

ORIGINAL ARTICLE

Prevalence of Helicobacter pylori in Children

Praveen Saraswat

Assistant Professor, Department of Paediatrics, Hind Institute of Medical Sciences, Barabanki, UP, India

ABSTRACT:

Background: The study was conducted to assess the prevalence of helicobacter pylori in children in a known population. **Material and methods:** A total of one hundred children were checked for gastrointestinal problems. Individuals under the age of one year, those above the age of fifteen, those with chronic illnesses of any kind, and those taking gastrointestinal-related medications were not included in the study. Questions concerning gastrointestinal symptoms and complaints, such as persistent stomach discomfort, nausea, vomiting, gastroesophageal reflux, and dyspeptic complaints, were asked of all the children. Patients were considered to be positive for H. pylori infection if they had both positive serology results and positive stool antigen test results. **Results:** This study comprised of 50 male subjects as well as 50 female subjects. H. pylori infection was present in 9 subjects out of 100 subjects. It was observed that out of 9 subjects having H. pylori infection, 6 subjects reported with recurrent abdominal pain, 2 subjects presented with nausea and vomiting and 1 subject presented with haematemesis. **Conclusion:** 9 out of 100 subjects had H. pylori infection. The most common clinical feature was recurrent abdominal pain.

Keywords: H. pylori, bacteria, children, abdominal pain, haematemesis

Corresponding author: Praveen Saraswat, Assistant Professor, Department of Paediatrics, Hind Institute of Medical Sciences, Barabanki, UP, India

This article may be cited as: Saraswat P. Prevalence of Helicobacter pylori in Children. J Adv Med Dent Scie Res 2014;2(3):400-402.

INTRODUCTION

Helicobacter pylori is a spiral gram-negative microorganism that is distributed worldwide. It is estimated that over 50% of the world population are infected with H. pylori.¹⁻³ H. pylori-associated infection is either usually clinically silent or its signs and symptoms are non-specific. Gastroesophageal reflux, esophagitis, delayed gastric emptying, and various motility disorders can be a sign or symptom of it.^{4,5} However, these symptoms are seen in many childhood illnesses. Younger children with peptic complaints may not have symptoms as clear as those of older children, and diagnosis of infection due to H. pylori is more difficult.⁶⁻⁸

The epidemiology of H. pylori-associated infection is variable, since the prevalence is significantly higher and infection occurs in earlier ages in developing or poor countries compared to developed countries.⁹⁻¹¹

MATERIAL AND METHODS

A total of one hundred children were checked for gastrointestinal problems. Individuals under the age of one year, those above the age of fifteen, those with chronic illnesses of any kind, and those taking gastrointestinal-related medications were not included in the study. Questions concerning gastrointestinal symptoms and complaints, such as persistent stomach discomfort, nausea, vomiting, gastroesophageal reflux, and dyspeptic complaints, were asked of all the children. Patients were considered to be positive for H. pylori infection if they had both positive serology results and positive stool antigen test results. All children had stool samples and blood sera examined. All children had their venous blood drawn in three

milliliters, and the sera were kept in a deep freezer at -20°C.

RESULTS

Table 1: Gender-wise distribution of subjects.

Gender	Number of subjects	Percentage
Males	50	50%
Females	50	50%
Total	100	100%

There were 50 males and 50 females in this study.

Table 2: Prevalence of Helicobacter pylori infection in children

Prevalence	Number of subjects	Percentage
Absent	91	91%
Present	09	09%
Total	100	100%

H. pylori infection was present in 9 subjects out of 100 subjects. It was observed that out of 9 subjects having H. pylori infection, 6 subjects reported with recurrent abdominal pain, 2 subjects presented with nausea and vomiting and 1 subject presented with haematemesis.

DISCUSSION

In 1983, Robin Warren, a pathologist in Perth, reported the presence of "curved bacterium" in the mucosal layer of the gastric biopsy specimen. Together with Barry Marshall, he subsequently isolated the organism from the gastric biopsy specimens and named it Campylobacter pyloridis (C. pylori),¹² which was ultimately named as Helicobacter pylori (H. pylori). Marshall and Warren also noted

that *C. pylori* (*H. pylori*) infection was associated with gastric and duodenal ulceration.¹³ In recognition with this imperative discovery, they were awarded the Nobel Prize for Medicine in 2005.

Their discovery initiated a new interest in previously neglected field of gastric microbiology. In 1994, the National Institute of Health consensus conference in USA declared an association between *H. pylori* and peptic ulcer disease.¹⁴ During the same year, *H. pylori* was identified as a carcinogen associated with gastric adenocarcinoma¹⁵ and gastric non-Hodgkin lymphoma.¹⁶ An association between *H. pylori* and gastric mucosa-associated lymphoid tissue lymphoma (MALToma) was identified in 1991.¹⁷

This study was conducted to assess the prevalence of *Helicobacter pylori* in children.

