

# Gastric T lymphocyte responses to *Helicobacter pylori* in patients with *H pylori* colonisation

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## Abstract

***Helicobacter pylori* has been identified as a dominant factor in the pathogenesis of duodenal ulcer. The aim of this study was to examine peripheral blood and gastric lymphocyte proliferation and cytokine production in patients with *H pylori* colonisation. Sixty five dyspeptic patients attending for endoscopy were studied; 35 of these were *H pylori* positive and 30 *H pylori* negative as assessed by culture, histology, and rapid urease test. *H pylori* antigen was capable of stimulating peripheral blood lymphocyte proliferative responses even in *H pylori* negative patients. Peripheral blood lymphocyte proliferative responses to *H pylori* (but not to purified protein derivative or phythaemagglutinin) were significantly lower in *H pylori* positive than *H pylori* negative patients. Similarly, antigen specific proliferative responses and interferon  $\gamma$  production by gastric lamina propria lymphocytes were also depressed in *H pylori* positive patients compared with *H pylori* negative patients. CD8 and CD22 positive lamina propria lymphocytes were increased in *H pylori* positive patients. These data show that antigen specific responses to *H pylori* are significantly lower in *H pylori* positive patients and could indicate activation of antigen specific suppression**

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*Helicobacter pylori* (*H pylori*, previously known as *Campylobacter pylori*) is a spiral organism<sup>1,2</sup> that colonises the human gastric epithelium.<sup>3</sup> It is strongly associated with chronic type B gastritis and peptic ulcer disease.<sup>4-7</sup> Eradication of the bacterium is associated with healing of gastric and duodenal ulcer and longterm remission suggesting that this is a dominant factor in the pathogenesis of peptic ulcer.<sup>8-10</sup> *H pylori* colonises the gastric antrum but can also be found in the corpus and duodenum in regions of gastric metaplasia.<sup>11</sup> Colonisation of the stomach with *H pylori* is accompanied in the acute stage by an increased number of neutrophils in the lamina propria. In the chronic stage, an increase in the number of lymphocytes, plasma cells, and eosinophils in the lamina propria is evident.<sup>12</sup> The infection is also associated with both a local and systemic specific antibody immune response.<sup>13-15</sup> The contribution, however, of T cell responses to host defences to *H pylori* is little understood. It has previously been reported that peripheral blood lymphocyte

responses to *H pylori* are reduced in *H pylori* positive patients.<sup>16,17</sup> It has been suggested that reduction in peripheral blood lymphocyte responses might simply reflect a selective accumulation of antigen specific T cells in mucosal sites. In this study, we have developed a system to examine antigen specific T cell responses to *H pylori* in the gastric lamina propria lymphocytes. Gastric lymphocytes were isolated by a modification of techniques developed for the isolation of small bowel lymphocytes from intestinal biopsy specimens.<sup>18</sup> Antigen specific responses could be analysed if an exogenous source of antigen presenting cells and cytokine was provided in the form of irradiated autologous peripheral blood mononuclear cells and low dose interleukin 2, respectively.

## Methods

### SUBJECTS

Sixty five dyspeptic patients attending for upper gastrointestinal endoscopy for dyspepsia were studied (35 females, 30 males, age range: 17-78 years; mean age: 49.6), all of whom had antral biopsies performed. None of the patients studied had received non-steroidal anti-inflammatory drugs, bismuth compounds or antibiotics recently. Patients with evidence of malignant disease or immunosuppression were excluded. Multiple biopsy specimens were obtained during upper gastrointestinal endoscopy from adjacent sites of the gastric antrum for *H pylori* culture, histological examination, and rapid urease test (CLO test). *H pylori* was identified histologically by a Giemsa stain. *H pylori* were grown on 7% lysed horse blood agar under microaerophilic conditions at 37°C for three days. Patients were designated as *H pylori* positive on the basis of CLO testing. Preliminary analysis on 37 patients showed that CLO testing had a 94% sensitivity for *H pylori* positivity with respect to culture and a 100% specificity. Table I summarises the patient characteristics.

