Modulation of host antimicrobial peptide (β-defensins 1 and 2) expression during gastritis

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Background: β-Defensins are a newly identified family of antimicrobial peptides that are expressed by epithelia on mucosal surfaces where their production is augmented by infection or inflammation. *Helicobacter pylori* colonises the gastric epithelium causing persistent gastric inflammation leading to antral and corpus gastritis, and peptic ulcer disease.

Aims: To evaluate the role of β-defensins in the innate immune response of the gastric epithelium to infection and inflammation, we have assessed mRNA expression and regulation of human β-defensins 1 and 2 (hBD1, hBD2) by *H pylori* and proinflammatory stimuli. We have also compared gene and peptide expression of these bactericidal agents in *H pylori* infected gastritis with that in normal gastric mucosa.

Methods: Modulation of expression of hBD1 and hBD2 by various stimuli was studied in three (AGS, MKN7, MKN45) gastric epithelial cell lines by quantitative competitive reverse transcription-polymerase chain reaction (RT-PCR). Defensin mRNA expression was measured by semiquantitative RT-PCR in gastritis tissue and compared with controls. Peptide localisation was assessed by immunohistochemistry.

Results: Cytotoxic *H pylori* and interleukin 1β (IL-1β) markedly upregulated expression of hBD2 in a dose and time dependent manner in both AGS and MKN7 cell lines. A modest increase in hBD1 expression was also noted during infection. Interestingly, induction of hBD1 gene expression by IL-1β was only observed in MKN7 cells. The magnitude of this response was delayed and reduced compared with hBD2 expression. In gastric biopsies, hBD2 was undetectable in normal gastric antrum but a marked increase was observed in *H pylori* positive gastritis compared with control tissue (p<0.001). Constitutive expression of hBD1 was observed in normal gastric mucosa and there was a significant increase in gastritis (p<0.05). Immunohistochemistry revealed a parallel increase in hBD1 and hBD2 peptide expression in gastritis tissue with positive staining confined to the surface epithelium of the gastric glands.

Conclusions: Modulation of β-defensin expression by pathogenic and/or inflammatory stimuli and their cellular localisation places these antimicrobial peptides in the front line of innate host defence in the human stomach.
increase in hBD1 and hBD2 mRNA and peptide expression in gastritis compared with control tissue. Our present study suggests that surface expression of hBD1 and hBD2 may form a critical component of the host innate defence to potential noxious stimuli in the human stomach.

MATERIALS AND METHODS

Cells and reagents

The human gastric cell lines AGS, MKN7, and MKN45 were maintained in RPMI 1640 culture medium supplemented with 10% heat inactivated fetal calf serum (FCS; Sigma, Poole, UK), 2 mM L-glutamine, 100 units/ml penicillin, 100 μg/ml streptomycin, and 1% non-essential amino acids (Life Technologies, Paisley, UK). Cells (1×10⁶) were seeded into 25 cm² area flasks and maintained at 37°C in 5% CO₂.

H pylori culture and cytokine stimulation protocols

H pylori strain NCTC 11637 (CagA/VacA positive) was grown on blood agar base 2 plates supplemented with 5% (v/v) horse serum and quantified by measurement of optical density at 450 nm. For bacterial/epithelial coculture experiments, the bacteria were resuspended in phosphate buffered saline and applied onto the confluent cell layer that was maintained in serum free (0.5% fetal calf serum) media as before, without antibiotics.

Prior to use, the bacteria were resuspended in phosphate buffered saline and quantified by measurement of optical density at 450 nm. For bacterial/epithelial coculture experiments, the confluent epithelial cell layer was maintained in RPMI 1640 media as before, without antibiotics.

For cytokine stimulation experiments, the confluent cell layer was maintained in serum free (0.5% fetal calf serum) media for 16 hours before addition of various cytokines. Recombinant interleukin (IL)-1β, tumour necrosis factor α (TNF-α), interferon γ (IFN-γ), and IL-17 were obtained from R&D Systems (Abingdon, UK).

