5% hexadecyltrimethylammonium bromide. Samples were centrifuged at 8000 g for 15 minutes at 4°C and the supernatants were saved. MPO activity was assessed by measuring \(\text{H}_2\text{O}_2\) dependent oxidation of 3,3′,5,5′-tetramethylbenzidine. One unit of enzyme activity was defined as the amount of MPO that caused a change in absorbance of 1.0/minute at 655 nm at 25°C.

**Statistical analysis**

All data are expressed as mean (SEM); \(p<0.05\) was considered to denote statistical significance. The data were analysed using one way analysis of variance followed by Scheffe’s multiple comparison test.

**RESULTS**

**cDNA and protein sequences of the Mongolian gerbil preproghrelin and characterisation of the peptide**

The isolated full length Mongolian gerbil preproghrelin cDNA was 460 bp long, consisting of 17 bp in the 5'-untranslated region, 354 bp of coding region, and 89 bp in the 3'-untranslated region. The predicted initiation methionine was located at nucleotide position 18–20. An AATAAA polyadenylation signal was identified in the 3' non-coding region (position 452–457) (fig 1A) (Suzuki et al, EMBL/GenBank: AF442491). The deduced amino acid sequence of the coding region indicated that gerbil preproghrelin was composed of 117 amino acid residues, the same length as that of the mouse, rat, and human preproghrelin (fig 1B) (AF442491). At the nucleotide level, the gerbil preproghrelin sequence exhibited an average homology score of 85% with human (AJ252278), 90% with mouse (AJ243503), and 92% with rat (AB029433) preproghrelin (fig 1B). The gerbil preproghrelin was highly conserved, the protein sharing 83% (97/117) identity with its human counterpart, 89% (104/117) identity with its mouse counterpart, and 88% (103/117) identity with its rat counterpart. The unidentified third residue of the purified peptide was determined to be serine, the same as that in the case of other mammalian ghrelin.

As gerbil ghrelin was determined to be entirely homologous with rat ghrelin apart from one amino acid (Thr12), two polyclonal antibodies against rat ghrelin were used to measure the gerbil ghrelin in the present study.

**GAPDH and characterisation of this peptide**

The isolated full length Mongolian gerbil GAPDH cDNA was 1265 bp long, consisting of 71 bp in the 5'-untranslated region, 1002 bp in the coding region, and 192 bp in the 3'-untranslated region (Suzuki et al, EMBL/GenBank: AY066007). The predicted initiation methionine was located at nucleotide position 72–74. The deduced amino acid sequence of the coding region indicated that the gerbil GAPDH may be composed of 333 amino acids residues, the same length as that of the mouse or rat GAPDH, and two amino acids shorter than the human GAPDH. The deduced amino acid sequence of gerbil GAPDH was 97.6% (325/333) identical to that of its mouse counterpart, 97.3% (324/333) identical to that of its rat counterpart, and 95.2% (319/333) identical compared with its human counterpart.

**Preproghrelin mRNA expression and ghrelin contents in the stomach**

The level of preproghrelin mRNA as a ratio of the level of GAPDH mRNA (preproghrelin/GAPDH mRNA) in the stomach was determined using the modified Lowry method,22 as described by Smith and colleagues.23

Myeloperoxidase (MPO) activity, an index of polymorphonuclear cell accumulation, was determined using a previously described method, with some modifications.24 Samples of mucosal homogenates were centrifuged at 8000 g for 15 minutes at 4°C to pellet the insoluble cellular debris. The pellet was then rehomogenised in an equal volume of 0.05 M potassium phosphate buffer (pH 5.4) containing 0.5% hexadecyltrimethylammonium bromide. Samples were centrifuged at 8000 g for 15 minutes at 4°C and the supernatants were saved. MPO activity was assessed by measuring \(\text{H}_2\text{O}_2\) dependent oxidation of 3,3′,5,5′-tetramethylbenzidine. One unit of enzyme activity was defined as the amount of MPO that caused a change in absorbance of 1.0/minute at 655 nm at 25°C.

Statistical analysis

All data are expressed as mean (SEM); \(p<0.05\) was considered to denote statistical significance. The data were analysed using one way analysis of variance followed by Scheffe’s multiple comparison test.
Amino acids constituting the matured ghrelin peptides are printed in bold type. The EMBL/GenBank accession number for this sequence is AF442491. (B) Alignment numbered on the left and right, respectively. The amino acid residues in the signal peptide sequence are printed in italics. The amino acid residues corresponding to the ghrelin peptide are shown as the shaded area.