### ANTIGEN AND MITOGEN PREPARATION

*H pylori* antigen was prepared from a mixture of *H pylori* cultures obtained from six patients. Bacteria cells were harvested in phosphate buffered saline and then washed ( $\times 3$ ) in phosphate buffered saline by centrifugation (12 000  $\times g$ ) for 15 minutes at 10°C. The cells were resuspended in phosphate buffered saline (1:2 vol/vol). This suspension was then sonicated on ice using 6  $\times$  15 second 100 Watt pulses, with 30 second cooling intervals in between, using a DAWE Soniprobe 7532A.<sup>19</sup>

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TABLE I Characteristics of the patients studied for peripheral blood mononuclear cells and lamina propria lymphocytes response. Patients were designated as *H pylori* positive on the basis of CLO testing. Preliminary analysis of 37 patients showed that CLO testing has a 94% sensitivity for *H pylori* positivity with respect to culture and a 100% specificity. Patient characteristics are summarised in the Table

Characteristics	HP+ve on the CLO test (n=33) (%)	HP-ve on the CLO test (n=32) (%)
Endoscopic gastritis	30/33 (91) including 2 cases of peptic ulcer	5/32 (16)
Histological gastritis	33/33 (100)	4/32 (13)
HP seen on staining	31/33 (94)	0/32 (0)

HP=*H pylori*.

The *H pylori* protein concentration in the preparation was measured by the method of Bradford using bovine serum albumin as the standard.<sup>20</sup> The sonicated preparation of *H pylori* was irradiated at 400 Gy and stored at  $-20^{\circ}\text{C}$ . Purified protein derivative (DK-2300 Copenhagen, Denmark) and phythaemagglutinin (Sigma Chemical, St Louis, MO, USA) were used at a concentration of 100 U/ml and 10  $\mu\text{g/ml}$ , respectively. *Escherichia coli* (*E coli*) antigen was prepared by inactivating the bacteria by irradiation and sonication as described above and used at a protein concentration of 2 mg/ml. The optimum concentration of *H pylori* for use in the lymphocyte proliferation studies was determined in preliminary experiments.

#### PERIPHERAL BLOOD LYMPHOCYTE PROLIFERATION STUDIES

Peripheral blood samples were obtained by venesection for isolation of peripheral blood mononuclear cells. Peripheral blood mononuclear cells were separated by Ficoll-hypaque density gradient centrifugation at  $250\times g$  for 30 minutes at  $4^{\circ}\text{C}$  (Lymphoprep Nycomed Pharma AS, Oslo, Norway), then washed ( $\times 3$ ) and resuspended in RPMI 1640 (Gibco, Life Technologies Ltd, Paisley, Scotland) medium containing 100 U/ml penicillin, 100  $\mu\text{g/ml}$  streptomycin, 2 mM L-glutamine, and 10% fetal calf serum. To assess antigen specific lymphocyte proliferation,  $1\times 10^6/\text{ml}$  peripheral blood mononuclear cells were cultured in a total volume of 200  $\mu\text{l}$  for five days with varying amounts of *H pylori*, purified protein derivative, phythaemagglutinin, *E coli* or without stimulant (spontaneous cultures).  $^3\text{H}$ -thymidine

TABLE II Proliferative response of peripheral blood mononuclear cells to *H pylori*. Results are expressed as  $^3\text{H}$ -thymidine incorporation (cpm) in peripheral blood mononuclear cells cultured for five days

	$^3\text{H}$ -thymidine incorporation (cpm)		p Value
	<i>H pylori</i> (+) (n=21)	<i>H pylori</i> (-) (n=16)	
Phythaemagglutinin (10 $\mu\text{g/ml}$ )	9510 (1552)	10 417 (2156)	0.72
Purified protein derivative (100 U/ml)	4771 (1613)	8 109 (2466)	0.08
<i>H pylori</i> (3 $\mu\text{g/ml}$ )	1601 (555)*	3 946 (778)*	0.018
<i>E coli</i> (2 $\mu\text{g/ml}$ )	529 (155)	752 (182)	0.39
Spontaneous RPMI	745 (190)	954 (221)	0.35

\*Proliferative responses were significantly lower ( $p<0.02$ ) in the *H pylori* positive group compared with *H pylori* negative patients. There were no significant differences between the two groups after stimulation with phythaemagglutinin, purified protein derivative, *E coli*, and RPMI control. Data shown as mean (SEM).