Patients and tissues

This study was approved by the Research Ethics Committee of the East London and City Health Authority. For reverse transcription-polymerase chain reaction (RT-PCR) analysis, mucosal antral biopsies were obtained from patients with histologically proved active gastritis (n=10, H pylori positive) and 12 control patients endoscoped for symptoms of non-ulcer dyspepsia or reflux in whom biopsies showed no significant histological abnormality. Biopsies were immediately snap frozen and stored at −70°C. Patients who had received anti-inflammatory, immune modulating, or antisecretory drugs in the previous month were excluded. H pylori infection was determined by biopsy urease test (CLOtest) and by histological examination of tissue samples using cresyl-fast violet staining as well as routine haematoxylin and eosin staining.

RNA standard for quantitative RT-PCR

To allow quantification of hBD1 and hBD2 mRNA expression by RT-PCR, we constructed a plasmid that encodes a standard hBD2 transcript. The above primer sequences were cloned into plasmid pMBEK as described in detail previously. To generate standard RNA, the plasmid was linearised with Hind III and transcribed in vitro using T7 RNA polymerase under in vitro transcribed in vitro using T7 RNA polymerase under standard conditions recommended by the supplier (Promega, Southampton, UK). Using the same primer set, RT-PCR for hBD1 yielded a standard PCR product of 434 bp whereas the target size was 253 bp. Similarly, the size difference between the standard hBD2 PCR product (486bp) versus the target tissue PCR product (276 bp) allowed quantification (number of mRNA transcripts/μg] total cellular RNA) of each of the β-defensins.

Semiquantitative RT-PCR was carried out on the gastric biopsy samples. Expression of hBD1 and hBD2 was normalised to expression of the housekeeping gene glyceraldehyde-3-phosphate dehydrogenase (GAPDH). GAPDH primers used in this study were: sense primer 5′-CACTGGCCGCTGGCAAGGC-3′ and antisense primer 5′-GCCCAGGAGGCTCCCACACACCTCGTG-3′.

RNA isolation and PCR amplification

All reagents for RNA extraction and RT-PCR analysis were obtained from Life Technologies. Total RNA isolation from cells (AGS, MKN7, and MKN45) and mucosal gastric biopsies were performed using a monophasic solution of phenol and guanidine thiocyanate (TRIzol), as recommended by the suppliers. Serial 10-fold dilutions of the standard (1 ng to 100 fg) were cotranscribed with total cellular RNA (~5 μg) to generate cDNA using Moloney murine leukaemia virus reverse transcriptase. PCR was routinely performed using 0.5 units of Taq polymerase, 20 pmol of each oligonucleotide primer, 1.5 mM MgCl₂, and 200 μM deoxynucleotide triphosphates in a final reaction volume of 50 μl. The amplification cycle was as follows: 94°C denaturation for 90 seconds; 58°C annealing for 90 seconds; and 72°C extension for 90 seconds; for 37 cycles. Following amplification, the PCR products were analysed on 2% agarose gels and band intensities were quantified by densitometry (Kodak Digital Science Electrophoresis Documentation and Analysis System 120; Life Technologies). The ratios of band intensities of the PCR products from the standard RNA and target RNA were plotted against the starting number of standard RNA molecules on a double logarithmic scale. In this way, the point at which the starting number of standard RNA transcripts is equal to the starting amount of cellular target RNA can be determined. This technique allows us to quantify the number of mRNA transcripts in a sample to a sensitivity of 10⁻¹⁴ molecules/μg total RNA.

Semiquantitative RT-PCR was performed on biopsy specimens. cDNA was prepared from the samples as described above and the product was divided into three aliquots. PCR for hBD1, hBD2, and the housekeeping gene GAPDH was terminated after 37 cycles of amplification where we obtained optimum product yield in the linear phase of the reaction. The resulting products were analysed as above.