This study comprised of 50 male subjects as well as 50 female subjects. *H. pylori* infection was present in 9 subjects out of 100 subjects. It was observed that out of 9 subjects having *H. pylori* infection, 6 subjects reported with recurrent abdominal pain, 2 subjects presented with nausea and vomiting and 1 subject presented with haematemesis. Singh M et al (2006)¹⁸ determined prospectively the prevalence of *H. pylori* infection in children and its association with gastroduodenal disease. They evaluated 240 children undergoing upper gastrointestinal endoscopy for *H. pylori* infection by rapid urease test, culture, ureA PCR and histopathology. Group I constituted 58 children with upper abdominal pain (UAP) and group II (controls) of 182 children without UAP who underwent diagnostic or therapeutic endoscopy for other reasons. *Helicobacter pylori*-positive children with UAP received anti-*H. pylori* therapy. *Helicobacter pylori* infection was significantly higher in children with UAP than controls (53.4% vs. 28%; $P < 0.001$) and overall prevalence increased with age. On follow-up endoscopy, *H. pylori* had been eradicated from 82% of children with UAP; it was eradicated from the remaining 18% after a second regimen. Treated *H. pylori*-positive children with UAP remained symptom-free for a median of 25 months. Control children remained chronically *H. pylori* infected. Chronic inflammation was present in all infected children, and active inflammation in 48.8%. The study shows *H. pylori* infection increases with age and is strongly linked to UAP in children. Ceylan A et al (2007)¹⁹ in their study determined the prevalence of *Helicobacter pylori* among children and their family members and to evaluate some epidemiologic characteristics. The study included 275 children, aged 1-15 year(s), suffering from different gastrointestinal complaints. Blood serology and stool antigen testing were used for the diagnosis of infection due to *H. pylori*. Sixty-five (23.6%) of the 275 children were positive for *H. pylori*, and this positivity had a significantly increasing correlation with age ($p < 0.001$). *H. pylori*-associated infection was observed among 45 (69.2%) and 17 (8%) mothers in the *H. pylori*-infected and

non-infected groups respectively ($p < 0.0001$). Most children and their families infected with *H. pylori* were living in an urban area. The findings suggest that infection due to *H. pylori* is a problem for this district area, and all children having any gastrointestinal complaints should be examined whether *H. pylori* was prevalent among their family members.

CONCLUSION

9 out of 100 subjects had *H. pylori* infection. The most common clinical feature was recurrent abdominal pain.

REFERENCES

- Herbst JJ. Ulcer disease. In: Behrman RE, Kliegman RM, Jenson HB, editors. Nelson Textbook of pediatrics. 16th ed. Philadelphia: Saunders; 2000. pp. 1147–8.
- Marshall B. The 1995 Albert Lasker Medical Research Award. *Helicobacter pylori*. The etiologic agent for peptic ulcer. *JAMA*. 1995;274:1064–6.
- Kutukculer N, Aydogdu S, Caglayan S, Yagci RV. Salivary and gastric fluid secretory immunoglobulin A and free secretory component concentrations in children with *Helicobacter pylori*-positive gastritis. *J Trop Pediatr*. 1998;44:178–80.
- Glassman MS. *Helicobacter pylori* infection in children. A clinical overview. *Clin Pediatr (Phila)* 1992;31:481–7.
- Go MF, Crowe SE. Virulence and pathogenicity of *Helicobacter pylori*. *Gastroenterol Clin North Am*. 2000;29:649–70.
- Ozturk H, Senocak ME, Uzunalimoglu B, Hascelik G, Büyükpamukcu N, Hicsönmez A. *Helicobacter pylori* infection in symptomatic and asymptomatic children: a prospective clinical study. *Eur J Pediatr Surg*. 1996;6:265–9.
- Us D, Hascelik G. Seroprevalence of *Helicobacter pylori* infection in an asymptomatic Turkish population. *J Infect*. 1998;37:148–50.
- Meyer B, Werth B, Beglinger C, Dill S, Drewe J, Vischer WA, et al. *Helicobacter pylori* infection in healthy people: a dynamic process? *Gut*. 1991;32:347–50.
- Biselli R, Fortini M, Matricardi PM, Stroffolini T, D'Ameli R. Incidence of *Helicobacter pylori* infection in a cohort of Italian military students. *Infection*. 1999;27:187–91.
- Blecker U, Lanciers S, Hauser B, Vandenplas Y. The prevalence of *Helicobacter pylori* positivity in a symptom-free population, aged 1 to 40 years. *J Clin Epidemiol*. 1994;47:1095–8.
- Graham DY, Adam E, Reddy GT, Agarwal JP, Agarwal R, Evans DJ, Jr., et al. Seroepidemiology of *Helicobacter pylori* infection in India. Comparison of developing and developed countries. *Dig Dis Sci*. 1991;36:1084–8.
- Warren JR, Marshall BJ. Unidentified curved bacilli on gastric epithelium in active chronic gastritis. *Lancet*. 1983;1:1273–5.
- Marshall BJ, Warren JR. Unidentified curved bacilli in the stomach of patient with gastritis and peptic ulceration. *Lancet*. 1984;1:1311–5.
- NIH consensus conference. *Helicobacter pylori* in peptic ulcer disease. NIH consensus development panel

- on *Helicobacter pylori* in peptic ulcer disease. *JAMA*. 1994;272:65–9.
15. World Health Organization/International Agency of Research on Cancer. Schistosomes, Liver flukes and *Helicobacter pylori*. Lyon: Lyon international agency for research on cancer; 1994. Infection with *Helicobacter pylori*; pp. 177–244.
 16. Personnet J, Hansen H, Rodriguez L, Gelb AB, Warnke RA, Jellum E, et al. *Helicobacter pylori* infection and gastric lymphoma. *N Engl J Med*. 1994;330:1267–71.
 17. Wotherspoon AC, Ortiz-Hidalgo C, Falzon MR, Issacson PG. *Helicobacter* associated gastritis and primary gastric B cell lymphoma. *Lancet*. 1991;338:1175–6.
 18. Singh M, Prasad KN, Yachha SK, Saxena A, Krishnani N. *Helicobacter pylori* infection in children: prevalence, diagnosis and treatment outcome. *Trans R Soc Trop Med Hyg*. 2006 Mar;100(3):227-33.
 19. Ceylan A, Kirimi E, Tuncer O, Türkdöğän K, Ariyuca S, Ceylan N. Prevalence of *Helicobacter pylori* in children and their family members in a district in Turkey. *J Health PopulNutr*. 2007 Dec;25(4):422-7. PMID: 18402185; PMCID: PMC2754017.