(1  $\mu\text{Ci/ml}$ , specific activity 74 GBq/mmol, NEN, Boston, MA) was added for the last 24 hours of culture. All samples were measured in triplicate. The cultured cells were harvested on Whatman paper using a cell harvester (Multimash 2000, Dynatech). Radioactivity incorporated into cells were determined by liquid scintillation counting using a Packard Tri-Carb scintillation counter. Lymphocyte proliferation assays were expressed as  $^3\text{H}$ -thymidine incorporation (cpm).

#### GASTRIC MUCOSAL LYMPHOCYTE CULTURE

Lamina propria lymphocytes were separated by modification of techniques developed for the isolation of lymphocytes from duodenal mucosal biopsy specimens.<sup>21</sup> Five gastric mucosa specimens were placed in Hanks's balanced salt solution without calcium and magnesium (CMF HBSS Gibco, +5% fetal calf serum) containing 1 mM dithiothreitol (Sigma) and 1 mM ethylenediamine tetraacetic acid (Analar, BDH Chemicals Ltd, Poole, England) in a 20 ml V bottomed container. The specimens were then agitated for one hour at  $37^{\circ}\text{C}$  (Matburn blood cell suspension mixer), to remove the epithelial layer and washed with RPMI 1640. The specimens were treated with collagenase, type I (120 U/ml, Sigma) for three hours at  $37^{\circ}\text{C}$  with agitation. The cellular supernatant was then washed three times and the number and viability ( $>85\%$ ) of the isolated lamina propria lymphocytes determined using acridine orange/ethidium bromide. Viability of lamina propria lymphocytes was consistently greater than 85%. Lymphocytes were cultured ( $4\times 10^5/\text{ml}$ ) with *H pylori* (300  $\mu\text{g/ml}$ ), phythaemagglutinin (5  $\mu\text{g/ml}$ ), and anti-CD3 (OKT3, 1:50) for three days (at  $37^{\circ}\text{C}$  in 5% carbon dioxide) with irradiated (250 Gy) autologous peripheral blood mononuclear cells ( $2\times 10^6/\text{ml}$ ) in a final volume of 200  $\mu\text{l}$  in RPMI 1640 medium containing interleukin 2 (2 U/ml) in 96-well U bottom microplates.  $^3\text{H}$ -thymidine was added for the last 24 hours of culture. All samples were measured in triplicate. The cultured cells were processed as described earlier.

#### B LYMPHOCYTE ELIMINATION FROM LAMINA PROPRIA

Lamina propria lymphocytes were incubated with Leu 14 (CD22): (5  $\mu\text{l}$  for every  $2\times 10^5$  positive cells) for 45 minutes at  $4^{\circ}\text{C}$  with mixing and then washed. Magnetic beads coated with goat antimouse immunoglobulin G (IgG) (Dynabeads M-450) were added to the cell suspension: (10 beads:1 positive cell) and the mixture was then incubated for 50 minutes at  $4^{\circ}\text{C}$  with mixing. RPMI 1640 (4 ml) was then added and the B cells were removed on a magnetic particle concentrator (DYNAL MPC 1).<sup>22 23</sup> Non-adherent cells were collected and CD22 positive cells quantitated by flow cytometry before and after CD22 depletion. The cellular supernatant was then washed three times and the number and

