



Perspective

Can Natural Products Suppress Resistant *Helicobacter pylori* to Fight Against Gastric Diseases in Humans?

Li-Shu Wang¹, Carla Elena Echeveste¹, Jianhua Yu², Yi-Wen Huang³, John Lechner¹, Ling Mei⁴, Patrick Sanvanson⁴, Martha Yearsley⁵, Chin-Kun Wang⁶, Gary Stoner^{7,*}

¹Division of Hematology and Oncology, Department of Medicine, Medical College of Wisconsin, Milwaukee, WI, USA

²Department of Hematology and Hematopoietic Cell Transplantation, City of Hope National Medical Center and Beckman Research Institute, Duarte, CA, USA

³Department of Obstetrics and Gynecology, Medical College of Wisconsin, Milwaukee, WI, USA

⁴Division of Gastroenterology and Hepatology, Department of Medicine, Medical College of Wisconsin, Milwaukee, WI, USA

⁵Department of Pathology, Ohio State University, Columbus, OH, USA

⁶School of Nutrition, Chung Shan Medical University, Taichung City, Taiwan

⁷Department of Microbiology and Immunology, Montana State University, Bozeman, MT, USA

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ABSTRACT

More than 50% of the world's population is infected with *Helicobacter pylori*. *H. pylori* is the major causative agent of gastric ulcers and gastric cancer. *H. pylori* eradication using antibiotics either alone or together with a proton pump inhibitor is the primary strategy to decrease the incidence of gastric cancer. Although eradication therapy is effective, there are significant adverse effects and more importantly, resistance to antibiotics occurs, which represents a major therapeutic challenge. Multiple natural products have been shown to suppress *H. pylori* both *in vitro* and in animal model systems. However, only a handful of natural products have been evaluated in human clinical trials. The focus of this review is to summarize the results of published human clinical trials to assess the ability of natural products to reduce or eliminate *H. pylori* infections. Current evidence suggests that these products appear to have great potential to be developed as pharmaceutical candidates for eradication of *H. pylori*, hopefully both antibiotic-sensitive and antibiotic-resistant strains. Frequent consumption of locally produced foodstuff for controlling *H. pylori* infection in different countries around the world may well be a feasible long-term solution to fight against this worldwide prevalent pathogen.

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1. INTRODUCTION

It was estimated that more than 50% of the world's population is infected with *Helicobacter pylori* [1]. The bacterium highly links to gastric ulcers as well as gastric cancer and in 1994, it was classified as a group I carcinogen by the World Health Organization (WHO) [1]. *H. pylori* can induce gastric inflammation which could progress to superficial gastritis, atrophic gastritis, intestinal metaplasia, dysplasia, and ultimately to gastric cancer [2]. Eradication therapy using antibiotics and Proton Pump Inhibitors (PPIs) is the standard therapy, but in some regions of the world it fails due to emergence of antibiotic-resistant strains of the organism [3].

Numerous plant-derived compounds have been reported to inhibit *H. pylori* such as various polyphenols, flavonoids, quinones, coumarins, terpenoids, and alkaloids [2]. For example, quercetin, tea products, garlic extract, apple peel polyphenols, ginger-root extract, and many others have been studied. Mechanisms by which these compounds exert their anti-*H. pylori* effects are as follows: (1) inhibition of *H. pylori* enzymes such as urease which decrease

the acidity of gastric juice; (2) inhibition of adhesion of the organism to gastric mucosa; and (3) reduction in oxidative radicals that damage cellular macromolecules. Several reviews have summarized the use of natural products in suppressing *H. pylori* infection and their mechanisms of action using *in vitro* cell culture and *in vivo* animal models [2–8]. Readers are referred to these comprehensive reviews for details. The present review summarizes the results of published human clinical trials and discusses current evidence of the ability of natural products to inhibit *H. pylori* in humans. Multiple medicinal plant products inhibit *H. pylori* growth and *H. pylori*-induced gastric inflammation in humans. These natural products appear to have significant potential to be developed as pharmaceutical lead candidates for *H. pylori* eradication and for prevention of gastric diseases induced by *H. pylori* infection.

2. GASTRIC CANCER AND *H. PYLORI* INFECTION

Gastric cancer is the third leading cause of cancer-related mortality worldwide [9]. Drs J.R. Warren and B.J. Marshall were awarded for the Nobel Prize for Physiology and Medicine for the isolation and culture of *H. pylori*, and demonstrating its relevance to peptic

*Corresponding author. Email: gstoner@montana.edu

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ulcer disease [10]. Several case-controlled studies have shown that *H. pylori* seropositivity is associated with a significantly increased risk for gastric cancer (2.1–16.7-fold greater than the seronegative individuals) [11]. Encouragingly, epidemiologic studies indicate that there was a progressive decline in *H. pylori* infection during the last century in developed countries [1]. Nevertheless, more than 50% of the world's population remains infected with this organism, with the highest infection rates and gastric cancer incidence rates reported in lower socioeconomic countries [12].

