Histological study of chronic gastritis from the United Arab Emirates using the Sydney system of classification

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Abstract

Aims—To determine the prevalence of *Helicobacter pylori* in five main nationality groups with gastric ulcer, duodenal ulcer, and non-ulcer dyspepsia; and to determine the histopathological types of gastritis and assess the graded variables of *Helicobacter* associated gastritis.

Methods—Gastric antral and corpus biopsy specimens from 437 patients were examined for the prevalence of *H pylori*, 337 of which were classified and graded histologically according to the Sydney system.

Results—The overall colonisation rate of *H pylori* was 90%, and there was no significant difference between groups of different ethnic origins. The colonisation rates were 99%, 89%, and 78% in patients with duodenal ulcer, non-ulcer dyspepsia, and gastric ulcer, respectively. *Helicobacter* associated gastritis was the most common form of chronic gastritis (87%). *H pylori* density was greater in the antrum than the body. Gastric atrophy in *helicobacter* associated gastritis was seen in 54% of the cases (43% grade I, 10% grade II, 1% grade III) and increased the older the patients. Atrophy of the corpus alone was very rare (1%). Atrophy and intestinal metaplasia were more prevalent in patients with gastric ulcer than duodenal ulcer.

Conclusion—The colonisation rate of *H pylori* was similar in the five groups studied and was almost invariably present in gastric biopsy specimens in patients with duodenal ulcer. *H pylori* associated gastritis was the most common form of gastritis. Atrophy was mainly of low grade and increased the older the patient.

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A large body of evidence indicates that *Helicobacter pylori* has a role in the pathogenesis of chronic gastritis¹⁻³ and peptic ulcer disease.⁴⁻⁷

Based on several published epidemiological studies, Graham⁶ reported that the prevalence of *H pylori* is higher in developing countries than industrialised countries. In developing countries acquisition of *H pylori* occurs at an early age: Klein *et al* have shown that the prevalence of *H pylori* is 48% in Peruvian children below the age of 12 years.⁸

The prevalence of *H pylori* in gastric mucosa from patients with ulcer and non-ulcer dyspepsia has also been studied.⁹ However, studies concerning the colonisation rate of *H pylori* in gastric biopsy specimens and its role in gastritis and peptic ulceration in patients from Middle Eastern countries are still few in number.¹⁰⁻¹²

There are several classifications for chronic gastritis: morphological¹³⁻¹⁴; topographical¹⁵⁻¹⁶; and combined morphological and topographical.¹⁷ The discovery of *H pylori* as a major cause of gastritis has led many authors to incorporate aetiology in the classification of chronic gastritis.¹⁸⁻²⁰ The Sydney system of classification of chronic gastritis has incorporated topography, morphology, and aetiology in one system.²¹

This system recognises acute, chronic, and special forms of chronic gastritis, permitting differentiation of distinct entities previously classified together under chronic gastritis. The most important characteristic of this system is its grading of five main histological features of gastritis, which allows the changes that occur in gastric mucosa to be assessed accurately.²¹

The aims of this study were to assess the colonisation rate of *H pylori* in gastric biopsy specimens from patients with peptic ulcers and non-ulcer dyspepsia in five main population groups currently living in the United Arab Emirates (UAE). The Sydney system was also applied to the material studied, and the graded variables in *Helicobacter* associated gastritis were assessed quantitatively.

Methods

In this study 437 patients with ulcer and non-ulcer dyspepsia attending five gastroenterology clinics in the northern region of the UAE were studied during December 1990 to May 1992. The mean age was 36.8 years (range 12-100 years) and 152 were female. Only patients with histological evidence of gastritis were included. Patients taking antibiotics before endoscopy over a period of at least six weeks and those with malignant diseases were excluded.

Patients were categorised into five main nationality groups:

- group 1 nationals of the UAE and some other neighbouring countries;
- group 2 nationals of India, Pakistan, and Sri Lanka;
- group 3 nationals of Bangladesh;

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Figure 1 Gastric biopsy specimen of the antrum showing mild atrophy represented by some loss of the gland in a background of mild inflammation (haematoxylin and eosin).