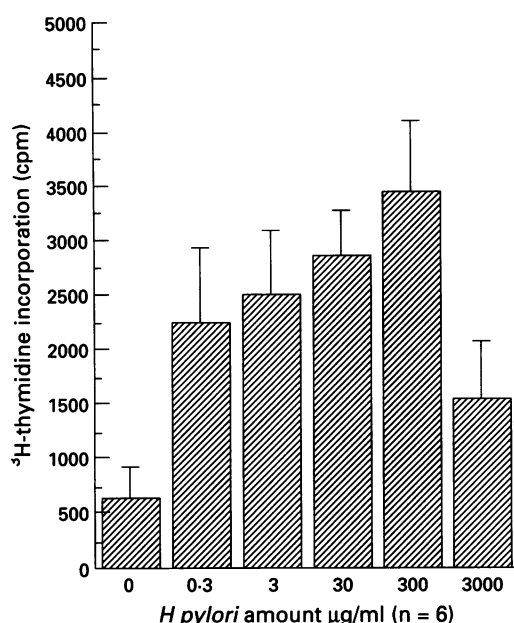


Figure 1: Dose response curve for the proliferation of lamina propria lymphocytes in response to H pylori. The effect of different amounts of H pylori on <sup>3</sup>H-thymidine incorporation (cpm) into lamina propria lymphocytes ( $4 \times 10^5$ /ml) was estimated in the presence of irradiated (250 Gy) autologous peripheral blood mononuclear cells ( $2 \times 10^6$ /ml) and interleukin 2 (2 U/ml) after a three day culture. Maximal proliferation was found at 300 µg/ml and this amount was used in subsequent experiments.

viability (>85%) of the isolated cells after CD22 depletion was determined.

#### INDUCTION AND ASSAY OF INTERFERON $\gamma$

Peripheral blood mononuclear cells ( $1 \times 10^6$ /ml) and lamina propria lymphocytes ( $4 \times 10^5$ /ml) were cultured with inactivated H pylori (3 µg/ml, 300 µg/ml), purified protein derivative (100 U/ml), phythaemagglutinin (10 µg/ml, 5 µg/ml), OKT3 (1:50), and RPMI 1640 only in RPMI 1640 for either five days or three days at 37°C in 5% carbon dioxide. The culture supernatants were collected and stored at -70°C.

Interferon  $\gamma$  was measured by enzyme linked immunosorbent assay (ELISA) using anti-interferon  $\gamma$  antibodies kindly donated by Dr Kingston Mills, National Institute for Biological Standards, UK. Purified mouse anti-interferon monoclonal Ab was coated (50 µl/well) onto 96 well round bottomed ELISA plates and incubated for two hours at 37°C. The wells were then blocked overnight

(4°C) with 150 µl phosphate buffered saline containing bovine serum albumin (0.5%, wt/vol).

The plates were washed four times with phosphate buffered saline-TWEEN 20 (200 µl/well). Supernatants were added (50 µl/well) in triplicate. Triplicate serial dilutions of interferon  $\gamma$  (100 IU to 1.5 IU/ml) were added to standard wells. The plates were incubated for one hour at 37°C and then washed four times with phosphate buffered saline-TWEEN 20.

Rabbit polyclonal anti-interferon  $\gamma$  was added (50 µl/well) in bovine serum albumin-phosphate buffered saline and incubated for one hour at 37°C. After washing the plates biotinylated antirabbit IgG (50 µl/well) was added in bovine serum albumin-phosphate buffered saline and incubated for one hour at 37°C. Streptavidin biotinylated horseradish peroxidase conjugate (50 µl/well) in bovine serum albumin-phosphate buffered saline was added and incubated for 30 minutes at 37°C, washed four times with bovine serum albumin-phosphate buffered saline followed by two washes with 0.1 M citrate phosphate buffer (pH 5.0).

The substrate, o-phenylenediamine (1 mg/ml in citrate phosphate buffer) was then added (50 µl/well) and the reaction was developed in the dark at room temperature. The reaction was stopped after 10 to 15 minutes by addition of 1 M H<sub>2</sub>SO<sub>4</sub> (50 µl/well). The plates were read using an ELISA reader at 450 nm. A standard curve was constructed and the amount of interferon  $\gamma$  present in the supernatant was determined with reference to this standard curve. The assay sensitivity was 2 IU/ml.<sup>24</sup>