Immunohistochemistry

Immunohistochemistry was performed on archival paraffin blocked full thickness samples of resection tissue and biopsies from patients with varying grades of gastritis. Dewaxed and rehydrated 4 μm sections were initially incubated for 15 minutes in a 3% H₂O₂/methanol mix to inhibit endogenous peroxidase activity. For both hBD1 and hBD2 staining, antigen retrieval was performed by microwaving sections in 0.1 M citrate buffer at pH 6.0 for 30 minutes before further incubation with neat horse serum for 15 minutes in order to reduce non-specific binding. Sections were incubated with primary polyclonal antibodies for hBD1 and hBD2 (raised in rabbit) at 1:800 dilution overnight at 4°C. Both positive (skin tissue) and negative (without primary antibody) controls were also included. After appropriate washes, the sections were incubated with biotinylated antirabbit IgG antibody, followed by avidin-peroxidase complex for 30 minutes at room temperature. Visualisation was performed using the 3, 3′-diaminobenzidine method. All reagents used in the above procedure were purchased from Vector Laboratories Ltd (Peterborough, UK).

Statistical analysis

Comparisons of hBD1 and hBD2 mRNA expression among the various groups of patients was performed using the Mann-Whitney U test.
with Helicobacter pylori

Figure 2

A representative quantitative reverse transcriptions/mRNA expression levels by Irrespective of this basal difference, the increase in hBD1 incubation period. Constitutive expression of hBD1 was both AGS (fig 1A) and MKN7 (fig 1B) cell lines over a 24 hour increasing number of infecting organisms was observed in A dose dependent increase in hBD1 mRNA expression with increasing multiplicity of infection (MOI) of 100. A representative quantative RT-PCR gel showing increased hBD1 mRNA expression (that is, target band) with increasing MOI (10 bacteria/cell). Similar high levels of hBD2 expression were achieved in each cell line. On maximal stimulation there was no significant difference between expression levels of hBD1 and hBD2. Interestingly, hBD1 and hBD2 expression was found to be constitutive and high (>10^7 transcripts/µg total RNA) in MKN45 cells (data not shown), suggesting dysregulation of defensin gene expression in this cell line.

RESULTS

A dose dependent increase in hBD1 mRNA expression with increasing number of infecting organisms was observed in both AGS (fig 1A) and MKN7 (fig 1B) cell lines over a 24 hour incubation period. Constitutive expression of hBD1 was sixfold lower in MKN7 cells compared with the AGS cell line. Irrespective of this basal difference, the increase in hBD1 mRNA expression levels by H. pylori was found to be similar in both cell lines. The maximum level of expression was observed at a multiplicity of infection (MOI) of 100. A representative quantitative RT-PCR gel showing increased hBD1 mRNA expression with increasing MOI is shown in fig 2. No hBD2 was detected in control cells but on infection with the bacterium there was a significant increase in hBD2 mRNA expression, even at low MOI (10 bacteria/cell). Similar high levels of hBD2 expression were achieved in each cell line. On maximal stimulation there was no significant difference between expression levels of hBD1 and hBD2. Interestingly, hBD1 and hBD2 expression was found to be constitutive and high (>10^7 transcripts/µg total RNA) in MKN45 cells (data not shown), suggesting dysregulation of defensin gene expression in this cell line.

Due to the low constitutive expression of hBD1 in MKN7 cells, this cell line was chosen for further analyses. The kinetics of induction of hBD1 and hBD2 were studied at an MOI of 100, as this ratio of infectivity was found to be most potent, as shown above. Time course experiments showed that expression of hBD2 began 3–4 hours after infection (fig 3), then markedly increased to attain maximal expression at eight hours, followed by declining levels at 12 and 24 hours. In contrast, induction of hBD1 mRNA expression was much slower than that for hBD2 with the first increase observed at eight hours and maximum expression occurring at 12 hours post-infection.