Figure 1 cDNA and protein sequences of preproghrelin and their homology with those of other species. (A) Nucleotides and amino acids are numbered on the left and right, respectively. The amino acid residues in the signal peptide sequence are printed in italics. The amino acid residues corresponding to the ghrelin peptide are shown as the shaded area.

Decreased in association with accumulation of inflammatory cells, ghrelin immunoreactive cells were still found to be localised in association with these H⁺-K⁺-ATPase positive cells (fig 4D). On the other hand, in H pylori colonised corpus mucosa showing severe atrophy, extensive disappearance of H⁺-K⁺-ATPase positive cells was noted and the number of ghrelin immunoreactive cells was also decreased (fig 4E).

Representative photomicrographs of ghrelin immunohistochemistry in the corpus revealed a decrease in the number of brown stained ghrelin immunoreactive cells in H pylori colonised mucosa (fig 5B, D).

Morphometric analysis also revealed that the number of ghrelin immunoreactive cells/mm² was significantly decreased in H pylori colonised mucosa of gastric corpus, both at 17 and 23 weeks after H pylori inoculation (table 1). In contrast, the total cell number was found to be higher in H pylori colonised mucosa (table 1), possibly due to inflammatory cell infiltration. Taking these parameters into consideration, it was found that the density of ghrelin immunoreactive cells (%) was decreased in H pylori infected gastric corpus mucosa (table 1), suggesting a possible relationship between the increase in the number of inflammatory cells and decrease in the number of ghrelin immunoreactive cells.

Gastric mucosal inflammation

During the observation period, persistent H pylori colonisation was confirmed in all H pylori inoculated gerbils. No significant difference in body weight was observed between
the control and \textit{H pylori} groups (table 2), suggesting that although \textit{H pylori} altered the ghrelin dynamics, it probably did not influence the eating behaviour or body mass of the animals in such a short period of time after the onset of infection. In this situation, some compensatory mechanism may be evoked to regulate ghrelin independent growth hormone secretion and body mass. In contrast, stomach wet weight was significantly increased in the \textit{H pylori} group (table 2), which could reflect the macroscopic findings of severe thickening and oedematous changes associated with severe inflammation observed in \textit{H pylori} colonised gastric mucosa. Consistent with the increase in stomach weight, gastric mucosal MPO activity, which reflects polymorphonuclear infiltration, was increased significantly compared with that in controls 17 and 23 weeks after \textit{H pylori} inoculation (table 2). At four weeks after inoculation, although the wet weight of the stomach was significantly increased in the \textit{H pylori} group (table 2), no significant increase in MPO activity was observed, reflecting the merely oedematous changes of the gastric mucosa in the acute phase of infection.

Notwithstanding the above data on stomach weight, total and active ghrelin levels in each whole stomach specimen were determined to be the same in the control and \textit{H pylori} groups (table 3).

\textbf{Plasma ghrelin levels} Although the fasting plasma total ghrelin level was significantly increased in the \textit{H pylori} group compared with that of controls at 17 weeks after \textit{H pylori} inoculation, the difference was not significant at four or 23 weeks (fig 6A). In contrast, fasting plasma active ghrelin levels were significantly increased in the \textit{H pylori} group at both 17 and 23 weeks after \textit{H pylori} inoculation compared with those in each control group, although no significant increase in plasma active ghrelin level was shown at four weeks (fig 6B). Gastric mucosal MPO activity was correlated with plasma levels of total and active ghrelin (fig 7).

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\textbf{Figure 3} Levels of ghrelin in the stomach of Mongolian gerbils. (A) Total ghrelin levels in the stomach of Mongolian gerbils with or without \textit{Helicobacter pylori} colonisation at 17 and 23 weeks after bacterial inoculation. \textit{**}p<0.01 compared with controls at each of the time points. (B) Active ghrelin levels in the stomach of Mongolian gerbils with or without \textit{H pylori} colonisation at 17 and 23 weeks after bacterial inoculation. \textit{p}<0.05 compared with controls at 17 weeks; \textit{†}p<0.01 compared with controls at 23 weeks.

the control and \textit{H pylori} groups (table 2), suggesting that although \textit{H pylori} altered the ghrelin dynamics, it probably did not influence the eating behaviour or body mass of the animals in such a short period of time after the onset of infection. In this situation, some compensatory mechanism may be evoked to regulate ghrelin independent growth hormone secretion and body mass. In contrast, stomach wet weight was significantly increased in the \textit{H pylori} group (table 2), which could reflect the macroscopic findings of severe thickening and oedematous changes associated with severe inflammation observed in \textit{H pylori} colonised gastric mucosa. Consistent with the increase in stomach weight, gastric mucosal MPO activity, which reflects polymorphonuclear infiltration, was increased significantly compared with that in controls 17 and 23 weeks after \textit{H pylori} inoculation (table 2). At four weeks after inoculation, although the wet weight of the stomach was significantly increased in the \textit{H pylori} group (table 2), no significant increase in MPO activity was observed, reflecting the merely oedematous changes of the gastric mucosa in the acute phase of infection.