Figure 2 Gastric biopsy specimen of the antrum showing moderate atrophy with more loss of gland in a background of mild inflammation and grade I intestinal metaplasia (AB/PAS stain).

group 4 nationals of Iraq, Iran, and countries north of Saudi Arabia;
group 5 nationals of the following three African countries—Egypt, Sudan, and Somalia.

All non-UAE patients were born in their original countries and had been resident in the UAE for a period of one to 20 years.

Patients were also grouped under three clinical conditions: non-ulcer dyspepsia, duodenal ulcer, and gastric ulcer.

Biopsy specimens were fixed in 10% formalin and routinely processed. Paraffin wax sections were cut at three levels of 4 μ m in thickness and stained with haematoxylin and eosin, Diff-3,²² and alcian blue/periodic acid-Schiff (AB/PAS) stains. All slides were examined for the presence of *H pylori*. Slides from 337 patients were graded according to the Sydney system.²¹ The remaining 100 biopsy specimens were superficial and used only to study the prevalence of *H pylori*. Graded variables were *H pylori* density, inflammation, activity, atrophy (figs 1–3) and intestinal metaplasia.

Reproducibility of grading was assessed blind by examining eight gastric biopsy specimens on 12 different occasions. The coefficient of variations (CV) for all graded variables in both antrum and body mucosa were: 0; 4.21; 4.76; 4.95; 5.10; 6.05; 6.41; and 6.64. Biopsy specimens with a small number of lymphocytes were considered normal apart from bile associated gastritis.²³ One or more aggregates of lymphocytes within the lamina propria with or without lymphoid follicles were regarded as significant.

The criteria described by other authors were used for the diagnosis of bile associated reactive gastritis,²³ drug associated reactive gastritis,²⁴ and lymphocytic gastritis.²⁵

Differences were evaluated for significance using the χ^2 test with the Yates' correction, and results were considered significant if $p < 0.05$ (two tailed probability) was reached.

Results

The colonisation rate of *H pylori* in gastric biopsy specimens from all 437 patients, in relation to the five different ethnic groups and three clinical conditions, is shown in table 1. The difference in the colonisation rates among all the groups was not significant and the overall prevalence was 90%. The colonisation rate was 99% in patients with duodenal ulcer, 89% in patients with non-ulcer dyspepsia, and 78% in those with gastric ulcer. The difference was significant between

Table 1 Prevalence of *H pylori* in 437 patients with ulcer and non-ulcer dyspepsia in relation to five main nationality groups and clinical diagnosis

	Total No of patients	Total positive	Percentage
Group I (UAE)	151	141	93.38
Group II (India)	29	113	87.60
Group III (Bangladesh)	42	38	90.48
Group IV (Jordan, Iraq, Iran)	61	56	91.80
Group V (Africa)	54	47	87.04
Non-ulcer dyspepsia	315	279	88.57*
Duodenal ulcer	99	98	98.99†
Gastric ulcer	23	18	78.26‡
Total	437	395	90.39

p value * $v \dagger p < 0.0015$

† $v \ddagger p < 0.0001$

* $v \ddagger p = \text{Ns}$

Figure 3 Gastric biopsy specimen of the antrum showing severe atrophy. Only a few normal glands remain in a background of mild inflammation and grade I intestinal metaplasia (haematoxylin and eosin).

patients with duodenal ulcer and gastric ulcer ($p < 0.0001$) and between patients with duodenal ulcer and non-ulcer dyspepsia ($p < 0.0015$).

The histological forms of chronic gastritis are shown in table 2. *Helicobacter* associated gastritis accounted for 87% of all cases. Idiopathic gastritis and bile associated reactive gastritis represented the other two major groups. Four patients had mucosal injury caused by non-steroidal anti-inflammatory drugs (NSAIDs), aspirin, and alcohol. Lymphocytic gastritis was seen in two patients who were also positive for *H pylori* (table 2). One patient showed granulomatous gastritis and was also positive for *H pylori*.

Table 3 shows the topographical patterns of *Helicobacter* associated gastritis in the three groups of patients: pangastritis (predominantly antral) was present in 54% of the

patients, followed by pangastritis (35%), antral gastritis (10%), and pangastritis (predominantly corporal) (1%).