IL-1β increases expression of hBD1 and hBD2 in MKN7 cells

Several previous studies have shown that proinflammatory cytokines such as IL-1β and TNF-α can increase β-defensin expression.21 We have also examined the effect of a panel of proinflammatory cytokines (IL-1β, TNF-α, IFN-γ, IL-17) on hBD1 and hBD2 mRNA expression in MKN7 cells. Among the cytokines tested, IL-1β (5 ng/ml) was found to be the only stimulant for hBD1 and hBD2 in this cell line. As observed during infection with H. pylori, IL-1β also caused an increase in hBD1 and hBD2 mRNA expression in a time dependent manner, with a similar earlier maximal increase in hBD2 expression (12 hours v 16 hours for hBD1; fig 4). However, the magnitude of induction of both peptides was much less and

![Figure 3](http://gut.bmj.com/) Time dependent increase in human β-defensin 1 and 2 (hBD1, hBD2) mRNA expression by Helicobacter pylori. MKN7 cells were coinfected at a multiplicity of infection (MOI) of 100 for 24 hours. hBD2 upregulation was initially noted at three hours with a maximum increase observed at eight hours. hBD1 mRNA expression was much slower with a maximum increase observed at 12 hours. Results are mean (SEM).

![Figure 4](http://gut.bmj.com/) Interleukin 1β (IL-1β) is a stimulant of human β-defensin 1 and 2 (hBD1, hBD2) mRNA expression. MKN7 cells were incubated with IL-1β (5 ng/ml) for 24 hours. As seen in fig 2, the kinetics of the increase in hBD2 expression were more rapid than for hBD1. The magnitude of response was much lower than that observed with Helicobacter pylori.
delayed in IL-1β treated cells when compared with *H pylori* infected cells.

**In vivo expression of hBD1 and hBD2**

To resolve the potential effect of *H pylori* colonisation on hBD1 and hBD2 mRNA expression in the gastric mucosa, semiquantitative RT-PCR analyses was performed on biopsies from patients with gastritis and compared with control tissue (fig 5A, B). Control biopsies showed constitutive expression of hBD1 whereas hBD2 mRNA was undetectable. Expression of hBD2 was markedly upregulated in *H pylori* positive gastritis (*p*<0.001). Upregulation of hBD1 also achieved statistical significance (*p*<0.05) in gastritis compared with control tissue.

To further confirm increased expression of hBD1 and hBD2 in the gastric epithelium of *H pylori* positive gastritis, immunohistochemistry was performed on archival paraffin sections of gastric biopsy specimens. Increased positive staining for hBD1 and hBD2 peptide expression was observed in gastritis (fig 6C, D) compared with control tissue (fig 6A, B). Staining was confined to the surface epithelium and the gastric pits of the gastric gland, with no staining detected in the lamina propria or in inflammatory cells of the gastric mucosa.

**DISCUSSION**

*Helicobacter pylori* plays a major role in the pathogenesis of chronic antral gastritis and peptic ulcer disease. There is accumulating evidence that both bacterial factors and subsequent host defence responses contribute to the ongoing inflammatory processes that determine the clinical manifestations of the disease.

During infection, detection of chemokines (C-C and C-X-C) and early recruitment of neutrophils in the gastric mucosa are indicative of the active participation of innate immune mechanisms. In the present study, we have expanded the repertoire of the innate immune responses by the host to *Helicobacter* infection. Studies to date have shown constitutive expression of hBD1 by various mucosal surface epithelia, suggesting a "surveillance-like" role for this peptide in maintaining tissue homeostasis in the absence of infection. In the present study, we show that the constitutive expression of hBD1 can be further modulated during infection and inflammation. Dale et al have also recently noted similar modulation of hBD1 expression in inflamed periodontal tissue. These two studies to date therefore imply a greater role for hBD1 in innate host defence than was previously thought. At present, regulation of hBD1 gene and peptide expression during infectious and inflammatory episodes remains unclear. Using the sensitive technique of competitive RT-PCR, we observed induction of hBD1 mRNA expression in AGS and MKN7 cells 24 hours after infection with *H pylori*. We observed great variation in basal expression of hBD1 in different gastric epithelial cell lines. For example, initial experiments conducted in MKN45 cells showed a spontaneous high level expression of
hBD1 and hBD2 in unstimulated cells which was not modulated by the bacterium. O’Neil et al have recently reported no change in constitutive expression of hBD1 in AGS cells by H pylori (MOI of 10). 16 In contrast, our coulture infection experiments were conducted at a greater multiplicity of infection (MOI 100), and the initial increase in hBD1 expression was first detected only at eight hours. The combined effect of variation in basal hBD1 levels in different cell lines, bacterial strains, and variable numbers of infecting organisms may partly explain the results obtained in different laboratories.