#### LAMINA PROPRIA LYMPHOCYTE PHENOTYPIC ANALYSIS

Lamina propria lymphocytes were isolated as described above. The number and viability of the isolated cells were determined using acridine orange/ethidium bromide. One  $\times 10^5$  cells were washed with phosphate buffered saline containing bovine serum albumin (1%, wt/vol) and sodium azide (0.02%, wt/vol). Monoclonal antibodies including Leu 4 (CD3), Leu 3a (CD4), Leu 2a (CD8), Leu M3 (CD14), Leu 14 (CD22), and Leu 19 (CD56), control phycoerythrin and fluorescein isothiocyanate were added (5 µl) to the lamina propria lymphocytes, respectively. Lymphocyte subsets were analysed by a combination of direct and indirect immunofluorescence. Data were acquired on a Beckton Dickinson FACSCAN and analysed using Lysis 3 software. Lamina propria lymphocyte phenotypic analysis was expressed as a percentage of the positive cells. All the monoclonal antibodies were obtained from Becton Dickinson Immunocytometry Systems, San Jose, CA, USA.

#### STATISTICAL ANALYSIS

Data are expressed as mean (SEM). Differences between the results obtained with H pylori

TABLE III Proliferation of lamina propria lymphocytes in gastric mucosa in H pylori positive (n=14) and negative (n=14) patients

	<sup>3</sup> H-thymidine incorporation (cpm)		
	H pylori (+)	H pylori (-)	p Value
Phythaemagglutinin (5 µg/ml)	7171 (1613)	7755 (156)	0.32
OKT3 (1:50)	6104 (493)	6213 (821)	0.25
H pylori (300 µg/ml)	1400 (448)*	3013 (418)*	0.015
Spontaneous RPMI	562 (192)	650 (281)	0.22
Irradiated feeder cells alone	435 (110)	502 (150)	0.43

Results are expressed as <sup>3</sup>H-thymidine incorporation (cpm) in lamina propria lymphocytes ( $4 \times 10^5$ /ml) cultured for 72 hours with autologous irradiated (250 Gy) peripheral blood mononuclear cells ( $2 \times 10^6$ /ml) containing interleukin 2 (2 U/ml). \*Lamina propria lymphocyte response to H pylori in culture was significantly lower (p<0.02) in H pylori positive subjects compared with H pylori negative subjects. Data shown as mean (SEM).

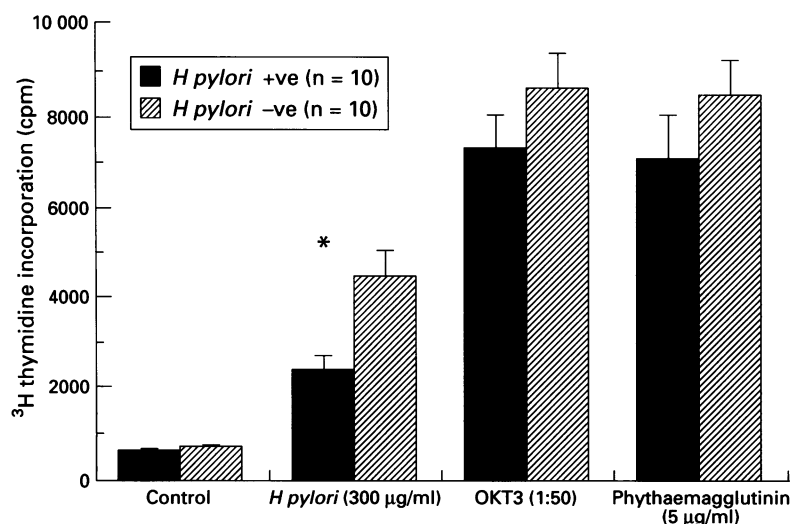


Figure 2: Response of B lymphocyte depleted lamina propria T lymphocytes to *H pylori* antigens. Lamina propria lymphocytes were depleted of B lymphocytes by magnetic bead separation. Proliferation was assessed in the presence of irradiated peripheral blood mononuclear cells as accessory cells and interleukin 2. \*Lamina propria T lymphocyte responses to *H pylori* (300 µg/ml) in culture were significantly lower in *H pylori* positive patients compared with *H pylori* negative patients ( $p < 0.02$ ).

positive and negative groups were analysed using the two tailed Mann-Whitney U test.