Figure 4 shows the three main topographical patterns of *Helicobacter* associated gastritis in relation to age. Pangastritis (predominantly antral) was the main pattern in patients over the age of 30, followed by pangastritis. Antral gastritis was the main pattern in patients under 20 years of age.

The distribution of *H pylori* in antral type and body type gastric mucosa is shown in table 4. One hundred and sixty four out of 294 (56%) patients had *H pylori* equally distributed between antrum and corpus. In 92 (31%) patients, the *H pylori* grade was predominantly higher in antral mucosa than body type mucosa. *H pylori* was identified in the antrum only in 19 (6%) patients. Two (1%) patients showed *H pylori* in the corpus only and atrophy and intestinal metaplasia in the antral mucosa.

Grades of atrophy in *H pylori* associated gastritis and idiopathic gastritis are shown in table 5. Chronic gastritis without atrophy was seen in 134 out of 294 (46%) patients with *H pylori* associated gastritis compared with 10 out of 24 (42%) of patients with idiopathic gastritis. Grades II and III atrophy were higher in idiopathic gastritis compared with *Helicobacter* associated gastritis.

Table 2 Histological forms of chronic gastritis in 337 patients with ulcer and non-ulcer dyspepsia

Gastritis	No (%) of cases
1 <i>Helicobacter</i> associated	294 (87.2)
2 Idiopathic	24 (7.1)
3 Bile associated reactive	12 (3.6)
4 Drug associated reactive	4 (1.2)
5 Lymphocytic	2 (0.6)
6 Granulomatous	1 (0.3)

Table 3 Topographical patterns of *Helicobacter* associated gastritis in 294 patients with ulcer and non-ulcer dyspepsia

Clinical condition	Total No	Pangastritis	Percentage	Pangastritis predominantly antral	Percentage	Pangastritis predominantly corporal	Percentage	Antral gastritis only	Percentage
Non-ulcer dyspepsia	197	73	37	102	52	4	2	18	9
Duodenal ulcer	79	26	33	44	56	0	0	9	11
Gastric ulcer	18	3	17	13	72	0	0	2	11
Total		102		159		4		29	
Percentage		35		54		1		10	

There were no significant differences between the groups.

Figure 4 The main topographical patterns of *Helicobacter* associated gastritis in relation to the age of patients (decade) in 294 patients with ulcer and non-ulcer dyspepsia.

Figure 5 Atrophic and non-atrophic gastritis in antral biopsy specimens in relation to the age of patients (decade) from 294 subjects with ulcer and non-ulcer dyspepsia.

Table 4 Prevalence and distribution of *H pylori* in antral type and body type gastric mucosa from *Helicobacter* associated gastritis in 294 patients with ulcer and non-ulcer dyspepsia

	Antrum and body equal grades	Antrum and body* predominantly antral	Antrum and body predominantly corporal	Antral only	Corporal only
No of patients	164	92	17	19	2
Percentage	56	31	6	6	1

*The predominance is determined by one-fold difference for *H pylori* grades between antrum and body.

Table 6 Grades of intestinal metaplasia in antral and body mucosa in 294 patients with *H pylori* associated gastritis

Grade	Antrum only	Body only	Antrum and body	Percentage
Grade I	22	2	0	8
Grade II	5	0	0	2
Grade III	1	0	1	1
Total	28*	2†	1	11

p value: *v † p < 0.0001.

Gastric atrophy of all grades was seen in 160 out of 294 (54%) patients with *H pylori* associated gastritis compared with 14 out of 24 (58%) of patients with idiopathic gastritis (not significant). In *H pylori* associated gastritis atrophy was seen in 114 out of 294 (39%) patients in the antrum compared with three out of 294 (1%) in the body (p < 0.0001). Atrophy in both antrum and body was seen in 43 out of 294 (15%) patients and was lower than that seen in the antrum (p < 0.0001) and higher than that seen in the body (p > 0.0001).

Figure 5 shows all types of non-atrophic and atrophic gastritis in relation to the age groups in the antral biopsy specimens of patients with *Helicobacter* associated gastritis. Non-atrophic gastritis was the main pattern in the first four decades of life. Grade I atrophy was the most common type in patients between the third and sixth decades of life. Grades II and III increased with the increased age of the patients.

