

Relationship between Disease Activity and *Helicobacter pylori* Infection in Patients with Vitiligo

Sir,
Vitiligo is a common dermatologic disease with a debated etiology. Infectious agents have been claimed to trigger disease development through antibody production, molecular mimicry, cytokine production, major histocompatibility complex activation, regulatory T-cell dysfunction, immune complex formation, and chronic inflammatory damage.^[1]

Helicobacter pylori (*H. pylori*) is one of the most common pathogens affecting humans, infecting approximately 50% of the world's population.^[2] The prevalence of infection varies in different governorates in Egypt. Reported prevalence in Delta is approximately 70%.^[3] Several immune-mediated skin disorders have been considered to have a relationship with *H. pylori* infection.

The aim of the present study was to evaluate if there is a relationship between active vitiligo and *H. pylori* infection.

Seventy-five patients with non segmental vitiligo were selected [51 (68%) cases with active disease and 24 (32%) with stable disease]. Seventy-five healthy, age and gender- matched subjects were enrolled as a control group. A written consent form approved by local Ethical Research Committee was obtained from every participant prior to study initiation.

The diagnosis of vitiligo was made on the basis of typical clinical features and Wood’s light examination. All studied cases were either newly diagnosed or old patients with completely depigmented lesions. Assessment of disease activity was done according to the Vitiligo Disease Activity (VIDA) score.^[4] Clinical data of selected cases and controls are summarized in Table 1.

Patients with dermatological diseases other than vitiligo, with autoimmune diseases or malignancy, those who took previous *H. Pylori* eradication treatment, and those using NSAIDs, antibiotics, or proton pump inhibitors in the last month before the study were excluded.

Diagnosis of infection was done by stool antigen test (SAT) which is a noninvasive method with good sensitivity and specificity, in global meta-analysis, in the diagnosis of *H. Pylori* infection.^[5] Measurement of serum IgG antibody was done to avoid the influence of different factors on SAT positivity.^[6]

H. pylori specific IgG antibody in serum and SAT were detected by enzyme-linked immunosorbent assay (ELISA). As the presence of antibody may denote active or past infection, included cases with both positive SAT and IgG were designated as having active infection.

For SAT, level <20 ng/ml denotes negative result and level >20 ng/ml denotes positive result.^[7]

For *H. pylori* IgG in serum, level <20 u/ml denotes negative result and level >20 u/ml denotes positive result.^[8]

Data were collected, tabulated, and statistically analyzed using a personal computer with SPSS version 11 software (SPSS Inc, Chicago, Illinois, USA). Fisher’s exact test was used for comparison of qualitative variables in 2 × 2 tables when expected cell count of more than 25% of cases was less than 5. Chi-square test (χ^2) was used to study the association between two qualitative normally distributed variables. Mann–Whitney U test was used for comparison between two groups not normally distributed having quantitative variables. Differences were considered statistically significant with $P < 0.05$.

H. pylori infection was positive in 49 (65.3%) vitiligo cases compared with 18 (24%) in the control group ($P = 0.001$) [Figure 1]. In cases, infection was significantly associated

with high VIDA score ($P = 0.009$) and with active disease ($P = 0.02$) [Table 2].

H. pylori may alter cellular immunity resulting in loss of self-tolerance and autoimmune destruction of melanocytes. This autoimmune reaction results from higher levels of C-reactive protein, chronic inflammatory condition, autoantibody production, and antigenic stimulation.^[9] In addition, local inflammation of gastric mucosa leads to release of inflammatory cytokines [tumor necrosis factor (TNF)- α , interleukin (IL)-6, IL-8, and IL-1 β] that produce systemic inflammation which subsequently affect skin microenvironment with loss of functional melanocytes.^[10]

The current study demonstrated significant association between active vitiligo and *H. pylori* infection. On the contrary, Dogan *et al.* reported absence of association between *H. pylori* positivity and vitiligo activity score.^[11] Çakmak *et al.* showed no significant relationship between *H. Pylori* SAT positivity and activity of vitiligo.^[12]

In the current work, positive cases for *H. Pylori* infection received eradication therapy consisting of omeprazole 20 mg BID, clarithromycin 500 mg BID, and metronidazole 500 mg BID for 14 days.^[13]

Follow-up 3 months later showed suppression of disease activity in (11/49) cases.

Non responding cases regarding disease activity may be explained by the multifactorial etiology of vitiligo.

Table 1: Clinical data of selected cases and control subjects

Variable	Cases (n=75)	Controls (n=75)	Test and P
Gender			U=1.15, P=0.2
Males	34 (45.3%)	9 (36%)	
Females	41 (54.7%)	16 (64%)	
Age (years)			$\chi^2=0.66$, P=0.4
X±SD	30.1±16.5	33.5±14.6	
Range	11-64	13-57	
Disease duration (months)			
X±SD	37.8±41.6		
Range	1-180		
Lesion extent (%)			
X±SD	27.1±25.2		
Range	5-90		
VIDA Score			
X±SD	2.37±1.61		
Range	-1-4		
Activity			
Active	51 (68%)		
Inactive	24 (32%)		