## Results

### PROLIFERATIVE RESPONSES OF PERIPHERAL BLOOD MONONUCLEAR CELLS IN RESPONSE TO *H PYLORI* ANTIGEN

Lymphocyte proliferative responses were performed using *H pylori* antigen (3 µg/ml), a dose found to give maximal peripheral blood mononuclear cells proliferation results in preliminary experiments. Lymphocyte proliferative responses were detected in both *H pylori* positive and negative subjects. Proliferative responses were significantly lower in the *H pylori* positive subjects ( $p < 0.02$ ). No significant differences were seen between the *H pylori*

positive and negative groups in the presence of purified protein derivative, phythaemagglutinin, and *E coli* (Table II). While proliferative response using purified protein derivative seems to be lower in the *H pylori* positive group, this did not show statistical significance.

### PROLIFERATIVE RESPONSES OF GASTRIC MUCOSA LAMINA PROPRIA LYMPHOCYTES IN RESPONSE TO *H PYLORI* ANTIGEN

A high dose of *H pylori* (300 µg/ml) resulted in optimal stimulation of lamina propria lymphocytes and this amount was used in subsequent experiments (Fig 1). This may reflect a lower efficiency of antigen processing by the irradiated antigen presenting cells used in these assays. In addition low amounts of interleukin 2 were required to induce proliferation in this system. Exogenous interleukin 2 was added at the lowest concentration that permitted measurement of proliferation. No proliferation was seen in the absence of irradiated peripheral blood feeder cells or exogenous interleukin 2. Also no proliferation was seen in the irradiated peripheral blood feeder cells alone in the presence of interleukin 2 and *H pylori* (Table III). Proliferation of gastric lamina propria lymphocytes was found to be maximal at three days in preliminary studies. The proliferative responses to *H pylori* (300 µg/ml) were significantly lower in the *H pylori* positive patients compared with negative patients ( $p < 0.02$ ). There was no significant difference, however, between the two groups in their responses to phythaemagglutinin and OKT3 (Table III).

### B CELL ELIMINATION

After B cell elimination from lamina propria lymphocytes, similar proliferative results were obtained. T lymphocyte responses to *H pylori* (300 µg/ml) were statistically lower mean (SEM) (2356 (342) v 4425 (626) cpm,  $p < 0.02$ ) in *H pylori* positive patients compared with the *H pylori* negative patients. There was no significant difference, however, between the two groups in their responses to phythaemagglutinin (7021 (1023) v 8456 (844),  $p = 0.325$ ) and OKT3 (7285 (783) v 8651 (744),  $p = 0.251$ ) (Fig 2).

### INTERFERON $\gamma$ PRODUCTION BY PERIPHERAL BLOOD MONONUCLEAR CELLS AND LAMINA PROPRIA LYMPHOCYTES IN CULTURE

Interferon  $\gamma$  secretion by peripheral blood mononuclear cells in response to *H pylori* antigen was significantly lower in *H pylori* positive patients compared with *H pylori* negative controls ( $p < 0.02$ ). Neither spontaneous production of the interferon  $\gamma$  or interferon  $\gamma$  production in response to phythaemagglutinin (10 µg/ml) stimulation were significantly different in the two groups (Fig 3A). Interferon  $\gamma$  secretion by lamina propria lymphocytes in response to *H pylori* antigen was significantly lower in patients with *H pylori* infection ( $p < 0.05$ ). Neither spontaneous production of interferon  $\gamma$  or interferon  $\gamma$  production in