The distribution of intestinal metaplasia in *Helicobacter* associated gastritis is shown in table 6. Intestinal metaplasia was seen in the antrum only in 28 patients out of 294, in the corpus only in two patients, and in both antrum and corpus in one patient out of 294 with *H pylori* associated gastritis.

Table 7 shows the mean scores of *H pylori*, atrophy, and intestinal metaplasia in non-ulcer dyspepsia, duodenal ulcer, and gastric ulcer. There was no significant difference in the *H pylori* grades in antrum or corpus among all clinical conditions. However, atrophy was more common in both antrum and corpus in patients with gastric ulcer than in patients with duodenal ulcer (p < 0.002, p < 0.015, respectively) and patients with non-ulcer dyspepsia (p < 0.002, p = NS, respectively). Intestinal metaplasia was similarly higher in the antrum in patients with a gastric ulcer than in those with duodenal

Table 5 Grades of atrophy in *Helicobacter* associated gastritis (294) in comparison with idiopathic gastritis (24) in 337 patients with ulcer and non-ulcer dyspepsia

	<i>Helicobacter</i> associated gastritis					<i>Idiopathic</i> gastritis				
	Antrum	Body	Antrum and body	Total	%	Antrum	Body	Antrum and body	Total	%
Grade O				134	46				10	42
Grade I	93	3	30	126	43*	9	0	0	9**	37
Grade II	19	0	11	30	10†	1	1	2	4‡	17
Grade III	2	0	2	4	1‡	0	0	1	1‡‡	4
All grades/site	114	3	43	160	54††	10	1	3	14§	58
Per cent/site	39△	1□	15○			42	4	12		

p < 0.005 ** v ††. p < 0.0001 † v ‡, † v ‡‡, * v ‡, □ v ○, △ v □, △ v ○.

Table 7 Scores of *H pylori*, atrophy, and intestinal metaplasia in three clinical conditions non-ulcer dyspepsia, duodenal ulcer, and gastric ulcer in 294 patients with *H pylori* associated gastritis

Clinical condition	Total No of patients	Mean score of <i>H pylori</i>		Mean score of atrophy		Mean score of intestinal metaplasia	
		Antrum	Body	Antrum	Body	Antrum	Body
Non-ulcer dyspepsia	197	1.53*	1.18†	0.64¶	0.22°	0.13■	0.04□
Duodenal ulcer	79	1.27†	0.88‡	0.61§	0.08‡	0.11∞	0.00
Gastric ulcer	18	1.33††	1.06**	1.22*¶¶	0.28¶	0.33▲	0.00

p < 0.0001: * v †, † v □, ¶ v ◇, § v ♀, ¶¶ v ||; p < 0.0005: ■ v □; p < 0.002: ¶ v *, § v *; p < 0.004: ◇ v ♀; p < 0.015: ♀ v ||; p < 0.02: ■ v ▲, ∞ v ▲; P.N.S.: ▲ v ■, ■ v ♀, ▲ v ♀, † v **, □ v **, ▲ v **, ▲ v ♀, ◇ v ||, ■ v ∞.

(Mean of score is obtained by dividing the total number of scores in each category by the total number of patients).

ulcer (p < 0.02) and non-ulcer dyspepsia (p < 0.02).

Discussion

The biopsy specimens used for this study were collected from patients in developing countries, from four cities in the northern region of the UAE, to give a representative sample of the heterogeneous population currently living there. The study has shown no significant variation in the prevalence of *H pylori* in gastric biopsy specimens among populations of different ethnic origins. The colonisation rates of *H pylori* varied between 87–93% (mean 90%). Most other authors have reported results comparable with those found in this study.^{9–12}

The results have also shown that in duodenal ulceration the gastric mucosa is colonised by *H pylori* in 99% of cases. However, in patients with gastric ulcer the colonisation rate (78%) was significantly lower (p < 0.0001). Dixon reviewed 12 studies from European and other developed and developing countries, and reported a colonisation rate of *H pylori* of 93% for patients with duodenal ulcer and 80% for patients with gastric ulcer.⁷ A prevalence of 100% in patients with duodenal ulcer has been reported in one study from the Ivory Coast.²⁶

H pylori associated gastritis was the most common form of gastritis encountered in this study and accounted for 87% compared with 80% in patients from developed countries.^{27–28} The next largest group was idiopathic gastritis (7.1%) and although no serological tests for *H pylori* were performed in patients of this group, some of these cases may have been due to previous infection with *H pylori*.²⁹ The development of gastric atrophy, which is higher in this group compared with *H pylori* associated gastritis and intestinal metaplasia, may have led to the disappearance of *H pylori* organisms within gastric biopsy specimens from some patients in this group.

