

Endoscopic Findings and the Prevalence of *Helicobacter Pylori* in Dyspeptic Monks at the Priest Hospital

Somkiat Kasemthamakhun, MD
Pratana Chaiyarerk, RN

Department of Surgery, Priest Hospital, Bangkok, Thailand

Abstract

A descriptive retrospective study was carried out in 312 monks presented to the Priest Hospital with upper abdominal pain and underwent upper gastrointestinal endoscopy and tissue sampling for *H. pylori* by the rapid urease test during March 1, 2008 and February 28, 2011. The average age was 62 + 12.7 years old (range 23-84). The result showed overall 122 infections (39.1%) with age over 50 year in 104 cases (40.9%). The endoscopic findings were gastritis (43.3%), gastric ulcer (10.9%), gastric cancer (3.5%), gastroduodenal ulcer (1.6%), duodenal ulcer (5.8%), duodenitis (7.7%) and non-ulcer dyspepsia (27.2%). Only the findings of gastric ulcer, gastric cancer, peptic ulcer and duodenal ulcer were significantly associated with *H. pylori* infection ($p < 0.05$).

Keywords: dyspepsia, *Helicobacter pylori*, monks, peptic ulcer

INTRODUCTION

Dyspepsia is a distressing symptom of upper abdomen^{1,2} such as pain, burning, early satiety, postprandial fullness, bloating, belching, anorexia and nausea³⁻⁵. It is a problem in 20-40%⁶ of normal population and one of chief presenting symptoms to the physician in 20-25%^{7,8}. Among possible intra-abdominal viscera, stomach, esophagus, and biliary system including gall bladder are the most culprits.^{9,10} Generally dyspepsia is chronically off and on and after history and physical exam taken, patients are usually prescribed symptom-relief medications for a certain period unless alarming features^{4,5,11} occur like ages older than 45 years, suspected gastrointestinal bleeding,

unexplained iron deficiency anemia, unintentional weight loss more than 10%, swallowing with difficulty or pain, frequent bouts of vomiting together with malignancy manifestations such as palpable abdominal mass, lymphadenopathy, jaundice and rectal shelf. Further standard investigation for these patients is upper gastrointestinal endoscopy¹² because of high reliance and safety.^{3,13,14} Most of endoscopic diagnoses range from normal mucosa, gastritis, gastric ulcers or duodenal ulcer, in addition to tissue sampling for *Helicobacter Pylori*. In Thailand, the best current method to identify the microorganism is by means of the rapid urease test¹⁵. The negative test result will need tissue sampling for pathological diagnosis while the positive

will not, except in case of ulcer that malignancy has to be ruled out by tissue study of the ulcer rim.

H. pylori is a spring-shaped, gram negative bacteria viable on gastric mucosa by producing urease enzyme which catalyzes urea into ammonia and carbon dioxide so that the bacteria could flourish amidst the hostile environment of strong acidity. Research revealed the oral route of bacterial transmission since childhood. Personal hygiene can lessen contact and transmission as evidenced in developing countries where poor sanitation entails in food and water contamination^{12,16}. *H. pylori* is taken for granted to play a major role in gastritis and one of the important causative factors of peptic ulcers and a risk to gastric cancer¹⁷. The Priest Hospital under Medical Division of the Public Health of Thailand has been serving 267,939 monks¹⁸ throughout the country for free. Due to the monastic life and religious practices like early morning receiving alms round, once or twice eating unrequested food offered before mid-day, it is interesting to study any association between ages, endoscopic findings in dyspeptic monks and *H. pylori* infection to contribute to the further care plan.