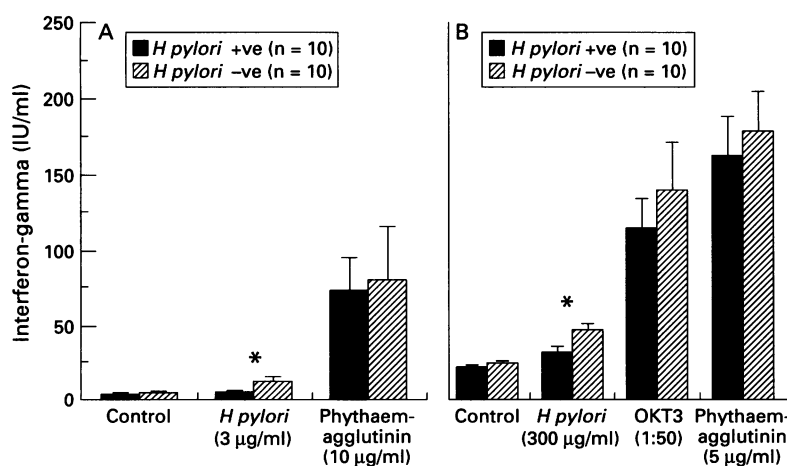


Figure 3: (A) Interferon  $\gamma$  production by peripheral blood mononuclear cells in culture in the presence of *H pylori*, phythaemagglutinin, and RPMI (control). Interferon  $\gamma$  was measured by ELISA. \* Interferon  $\gamma$  production was significantly lower in *H pylori* positive patients ( $p < 0.02$ ) after stimulation with *H pylori* compared with negative controls. (B) Interferon  $\gamma$  values in the supernatant of cultured lamina propria lymphocytes. Interferon  $\gamma$  was measured by ELISA. \* Interferon  $\gamma$  production by lamina propria lymphocytes in response to *H pylori* was significantly lower in *H pylori* infected patients ( $p < 0.05$ ) compared with *H pylori* negative subjects.

TABLE IV Characterisation of lamina propria lymphocyte population

	Percentage of positive cells					
	CD3	CD4	CD8	CD22	CD14	CD56
HP+ve (n=8)	50 (4.9)	19.2 (1.6)	28 (3.1)*	21 (5.2)*	7.7 (1.2)	<2
HP-ve (n=8)	55 (5.1)	21.3 (3.6)	21 (2.1)*	8.4 (1.5)*	6.4 (1.4)	<2

Results are expressed as a percentage of positive cells in lamina propria lymphocyte suspensions. All cells were quantitated by flow cytometry. \*Note that there was a significant ( $p < 0.05$ ) increase in CD8<sup>+</sup> and CD22<sup>+</sup> cells in lamina propria lymphocyte populations from *H pylori* positive subjects, but no significant difference in the number of CD3<sup>+</sup>, CD4<sup>+</sup>, CD14<sup>+</sup>, and CD56<sup>+</sup> cells from either *H pylori* positive or negative groups. Data shown as mean (SEM).

response to phythaemagglutinin (5 µg/ml) and OKT3 (1:50) stimulation were significantly different in the two groups (Fig 3B).

#### CHARACTERISATION OF LAMINA PROPRIA LYMPHOCYTE POPULATIONS

The percentage of CD3, CD4 (helper), CD8 (suppressor), CD22, CD14, and CD56 positive cells in lamina propria lymphocyte populations from *H pylori* positive and negative patients was assessed by flow cytometry. There was a significant ( $p < 0.05$ ) increase in CD8<sup>+</sup> and CD22<sup>+</sup> cells present in lamina propria lymphocytes subpopulations in *H pylori* positive subjects, but no significant difference in the number of CD3<sup>+</sup>, CD4<sup>+</sup>, CD14<sup>+</sup>, and CD56<sup>+</sup> cells in *H pylori* positive and negative groups (Table IV).

#### Discussion

In this study we have shown that an inactivated *H pylori* preparation is capable of stimulating T cell activation as reflected in the secretion of the T cell cytokine interferon  $\gamma$ , as well as T cell proliferation. This activation is seen not only in subjects with previous contact with *H pylori* and established serological immunity but also in subjects who are negative for *H pylori* colonisation. Furthermore, our data suggest a significantly attenuated response to *H pylori* in *H pylori* positive patients. It is conceivable that the observed lymphocyte response reported in *H pylori* negative patients resulted from cross reactivity with certain ubiquitous bacterial antigens. Lymphocyte proliferative responses to *E coli*, purified protein derivative or phythaemagglutinin were not different, however, when compared between the two groups.