Unlike hBD1, hBD2 peptide is not expressed by epithelia in the absence of infection. As expected, no hBD2 was detected in control gastric cell lines and in control gastric biopsy tissue. We have confirmed upregulation of hBD2 mRNA by H pylori, as reported previously. 17 18 In time course studies, we observed marked upregulation of hBD2 expression as early as three hours post-infection in MKN7 cells. The kinetics of hBD1 induction in comparison were much slower as the first increase was noted only at eight hours post-infection. At present, the identity of both bacterial virulence factors and host signalling events involved in the induction of defensin expression is limited. Although a crucial role for the bacterial “cag pathogenicity island” in induction of hBD2 mRNA expression has been observed, 19 its potential role in the regulation of hBD1 expression has not been studied to date. Wada et al have recently shown unequivocally the involvement of the transcription factor nuclear factor κB in induction of hBD2. 20 In contrast, no nuclear factor κB binding sites have been identified in the promoter sequence of hBD1. 21 The molecular mechanism(s) that may play a part in the activation of these peptides requires further investigation.

Several studies have shown that proinflammatory cytokines such as IL-1β and TNF-α can directly increase expression of mammalian defensins. 15 16 22 In the present study, we explored the effects of various cytokines on expression of hBD1 and hBD2 in gastric epithelial cell lines and found IL-1β to be a potent stimulant of hBD2 in all three gastric cell lines tested. In contrast, the effect of IL-1β on hBD1 expression was only observed in MKN7 cells with no induction in AGS and MKN45 cells. It is probable that the high basal levels of hBD1 observed in AGS and MKN45 cell lines may contribute to masking of cytokine response. Interestingly, two previous studies have reported no effect of IL-1β on hBD1 expression. 23 24 As different cancer cell lines were used in the various studies, the data taken together suggest that the degree of cellular differentiation may play a role in β-defensin gene regulation. Recent studies by El-Omar et al have suggested that the presence of IL-1 gene cluster polymorphism in susceptible individuals is associated with an increased risk of H pylori induced hypochlorhydria and gastric cancer. 25 At present, the role of β-defensins during inflammatory episodes is unclear but specific induction by IL-1β may have major consequences in the innate host defence response to bacterial colonisation and pathogenesis of gastritis and related complications.

To confirm our observations of increased hBD1 and hBD2 expression by H pylori in tissue culture cell lines, we analysed mRNA and peptide expression of both peptides in gastric biopsy specimens. Using semiquantitative RT-PCR, we found increased expression of hBD1 and hBD2 in H pylori mediated gastritis. As no hBD2 mRNA was detectable in control tissue, the increase observed in gastritis was found to be much greater than that observed for hBD1. Immunohistochemical analyses revealed a parallel increase in both hBD1 and hBD2 peptide expression during bacterial induced inflammation. This pilot study on analyses of hBD1 and hBD2 expression from human gastric biopsies suggests a potential role for these peptides in host innate defence to infection and inflammation. H pylori induced chronic inflammation is dependent on the persistence of the organism in the gastric antrum and/or fundus. Several factors such as the low immunological activity of H pylori lipopolysaccharide, bacterial expression of Lewis antigen, interference with antigen processing and presentation, and increased expression of IL-10 have all been implicated in the increased colonisation and ineffective clearance of H pylori from the gastric mucosa. The potential role and contribution of antimicrobial peptides at the surface of the gastric epithelium and their interactions with factors that favour colonisation of the bacterium are still unclear and warrant further investigation.

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