MATERIALS AND METHODS

A retrospective study was carried out in 312 dyspeptic monks who received care at the Priest Hospital during March 1, 2008 and February 28, 2011. After approval by the Ethics Committee of the Priest Hospital, their medical records were reviewed including endoscopic data from Surgery Department. Exclusion criteria were upper gastrointestinal bleeding, chronic liver disease, anticoagulant use and advanced cancer. Each patient underwent tissue sampling from gastric antrum and body via the endoscopy for the study of *H. pylori* by rapid urease test and in the case of negative

result, the tissue was sent for confirmation by pathological exam. Data in terms of ages, endoscopic findings grouped into normal mucosa, gastritis, gastric ulcer, etc were compared with test results of *H. pylori* and analyzed by both descriptive statistics for demographic data and analytic statistics, Chi-square, for any association between independent and dependent variables with $p < 0.05$ as the statistical significance.

RESULTS

The data of 312 dyspeptic monks who underwent physical exam, upper gastrointestinal endoscopy and tissue sampling for the rapid urease test showed an average age of 62 ± 12.7 years old (range 23-84), 122 infections of *H. pylori* (39.1%). Fifty eight monks under 50 years old (18.6%) of which 18 (31%) were infected while those 254 monks over 50 years old (81.4%) had 104 infected cases or 40.9%. No association was found between ages and *H. pylori* infection ($p = 0.163$; Table 1).

Upper gastrointestinal endoscopic findings were as follows: 135 gastritis (43.3%) of which 42 were infected with *H. pylori* (31.1%), 34 gastric ulcers (10.9%) of which 26 were infected (76.5%), 11 gastric cancer (3.5%) of which 8 were infected (72.7%), 5 gastroduodenal ulcers (1.6%) of which all were infected (100%), 18 duodenal ulcers (5.8%) of which 16 were infected (88.9%), 24 duodenitis (7.7%) of which 10 were infected (41.7%) and 85 non-ulcer dyspepsia (27.2%) of which 15 were infected (17.6%). The upper gastroscopic findings were significantly associated with *H. pylori* infection only in terms of gastric ulcer, gastric cancer, gastroduodenal ulcer and duodenal ulcers ($p < 0.05$; Table 2).

Table 1 Association between ages and *H. pylori* infection

Age (years)	Cases (%)	H. pylori positive	pylori negative	p
		n = 122 (%)	n = 190 (%)	
< 50	58 (18.6)	18 (31.0)	40 (69.0)	0.163
> 50	254 (81.4)	104 (40.9)	150 (59.1)	

Table 2 The associations between endoscopic findings and *H. pylori* infection

n = 312	cases (%)	presence	H. pylori positive (%)	H. pylori negative (%)	p
Gastritis	135 (43.3)	yes	42 (31.1)	93 (68.9)	0.126
		no	80 (45.2)	97 (54.8)	
Gastric ulcer	34 (10.9)	yes	26 (76.5)	8 (23.5)	0.004*
		no	96 (34.5)	182 (65.5)	
Gastric cancer	11 (3.5)	yes	8 (72.7)	3 (27.3)	0.02*
		no	114 (37.9)	187 (62.1)	
Gastroduodenal ulcer	5 (100)	-			0.009*
		no	117 (38.1)	190 (61.9)	
Duodenal ulcer	18 (5.8)	yes	16 (88.9)	2 (11.1)	0.001*
		no	106 (36.1)	188 (63.9)	
Duodenitis	24 (7.7)	yes	10 (41.7)	14 (58.3)	0.789
		no	112 (38.9)	176 (61.1)	
Normal	85 (27.2)	yes	15 (17.6)	70 (82.4)	0.068
		no	107 (47.1)	120 (52.9)	

*statistical significance

DISCUSSION

The incidence of *H. pylori* infection in this study was 39.1%, less than those reported by A Kiatpanapikul et al¹⁹ and U Kachinthorn et al²⁰. Monk activities, monastic life, abbot environment and one or twice eating before mid-day are explanatory. Foods offered for alms round, even unrequested for, are usually selected to be clean and well-done since Thai people traditionally believe in good deeds from putting good food into alms bowl. Besides, monks at large have foods with a serving spoon or eat from their own personal bowls, lowering risk to infection. Similarly to the studies of A Kiatpanapikul et al¹⁹ and P Choomsri et al,²¹ ages are not significantly associated with *H. pylori* infection. From the age of 50 the incidence is increasing, according to the previous reports that the microorganism is found in all ages, but trended upwards in the elderly, probably due to longer exposure especially in developing countries^{12,22}.