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\textbf{Figure 4} Colocalisation of ghrelin immunoreactive cells with parietal cells in the gastric fundic mucosa of Mongolian gerbils. The brown coloured cells represent ghrelin immunoreactive cells and the pink coloured cells represent \(H^+K^+\)-ATPase positive parietal cells. (A) Control fundic mucosa; magnification 20\(\times\). (B) Control fundic mucosa; magnification 100\(\times\), longitudinal view. (C) Control fundic mucosa; magnification 100\(\times\), cross sectional view. (D) \textit{Helicobacter pylori} colonised fundic mucosa showing inflammatory cell infiltration; magnification 20\(\times\). (E) \textit{H pylori} colonised fundic mucosa showing severe gastric mucosal atrophy; magnification 20\(\times\).
Ghrelin levels in the duodenum and jejunum

Figure 8 demonstrates levels of total ghrelin in the stomach, duodenum, and jejunum in both the control and H pylori groups at four weeks after H pylori inoculation. Although no significant difference in the content of ghrelin in any part of the intestine was observed between the control and H pylori groups at this time point, levels were significantly reduced to 0.55% and 0.16% in the duodenum and jejunum, respectively, compared with stomach levels, suggesting predominant localisation of ghrelin in the stomach of gerbils.

DISCUSSION

Modified ghrelin dynamics in H pylori infection were demonstrated in the present study using an experimental animal model, and reduced preproghrelin mRNA expression and ghrelin levels were clearly shown in the stomach of gerbils with H pylori colonisation, in association with a significant increase in inflammatory cell infiltration.

The amino acid sequence corresponding to the ghrelin peptide in the gerbil was 93% identical to that of humans and 96% identical to that of its mouse or rat counterpart (fig 1B).

Table 1  Ghrelin immunoreactive cells in the fundic mucosa of Mongolian gerbils

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<th>17 weeks</th>
<th>23 weeks</th>
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<tr>
<td></td>
<td>Control (n = 9)</td>
<td>H pylori (n = 6)</td>
</tr>
<tr>
<td>Ghrelin immunoreactive cells (No/0.1 mm²)</td>
<td>16.1 (1.0)</td>
<td>10.1 (1.0)*</td>
</tr>
<tr>
<td>Total cell number (No/0.1 mm²)</td>
<td>677.2 (21.4)</td>
<td>792.3 (15.0)</td>
</tr>
<tr>
<td>Ghrelin immunoreactive cells (%)</td>
<td>2.4 (0.1)</td>
<td>1.3 (0.1)**</td>
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* p < 0.05, ** p < 0.01 compared with controls at 17 weeks; *** p < 0.01 compared with control at 23 weeks.

Table 2  Body weight, stomach weight, and gastric myeloperoxidase (MPO) activity of Mongolian gerbils

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<th>4 weeks</th>
<th>17 weeks</th>
<th>23 weeks</th>
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<tr>
<td></td>
<td>Control (n = 10)</td>
<td>H pylori (n = 10)</td>
<td>Control (n = 9)</td>
</tr>
<tr>
<td>Body weight (g)</td>
<td>58.3 (0.7)</td>
<td>55.5 (0.8)</td>
<td>76.9 (2.2)</td>
</tr>
<tr>
<td>Wet weight of stomach (mg)</td>
<td>632.1 (14.1)</td>
<td>765.2 (53.9)**</td>
<td>619.1 (16.2)</td>
</tr>
<tr>
<td>Gastric MPO activity (mU/mg protein)</td>
<td>0.9 (0.2)</td>
<td>0.2 (0.1)</td>
<td>0.6 (0.1)</td>
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* p < 0.05 compared with controls at four weeks; ** p < 0.001 compared with control at 17 weeks; *** p < 0.001 compared with control at 23 weeks.
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suggesting functional homogeneity of this peptide in various animals. Differences in the amino acid sequences among the four species examined were found only at position 34 or 35 (fig 18), a rather small segment of the molecule, probably insignificant in relation to biological function. Indeed, studies on the structure-activity relationship of the peptide showed that the octanoyl group on Ser is the essential moiety for biological activity of this peptide,1 and that the N-terminal fragments showing conservation of the first five amino acids exhibited full functional activity.25

Using the same experimental animal, we previously reported a significant increase in the extent of inflammatory cell infiltration and oxidative stress in *H pylori* colonised gastric mucosa,15 which validated the usefulness of this model for the study of *H pylori* induced following infection.26 Alike cells induced by inflammatory stimuli, such as cytokines26 or free radicals,15 as plasma ghrelin elevation producing A-like cells may play a role in the enhanced ghrelin release noted following food deprivation.