To establish persistent infection, *H. pylori* escapes the immune system by mimicking host antigens [12]. This pathogen has the ability to escape the immune system and persist within host cells of the stomach leading to immune tolerance. Although establishment of *H. pylori* as a risk factor permits an approach to identify persons at increased risk for gastric adenocarcinoma, only a fraction of *H. pylori*-colonized persons develop the disease [12]. Recent data suggest that microbial–host interactions governed by differences in genotype acting in combination with permissive environmental conditions determine *H. pylori*-induced gastric cancer risk [11]. Therefore, it is not surprising that reducing an environmental factor such as salt intake may decrease the risk of gastric cancer irrespective of the presence of *H. pylori* [13].

3. CURRENT THERAPIES, EFFECTIVENESS, AND RESISTANCE

Helicobacter pylori is known to induce chronic inflammation in the gastric mucosa through its production of agents such as superoxide that induce DNA damage [14]. *H. pylori* eradication using antibiotic treatment in combination with a PPI is the major chemoprevention strategy to decrease the incidence of gastric cancer [11]. Eradication therapy is effective, but there is a steady increase in the emergence of antibiotic-resistant strains of the organism. A meta-analysis used data from 20 nationwide trials (3624 men and women) of *H. pylori* eradication in the United States to estimate the prevalence of *H. pylori* resistance to antibiotics and to characterize factors associated with resistance [15]. These factors included geographic region, age, sex, study year, ethnicity, ulcer status, and test method. The overall resistance to clarithromycin, metronidazole, and amoxicillin was 10.1% [95% Confidence Interval (CI), 9.1–11.1% (360 of 3571 patients)], 36.9% [95% CI, 35.1–38.7% (1063 of 2883 patients)], and 1.4% (95% CI, 1.0–1.8% (48 of 3486 patients)), respectively. Factors associated with resistance to different antibiotics varied. For example, clarithromycin resistance was associated with geographic region, older age, female sex, and inactive ulcer disease. Metronidazole resistance was associated with female sex, earlier year of study enrollment, Asian ethnicity, and use of an Epsilometer test. Amoxicillin resistance was low and was not significantly associated with any risk factor. These data are valuable for clinicians to consider when they decide which patients should have susceptibility testing when choosing appropriate anti-*H. pylori* treatments.

An important area of controversy in *H. pylori* eradication is the apparent difference in eradication rates seen in different countries. A recent meta-analysis showed that several factors may affect the outcome of therapy in different countries [16]. For example, individuals residing in northeast Asia had higher eradication rates than

those residing in Europe or other areas of Asia. In Japan, eradication therapy significantly diminished the prevalence of gastric cancer by around one-third [14,17]; however, studies from China [18] and the United States [19] reported different negative results [17]. Subgroup analysis suggests that eradication of *H. pylori* decreases the risk for development of gastric cancer and is most effective in *H. pylori* carriers who do not have atrophic gastritis or precancerous lesions [18].

It should be noted that other strains of *Helicobacter* such as *Helicobacter suis*, *Helicobacter heilmannii sensu stricto*, and *Helicobacter felis* may reside in the stomach of humans and animals [20]. Recent advancements in *in vitro* cultivation of these strains and recent reports from *in vivo* preclinical studies suggest that these strains may have more clinical significance than thought in the past. Additional studies of the health effects of these strains are warranted.

In addition to eradication therapy, vaccines against *H. pylori* have been developed in animals and tested in phase 3 clinical trials in humans. These trials have been largely unsuccessful, although *H. pylori*-specific immune responses have been induced [21]. New insights into promoting immunity and overcoming the immunosuppressive nature of *H. pylori* infections are required to improve the efficacy of the vaccines [21].

Recurrence of *H. pylori* infection after successful eradication occurs and is associated with relapse of gastroduodenal diseases [22]. Individual susceptibility and re-exposure to *H. pylori* are suggested as two major causes of reinfection. This represents a major concern that should be addressed by the research community.

4. HUMAN CLINICAL TRIALS OF NATURAL COMPOUNDS AGAINST *H. PYLORI*

There are multiple reports from around the world of the ability of natural products to inhibit *H. pylori* [7]. In the following sections, we summarize the ability of natural products to suppress *H. pylori* as evidenced from human clinical trials (Table 1), and we discuss their mechanisms of action (Figure 1).