Our upper gastrointestinal endoscopic findings, not different from previous reports,^{23,24} were 43.3% gastritis, 10.9% gastric ulcer, 3.5% gastric cancer, 1.6% gastroduodenal ulcer, 5.8% duodenitis with *H. pylori* infection 31.1, 76.5, 72.7, 100, 8.9 and 41.7% respectively. In Thailand U Kachinthorn et al²⁰ studied

the incidence of the infection to be 45% in either normal or gastritis, 55% in gastric ulcer and 60% in duodenal ulcer while abroad it was reported as high as 80-90%¹⁰⁻¹² in peptic ulcer and gastritis patients. From the study of A Kiatpanapikul et al,¹⁹ the incidence of *H. pylori* was 66.5% in gastritis, 77.5% in gastric ulcer and 90.2% in duodenal ulcer. For non-ulcer dyspepsia, it was 17.6% approximately the same that was reported to be 10-20% by Zagari RM et al.⁹ McCarthy C et al²⁵ found almost all patients with non-ulcer dyspepsia infected with *H. pylori* got better after treatment and supported by other studies to be cost-effective²⁶ which could be long term management to reduce risk of gastric ulcer, gastritis or gastric cancer in the future^{27,28}. There was a report of 6 - folded increase in cancer for patients infected with this microorganism over 10 years²⁹.

The association between our endoscopic findings in the upper gastrointestinal tract and *H. pylori* infection revealed a statistical significance to be the only involved factor in monks with gastric ulcer, gastric cancer, gastroduodenal ulcer and duodenal ulcer ($p < 0.05$).

CONCLUSION

Monks presenting with dyspepsia, undergoing upper gastrointestinal endoscopy and tissue sampling

were found to have *H. pylori* infection incidence of 39.1% which arose with age over 50 years. Findings of only gastric ulcer, gastric cancer, gastroduodenal ulcer and duodenal ulcer were significantly associated with the infection. The knowledge has contributed to the care plan for monks in Thailand.