These results from peripheral blood cultures are in agreement with those previously described.<sup>16 17</sup> In addition to the results obtained in peripheral blood lymphocyte populations, we have now been successful in examining the proliferative responses of gastric mucosal lymphocytes isolated from endoscopic biopsy specimens. This has permitted direct analysis of antigen specific responses at the site of mucosal inflammation. Our studies have shown an increase in local T cells adjacent to gastric epithelial cells in response to *H pylori* infection.<sup>25</sup> In the development of this assay we have found that irradiated antigen presenting cells and exogenous interleukin 2 were necessary for the stimulation of proliferation. No proliferation was seen in the absence of exogenous interleukin 2 or in the presence of

interleukin 2 U/ml alone (data not shown). No proliferation was seen in the irradiated peripheral blood feeder cells alone in the presence of interleukin 2 and *H pylori*. Gastric mucosal lymphocytes were stimulated by inactivated *H pylori* preparation in both *H pylori* positive and *H pylori* negative patients. Lymphocyte proliferation in *H pylori* positive patients was, however, significantly less intense when compared with *H pylori* negative patients. We excluded the possibility that this was a non-specific B cell response to lipopolysaccharide as we have performed these studies in lymphocyte preparations depleted of B cells by magnetic bead isolation. These results show that lamina propria T lymphocytes exhibit reduced proliferation in response to *H pylori* in *H pylori* positive patients. In addition, interferon  $\gamma$  production was also reduced in *H pylori* positive subjects. Interferon  $\gamma$  production was significantly higher in lamina propria lymphocytes than in peripheral blood mononuclear cells. This may reflect a differing clonal distribution of lymphocytes producing this cytokine at these sites. Data in intestinal T cell clones suggest that the majority of such T cells produce interferon  $\gamma$ . Similarly, intestinal lamina propria lymphocytes seem to be capable of expressing interleukin 2 receptor without proliferating, a finding that suggests that little interleukin 2 is available for stimulation.<sup>26</sup>

There are a number of potential mechanisms for this attenuated response to *H pylori*. Firstly, *H pylori* might produce an inhibitory factor or toxin in vivo, which blocks lymphocyte proliferation. Lymphocyte proliferation in response to the T cell mitogen phythaemagglutinin and anti-CD3 is unaltered, however, by *H pylori* infection suggesting that a non-specific toxin is unlikely. A second mechanism could be antigen specific suppression mediated by CD8<sup>+</sup> suppressor cells. In this regard it is notable that we have shown an increase in the proportion of CD8<sup>+</sup> cells in the lamina propria of *H pylori* positive patients. It is as yet unclear whether these cells have cytotoxic potential or whether they are functioning as suppressor cells. Further studies will be directed to finding out if antigen specific CD8<sup>+</sup> suppressor cells within the lamina propria may be implicated in attenuated responses to *H pylori*. Such cells have been shown in the cutaneous lesion of lepromatous leprosy<sup>27</sup> and are thought to play a part in the nature of the tissue response to *M leprae*.

Many studies have shown the structure and antigenicity of *H pylori* whole cell, outer membrane, acid extractable surface protein, and proteinase K treated whole cell lysate preparations from *H pylori* strains. Antibody reactivity is quite diverse and recognises a wide range of proteins.<sup>28</sup> It is possible that *H pylori* positive patients recognise different epitopes than *H pylori* negative patients. It has been reported that the presence of antibodies reacting with a 120 kDa protein is associated with duodenal ulcer and gastric cancer and it seems that this antigen is associated with strains of increased cytotoxicity.<sup>29 30</sup> It is

possible that exposure to a potent immunogen from a low virulence strain may confer T cell reactivity and protection from infection.

In assessing these data, it is important to consider the question of previous contact with *H pylori*. *H pylori* is an extremely prevalent organism with a high rate of recurrence in the year after eradication.<sup>31</sup> Furthermore, infection may be intrafamilial or sporadic within families suggesting a wide environmental exposure.<sup>32</sup> As yet there is little evidence for elimination of the bacteria in non-susceptible patients. The finding of lower responses to *H pylori*, which we and others have described, however, suggest the possibility that poor T cell responses may be associated with an inability to clear the organism.

DK is a Wellcome Senior Fellow in clinical science.

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