U: Mann-Whitney test, χ^2 : Chi-square test, X±SD: Mean=standard deviation, VIDA: Vitiligo Disease Activity

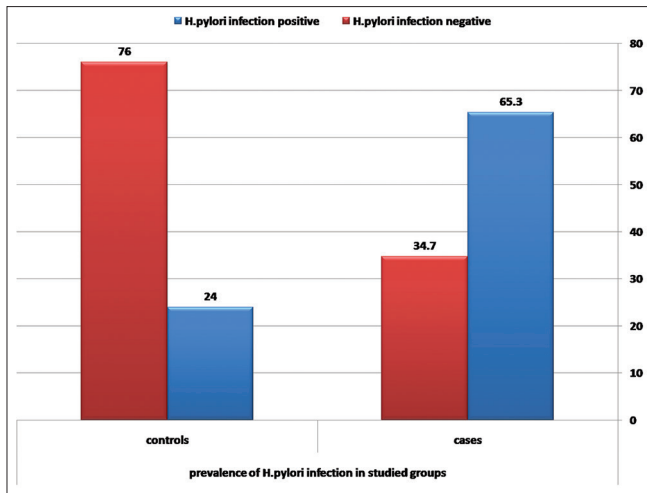


Figure 1: Prevalence of *H. pylori* infection in studied cases and controls

Table 2: Association between *H. pylori* infection and disease activity

Variables	<i>H. pylori</i> infection in studied cases		Test and P
	Positive (n=49)	Negative (n=26)	
VIDA Score			$\chi^2=2.6,$
X±SD	3.03±1.34	2.02±1.63	P=0.009*
Disease activity			$\chi^2=5.4,$
Active	41 (83.6%)	15 (57%)	P=0.02*
Inactive	8 (16.4%)	11 (43%)	

*Significant, χ^2 : Chi-square test, X±SD: Mean±standard deviation, VIDA: Vitiligo Disease Activity

Therefore, it may be useful to search for *H. Pylori* infection in patients with vitiligo, especially in those with active disease. Clinical trials to evaluate the effect of *H. Pylori* eradication in infected vitiligo patients are also needed.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

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References

- Kivity S, Arango MT, Ehrenfeld M, Tehori O, Shoenfeld Y, Anaya JM, *et al.* Infection and autoimmunity in Sjogren's syndrome: A clinical study and comprehensive review. *J Autoimmun* 2014;14:53-5.
- Windle H, Kelleher D. Identification and characterization of meyalloprotease activity from *H. Pylori*, which is capable of degrading immunoglobulins (abstract). In: 8th international workshop on gastroduodenal pathology and *H. Pylori*. Edinbrugh Scotland 1995; 340:43-49.
- Sabah AA, Gneidy MR, Saleh NM. Prevalence of *Helicobacter pylori* infection among adult patients with different gastrointestinal parasites in Tanta City. *J Egypt Soc Parasitol* 2015;45:101-6.
- Shankar DS, Shashikala K, Madala R. Clinical patterns of vitiligo and its associated co morbidities: A prospective controlled cross-sectional study in South India. *Indian Dermatol Online J* 2012;3:114-8.
- Gisbert JP, de la Morena F, Abaira V. Accuracy of monoclonal stool antigen test for the diagnosis of *H. pylori* infection: A systematic review and meta-analysis. *Am J Gastroenterol* 2006;101:1921-30.
- Chey W, Wong B. American College of Gastroenterology guideline on the management of *Helicobacter pylori* infection. *Am J Gastroenterol* 2007;102:1808-25.
- Cutler AF. Testing for *Helicobacter pylori* in clinical practice. *Am J Med* 1996;100:35-41.
- Blaser MJ. *Helicobacter pylori* and associated diseases. *BMJ* 1998;316:1507-10.
- Gasbarrini A, Franceschi F, Gasbarrini G, Pola P. Extraintestinal pathology associated with *Helicobacter* infection. *Eur J Gastroenterol Hepatol* 1997;9:231-3.
- Moss S, Calm J. *H. pylori* and peptic ulcer: The present position. *Gut* 1994;33:289.
- Doğan Z, Özdemir P, Ekşioğlu M, Filik L. Relationship between *Helicobacter pylori* infection and vitiligo: A prospective study. *Am J Clin Dermatol* 2014;15:457-62.
- Çakmak SK, Tantoglu BH, Onan D. The frequency of *Helicobacter pylori* infection in vitiligo patients. *Pigment Int* 2015; 2:81-4.
- Malferttheiner P, Mégraud F, O'Morain C, Bell D, Bianchi Porro G, Deltenre M, *et al.* Current European concepts in the management of *Helicobacter pylori* infection - The Maastricht Consensus Report. The European *Helicobacter Pylori* Study Group (EHPSG). *Eur J Gastroenterol Hepatol* 1997;9:1-2.

Access this article online

Website:
www.idoj.in

DOI:
10.4103/idoj.IDOJ_77_17

Quick Response Code



How to cite this article: Bakry OA, Basha M, El Hefnawy S, Mekkawy S. Relationship between disease activity and *Helicobacter pylori* infection in patients with vitiligo. *Indian Dermatol Online J* 2018;9:59-61.

Received: April, 2017. **Accepted:** May, 2017.

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