These contentions are endorsed by the significant correlation between gastric MPO activity (table 1) and plasma ghrelin levels (fig 7). Even though the underlying mechanisms have still to be elucidated in detail, there may be a compensatory increase in levels of plasma ghrelin following food deprivation in *H pylori* colonised gerbils, in association with a decrease in gastric ghrelin density. Such elevation of plasma ghrelin levels may attenuate the reduction in appetite and

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<th>Table 3</th>
<th>Ghrelin contents (nmol) per one stomach in Mongolian gerbils</th>
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<tr>
<td></td>
<td>17 weeks</td>
</tr>
<tr>
<td>Total ghrelin contents/stomach (nmol)</td>
<td>2.91 (0.38)</td>
</tr>
<tr>
<td>Active ghrelin contents/stomach (nmol)</td>
<td>1.10 (0.46)</td>
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![Figure 6](image-url)

**Figure 6** Plasma levels of ghrelin in Mongolian gerbils. (A) Total plasma ghrelin levels in Mongolian gerbils with or without *Helicobacter pylori* colonisation at 17 and 23 weeks after *H pylori* inoculation. *p*<0.05 compared with controls at 17 weeks. (B) Plasma active ghrelin levels in Mongolian gerbils with or without *H pylori* colonisation at 17 and 23 weeks after *H pylori* inoculation. *p*<0.05 compared with controls at the time points indicated.

![Figure 7](image-url)

**Figure 7** Relationship between gastric myeloperoxidase (MPO) activity and plasma ghrelin level. (A) Gastric mucosal MPO activity was well correlated with total plasma levels of ghrelin. (B) Gastric mucosal MPO activity was well correlated with plasma levels of active ghrelin.

![Figure 8](image-url)

**Figure 8** Ghrelin contents in the glandular stomach, duodenum, and jejunum, four weeks after *Helicobacter pylori* inoculation (fmol/mg).
body weight in H pylori infected gerbils until 23 weeks (table 2). As longer term H pylori infection (72 weeks) than in the cohorts examined in the present study (23 weeks) reportedly evoked significant weight loss in gerbils,29 further extension of gastric inflammation and atrophy may nullify the effects of the compensatory increase in plasma ghrelin levels and lead to reduction of both appetite and body weight.

As the average intestinal wet weight in uninfected gerbils at four weeks after inoculation was 1540 mg, the total ghrelin content in the small intestine was computed to be less than 0.029 nmol (fig 8). In contrast, the average wet weight of the stomach in uninfected gerbils at four weeks was 632.1 mg (table 2) and the average gastric total ghrelin content was computed to be 2.168 nmol. The gastric total ghrelin content was estimated to be at least 74-fold greater than that of the small intestine. Therefore, it would be difficult to consider an extragastric source of ghrelin as the origin of the increased plasma ghrelin in H pylori colonised gerbils.

The present data in gerbils were inconsistent with the results of two recent clinical studies,30,31 the results of which were also discrepant. These discrepancies might be attributable to differences in the methods of measurement of ghrelin peptide or patient selection, or interspecies difference in the distribution of extragastric ghrelin.

In the present study, plasma total ghrelin levels were not significantly increased in H pylori colonised cohorts at 23 weeks after H pylori inoculation (fig 6). Among the H pylori colonised cohorts at 23 weeks, four of 11 gerbils showed plasma total ghrelin levels below the average value observed in controls at the corresponding time point. Among these four gerbils with H pylori infection, two showed severe gastric atrophy, with the number of ghrelin immunoreactive cells being less than 1.2%, as well as lower than average values for gastric ghrelin and preproghrelin mRNA. Severe atrophy might account for the absence of a statistically significant difference between infected and uninfected gerbils at 23 weeks. These results suggest that the ghrelin content in the stomach decreases in response to H pylori infection, and with further extension of gastric atrophy, plasma ghrelin levels also decrease. Although there does seem to be differential regulation of preproghrelin mRNA, mucosal ghrelin, and possibly secretion of active ghrelin, the molecular physiology underlying ghrelin dynamics is outside the purview of the current study. Further investigation of ghrelin dynamics based on long term H pylori infection could help to clarify these issues.

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