Bile associated and drug associated reactive gastritis accounted together for 4.8% of the cases compared with 15% in patients from developed countries.²³ In general, special forms of chronic gastritis accounted for 0.9% of the cases, representing a very small group of chronic gastritis. Lymphocytic gastritis is a newly described entity with one large study reporting an incidence of 1.4%.²⁵ In this study 0.6% of the cases showed evidence of lymphocytic gastritis.

In 93% of the patients *H pylori* organisms were seen in both antrum and body mucosa.

Infection of antral mucosa only by *H pylori* was observed in 6% of the cases while infection of the corporal mucosa only was seen in 1% of the cases. This indicates that *H pylori* is likely to be missed in 6% of cases if the biopsy specimens included body-type gastric mucosa only, and in 1% of cases if the biopsy specimens were taken from the antrum only. Thus taking two biopsy specimens from the antrum and body of the anterior and posterior walls of the stomach²¹ will largely eliminate sampling error for detecting *H pylori*. Antral mucosa is the main target for *H pylori* colonisation. Similar results have been reported by other workers.³⁰

The study has shown that *H pylori* associated gastritis is accompanied by atrophy in 54% of the cases (of these 80% were grade I, 18% grade II, and 2% grade III). This is not surprising, as epidemiological studies from developing countries^{8,12} have suggested that the acquisition of infection by *H pylori* is common at an early age of life, compared with lower incidences in populations from developed countries.⁶ The low percentage of grade III atrophy indicates that progression of atrophy from grade I to grade II and grade III is very small. Ihamaki *et al* found that in 42% of Finnish subjects with specific gastritis, atrophic gastritis developed over three decades.³¹ They found that antral mucosa often regressed histologically whereas superficial body gastritis often progressed towards atrophy.

In another study Sipponen *et al* described three topographic patterns of gastritis with atrophy in the same populations: antral gastritis with atrophy (50%); corporal gastritis with atrophy (17%); and pangastritis with atrophy (33%). Three topographic patterns of *H pylori* associated gastritis with atrophy were encountered in this study: antral gastritis with atrophy (39%); pangastritis with atrophy (15%); and chronic corporal gastritis with mild atrophy, representing only a minor proportion (1%) of the patients in this study. This indicates that most cases of gastritis with atrophy are linked to the acquisition of *H pylori* infection.

In developed countries atrophic body gastritis can be seen in three subtypes. The first is autoimmune associated gastritis which is a rare cause of corporal atrophy³² and is associated with pernicious anaemia.^{33–35} The second type of atrophic corporal gastritis can be seen in some patients with *H pylori* infection and may represent an end stage of *H pylori* associated superficial gastritis.²⁹ The third type may

be seen as a part of severe pangastritis, with atrophy as a consequence of *H pylori* infection.³ The data from this study have shown a very low incidence of corporal gastritis with atrophy associated with *H pylori* infection (1%). None of the patients has shown histological evidence of autoimmune associated corporal gastritis. Thus corporal gastritis with atrophy represents a small part of *Helicobacter* associated gastritis, and occurred alone (1%) or in combination with atrophic antral gastritis (15%).

Correa described three topographic patterns of chronic atrophic gastritis: diffuse corporal, which occurs mainly in northern Europe; diffuse antral, which occurs in urban centres; and multifocal atrophic gastritis, which occurs in northern Europe, China, Japan and the Andes.¹⁸ As the risk of carcinoma of stomach is the lowest in the second pattern (atrophy of the antrum), then it is tempting to suggest that the risk of gastric carcinoma in the population studied would be far less than in the populations of northern Europe, Japan, and China.

In this study intestinal metaplasia occurred in 11% of the cases compared with 25% of cases reported by other workers from developed countries.³⁶ Intestinal metaplasia was of grade I and confined to the antrum in most cases (9.5%).