REFERENCES

1. Rabeneck L, Nelda PW, Graham DY. Managing dyspepsia: what do we know and what do we need to know?. *Am J Gastroenterol* 1998;93:920-4.
2. Holtmann G, Stanghellini V, Talley N. Nomenclature of dyspepsia, dyspepsia subgroups and functional dyspepsia: clarifying the concepts. *Baillieres Clin Gastroenterol* 1998;12:417-33.
3. Talley NJ, Silverstein MD, Agreus L, Nyren O, Sonnenberg A, Holtmann G. AGA technical review: evaluation of dyspepsia. *Gastroenterology* 1998;114:582-95.
4. Ikenberry SO, Harrison ME, Lichtenstein D, Dominitz JA, Anderson MA, Jagannath SB, et al. The role of endoscopy in dyspepsia. *Gastrointest Endosc* 2007;66:1071-5.
5. Talley NJ, Vakil N. Practice Parameters Committee of the American College of Gastroenterology. Guidelines for the management of dyspepsia. *Am J Gastroenterol* 2005;100:2324-37.
6. Grainger SL, Klass HJ, Rake MO. Prevalence of dyspepsia: the epidemiology of overlapping symptoms. *Postgrad Med J* 1994;70:154-61.
7. Tebaldi M, Heading RC. Clinical economics review: functional (non-ulcer) dyspepsia. *Aliment Pharmacol Ther* 1998;12:11-9.
8. Talley NJ, Weaver AL, Zinsmeister AR. Impact of functional dyspepsia on quality of life. *Dig Dis Sci* 1995;40:584-9.
9. Zagari RM, Fuccio L, Bazzoli F. Investigating dyspepsia. *BMJ* 2008;337:1400.
10. Dempsey DT, Stomach. In: Brunicaardi FC, editor. *Schwartz's principle of surgery*. 8th ed. New York: McGraw-Hill; 2005.
11. Talley NJ, American gastroenterological association. American gastroenterological association medical position statement: evaluation of dyspepsia. *Gastroenterology* 2005;129:1753-5.
12. Talley NJ, Vakil NB, Moayyedi P. American gastroenterological association technical review on the evaluation of dyspepsia. *Gastroenterology* 2005;129:1756-80.
13. Hart R, Classen M. Complications of diagnostic gastrointestinal endoscopy. *Endoscopy* 1990;22:229-33.
14. Daneshmend TK, Bell GD, Logan RF. Sedation for upper gastrointestinal endoscopy: results of a nationwide survey. *Gut* 1991;32:12-15.
15. Research group of peptic ulcer. Guidelines for diagnosis and treatment for *Helicobacter pylori* infection: Guidelines for diagnosis and treatment for dyspeptic patients and *Helicobacter pylori* infection in Thailand. 1st ed. Bangkok : The Gastroenterological Association of Thailand; 1999.
16. Rowland M, Kumar D, Daly L, O'connor P, Vaughan D, Drumm B. Low rates of *Helicobacter pylori* reinfection in children. *Gastroenterology* 1999;117:336-41.
17. Clark CJ, Thirlby RC, Picozzi V Jr, Schembre DB, Cummings FP, Lin E. Current problems in surgery : gastric cancer. *Curr Probl Surg* 2006;43:566-670.
18. Office of National Buddhism. Statistics the number of monks since 2009 -2010.2011 (cited 2012 March 28). Available from: <http://www.onap.go.th/Data Monk/>
19. Kiatpanabhikul A, Yodavudh S. Association of gastroscopic features with *Helicobacter Pylori* infection in dyspeptic patients at Charoenkrungpracharak Hospital. *J C H* 2008;4,12-19.
20. Kanchintorn U, Lueugrojanakul P, Atisook K, Theerabuttra C, Tauwandee T, Beenyapisit S. *Helicobacter pylori* and peptic ulcer diseases: prevalence and association with antral gastritis in 210 patients. *J Med Assoc Thai* 1992;75:386-92.
21. Choomsri P, Bumpenboon W, Wasuthit Y, Euanorasetr C, Sumritpradit P, Suwanthunma W, Lertsithichai P. Upper gastrointestinal endoscopy findings in patients presenting with dyspepsia. *Thai J Surg* 2010;31:7-12.
22. Talley NJ, Hunt RH. What role does *Helicobacter pylori* play in dyspepsia and nonulcer dyspepsia? Arguments for and against *H. pylori* being associated with dyspeptic symptoms. *Gastroenterology* 1997;113:S67-S77.
23. Tytgat G N J. Role of endoscopy and biopsy in the work up of dyspepsia. *Gut* 2002;50:13-16.
24. The role of endoscopy in dyspepsia. *Gastrointestinal Endoscopy* 2007;66:1071-5.
25. McCarthy C, Patchett S, Collins RM, Beattie S, Keane C, O'Morain C. Long-term prospective study of *Helicobacter pylori* in non-ulcer dyspepsia. *Dig Dia Sci* 1995;40:114-9.
26. Moayyedi P, Soo S, Deeks J, Delaney B, Harris A, Innes M, et al. Eradication of *Helicobacter pylori* for non-ulcer dyspepsia. *Cochrane Database Syst Rev* 2006;2:CD002096.
27. Spiegel BM, Vakil NB, Ofman JJ. Dyspepsia management in primary care: a decision analysis of competing strategies. *Gastroenterology* 2002;122:1270-85.
28. Malfertheiner P, Sipponen P, Naumann M, Moayyedi P, Megraud F, Xiao SD, et al. *Helicobacter pylori* eradication has the potential to prevent gastric cancer: a state-of-art critique. *Am J Gastroenterol* 2005;100:2100-15.
29. *Helicobacter and cancer collaborative group*. Gastric cancer and *Helicobacter pylori*: a combined analysis of 12 case control studies nested within prospective cohorts. *Gut* 2001;49:347-53.