This study has shown no significant differences in the topographic patterns of *H pylori* associated gastritis in non-ulcer dyspepsia, duodenal ulcer, and gastric ulcer. Differences in the patterns of gastritis in peptic ulcer diseases have been reported by some authors.^{7 37 38} However, the prevalence of glandular atrophy and intestinal metaplasia were higher in patients with gastric ulcer than in those with duodenal ulcer. The findings agree with those reported by other workers.^{7 37}

It is concluded that the prevalence of *H pylori* is higher in the UAE than that reported from developed countries, and that *H pylori* associated gastritis is the predominant type of gastritis. *Helicobacter* associated gastritis with atrophy, although representing 50% of the cases, mainly includes atrophy of grade I and a very small proportion of cases progressing to grade III atrophy. Intestinal metaplasia and gastric atrophy were more common in patients with gastric ulcer than duodenal ulcer.

- Rauws EAJ, Langenberg W, Houthoff HJ, Zanen HC, Tytgat GNJ. Campylobacter pyloridis-associated chronic active antral gastritis. *Gastroenterology* 1988;94:33-40.
- Wyatt JL, Dixon MF. Campylobacter associated chronic gastritis. *Pathol Annu* 1990;25(Pt 1):75-89.
- Sipponen P, Kekki M, Siurala M. The Sydney system: epidemiology and natural history of chronic gastritis. *J Gastroenterol Hepatol* 1991;6:244-51.
- Goodwin CS, Armstrong JA, Marshall BJ. Campylobacter pyloridis, gastritis and peptic ulceration. *J Clin Pathol* 1986;39:353-65.
- Blaser MJ. *Helicobacter pylori* and the pathogenesis of gastro-duodenal inflammation. *J Infect Dis* 1990;161:626-33.
- Graham DY. *Helicobacter pylori*: Its epidemiology and its role in duodenal ulcer disease. *J Gastroenterol Hepatol* 1991;6:105-13.
- Dixon MF. *Helicobacter pylori* and peptic ulceration: Histopathological aspects. *J Gastroenterol Hepatol* 1991;6:125-30.
- Klein PD, Graham DY, Gaillour A, Opekun A, Smith EO. Water source as risk factor for *Helicobacter pylori* infection in Peruvian children. *Lancet* 1991;337:1503-6.
- Marshall BJ, McGeechie DB, Rogers PA, Glancy RJ. Campylobacter pylori infection and gastroduodenal disease. *Med J Aust* 1985;142:439-44.
- Britt DP, Barakat MH, Tunekar MF, Painchaud SM, Adlouni M, Kern K, et al. *Helicobacter pylori* in dyspeptic patients in Kuwait. *J Clin Pathol* 1990;43:987-91.
- El-Guneid A, El-Sherif AM, Murray-Lyon IM, Zureikat N, Shousha S. Effect of chewing Qat on mucosal histology and prevalence of *Helicobacter pylori* in the oesophagus, stomach and duodenum of Yemeni patients. *Histopathology* 1991;19:437-43.
- Al-Moagel MA, Evans DG, Abdul Ghani ME, Adams E, Evans DJ, Malaty HM, et al. Prevalence of *Helicobacter* (formerly Campylobacter) pylori infection in Saudi Arabia: Comparison of those with and without upper gastrointestinal symptoms. *Am J Gastroenterol* 1990;85:944-8.
- Whitehead R, Truelove SC, Gear MWL. The histological diagnosis of chronic gastritis in fibreoptic gastroscope biopsy specimens. *J Clin Pathol* 1972;25:1-11.
- Cheli R, Giacosa A. Chronic atrophic gastritis and mucosal atrophy: One and the same. *Gastrointest Endosc* 1983;29:23-5.
- Strickland RG, Mackay IR. A reappraisal of the nature and significance of chronic atrophic gastritis. *Dig Dis* 1973;18:426-40.
- Glass GB, Pitchumoni CS. Atrophic gastritis. *Hum Pathol* 1975;6:219-50.
- Kekki M, Siurala M, Varis K, Sipponen P, Sistonen P, Nevanlinna RH. Classification principles and genetics of chronic gastritis. *Scand J Gastroenterol* 1987;22(Suppl):1-28.
- Correa P. Chronic gastritis: A clinico-pathological classification. *Am J Gastroenterol* 1988;83:504-9.
- Wyatt JL, Dixon MF. Chronic gastritis: a pathogenic approach. *J Pathol* 1988;154:113-24.
- Stolte M, Heilmann KL. New classification of gastritis. *Leber Magen Darm* 1989;19:220-6.
- Price AB. The Sydney system: Histological division. *J Gastroenterol Hepatol* 1991;6:209-22.
- Zaitoun AM. Use of Romanowsky type (Diff-3) stain for detecting *Helicobacter pylori* in smears and tissue sections. *J Clin Pathol* 1992;45:448-9.
- Sobala GM, King RFG, Axon ATR, Dixon MF. Reflux gastritis in the intact stomach. *J Clin Pathol* 1990;43:303-6.
- Taha AS, Nakshabendi I, Lee FD, Sturrock RD, Russell RI. Chemical gastritis and *Helicobacter pylori* related gastritis in patients receiving non-steroidal anti-inflammatory drugs: Comparison and correlation with peptic ulceration. *J Clin Pathol* 1992;45:135-9.
- Haot J, Hamichi L, Wallez L, Mainguet P. Lymphocytic gastritis: a newly described entity: a retrospective endoscopic and histological study. *Gut* 1988;29:1258-64.
- Biomande MI, Flejou JF, Potet F, Dago-Akribi A, Ouattara D, Kadjo K, et al. Chronic gastritis and *Helicobacter pylori* infection on the Ivory Coast. A series of 277 symptomatic patients. *Gastroenterologie Clinique et Biologique* 1991;15:711-6.
- Blaser MJ. Gastric Campylobacter-like organisms, gastritis and peptic ulcer disease. *Gastroenterology* 1987;93:371-83.
- Yardley JH, Paull G. Campylobacter pylori: A new recognised infectious agent in the gastrointestinal tract. *Am J Surg Pathol* 1988;12(Suppl):89-99.
- Karnes WEJ, Samloff IM, Siurala M, Kekki M, Sipponen P, Kim SWR, et al. Positive serum antibody and negative tissue staining for *Helicobacter pylori* in subjects with atrophic body gastritis. *Gastroenterology* 1991;101:167-74.
- Maaroos HI, Kekki M, Villako K, Sipponen P, Tamm A, Sadeniemi L. The occurrence and extent of *Helicobacter pylori* colonisation and antral and body gastritis profiles in Estonian population sample. *Scand J Gastroenterol* 1990;25:1010-7.
- Ihamaki T, Kekki M, Sipponen P, Siurala M. The sequelae and course of chronic gastritis during a 30 to 34 year biopsic follow-up study. *Scand J Gastroenterol* 1985;20:485-91.
- Strickland RG. Autoimmune gastritis. *J Gastroenterol Hepatol* 1991;6:238-43.
- O'Connor HJ, Axon ATR, Dixon MF. Campylobacter-like organisms unusual in type A (pernicious anaemia) gastritis. *Lancet* 1984;ii:1091.
- Flejou JF, Bahame P, Smith AC, Stockruger RW, Rhode J, Price AB. Pernicious anaemia and Campylobacter-like organisms: is the gastric antrum resistant to colonisation? *Gut* 1989;30:60-4.
- Feng TL, Dooley CP, Dehesa M, Cohen H, Carmel R, Fitzgibbons PL, et al. *Helicobacter pylori* infection in pernicious anaemia: a prospective controlled study. *Gastroenterology* 1991;100:328-32.
- Rugge M, Dimario F, Cassaro M, Baffa R, Farinati F, Rubio J, et al. Pathology of the gastric antrum and body associated with *Helicobacter pylori* infection in non ulcer patients: is the bacterium a promoter of intestinal metaplasia? *Histopathology* 1993;22:9-15.
- Sipponen P, Seppala K, Aarjnen M, Helske T, Kettunen P. Chronic gastritis and gastroduodenal ulcer; a case control study on the risk of coexisting duodenal or gastric ulcer in patients with gastritis. *Gut* 1989;30:922-9.
- Trier JS. Morphology of the gastric mucosa in patients with ulcer diseases. *Dig Dis* 1976;21:138-40.