4.1. Burdock Complex

Burdock complex contains burdock (*Arctium lappa*), angelica (*Angelica sinensis*), gromwell (*Lithospermum erythrorhizon*), and sesame (*Sesamum indicum*) oil commonly used in traditional Chinese medicine for treating various disorders [23]. Burdock complex was tested on asymptomatic *H. pylori*-infected patients in a randomized, double-blind, placebo-controlled clinical trial in Taiwan [23]. *H. pylori*-positive patients [Urea Breath Test (UBT) > 10%, $n = 36$] were enrolled and consumed burdock complex ($n = 19$) or placebo ($n = 17$) for 8 weeks. Antioxidant capacity, total phenol, UBT, and various inflammatory markers were analyzed at baseline and on the 4th, 8th, and 10th weeks after intervention. In addition, endoscopic examinations were carried out at baseline and at the 10th week. *H. pylori*-infected patients who had consumed burdock complex for 8 weeks had significantly decreased UBT values. In addition, the inflammatory markers (interleukin-8 and tumor necrosis factor- α) were suppressed, antioxidant activity

Table 1 Effects of natural products against *Helicobacter pylori* in humans

Natural product	Human patient	Treatment duration	Location	Effects	References
Burdock complex	Adults (19 treated, 17 placebo) who were asymptomatic but with <i>H. pylori</i> infection	8 weeks	Taiwan	Decreased UBT value and inflammatory markers, increased antioxidant activity.	[23]
Comifer green needle complex	Adults (26 treated, 24 no treatment) with precancerous gastric lesions	6 months	Russia	Partial regression of dyspeptic symptoms, eradication of <i>H. pylori</i> infection, reduction in endoscopic signs of gastritis, an increase of pepsinogen-pepsin in gastric juice, and total regression or reduction in the degree of intestinal metaplasia and lymphoplasmacytic infiltration.	[24]
Bovine lactoferrin	24 Adults and 25 children (aged 4–13 yr) who were asymptomatic and with minimal upper gastrointestinal symptoms, positive for <i>H. pylori</i> infection	12 weeks	Japan	Decreased UBT value at week 12, but it increased 4 weeks after stopping intervention.	[25]
Bovine lactoferrin	74 Adult <i>H. pylori</i> -positive patients with symptomatic dyspepsia, with or without active ulcer	Group A: 24 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) plus lactoferrin for 7 days; Group B: 26 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) for 7 days; Group C: 24 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) for 10 days	Italy	The eradication rates in groups A, B, and C were 100% (24/24), 76.9% (20/26; 95% CI, 61–93%), and 70.8% (17/24; 95% CI, 53–89%), respectively. Significant differences in eradication rates were found between group A and group B, and group A and group C, but not between group B and group C.	[26]
Bovine lactoferrin	402 Adult <i>H. pylori</i> -positive patients	Group A: 136 patients received triple therapy (esomeprazole, clarithromycin, and tinidazole) for 7 days; Group B: 132 patients received lactoferrin for 7 d followed by triple therapy (esomeprazole, clarithromycin, and tinidazole) for 7 days; Group C: 134 patients received triple therapy (esomeprazole, clarithromycin and tinidazole) plus lactoferrin for 7 days	Italy	The eradication rate was 77% (105/136) in group A, 73% (97/132) in group B, and 90% (120/134) in group C. The incidence of side effects was 9.5%, 9%, and 8.2% in groups A, B, and C, respectively.	[28]
Cranberries	295 Asymptomatic children (aged 6–16 years)	Cranberry juice and the probiotic <i>Lactobacillus johnsonii</i> La1 (La1) alone or in combination for 3 weeks	Chile	The combination group showed a slight but nonsignificant increase in suppression of <i>H. pylori</i> when compared with either intervention alone. When treatment was stopped, <i>H. pylori</i> was positive again in 80% of the negative patients who received cranberry and/or La1 previously. Negative test results: 14/97 in the cranberry juice group versus 5/92 in the placebo group.	[30]
Cranberries	189 Adults with <i>H. pylori</i> infection: $n = 97$ for the cranberry juice group and $n = 92$ for the placebo group	90 days	China		[31]
Probiotics	A meta-analysis from 33 randomized, controlled trials involving a total of 4459 patients	Multiple durations	Multiple countries	Supplementation of specific strains of probiotics with eradication therapy may be considered as an addition for increasing eradication rates. These probiotics are <i>Lactobacillus acidophilus</i> , <i>Lactobacillus casei</i> DN-114001, <i>Lactobacillus gasseri</i> , and <i>Bifidobacterium infantis</i> 2036.	[32]

Continued

Table 1 | Effects of natural products against *Helicobacter pylori* in humans—Continued

Natural product	Human patient	Treatment duration	Location	Effects	References
Vitamins and garlic	3365 Adult patients (2258 <i>H. pylori</i> positive, 1107 <i>H. pylori</i> negative)	Vitamin supplement: vitamin C, vitamin E, and selenium for 7.3 years Garlic supplement: aged garlic extract and steam-distilled garlic oil for 7.3 years	China	Long-term vitamin or garlic supplementation alone or given after eradication therapy did not have beneficial effects on the prevalence of precancerous gastric lesions or incidence of gastric cancer.	[33]
Garlic	A meta-analysis of 18 studies (142,921 patients)	Multiple durations	Multiple countries	Current epidemiological evidence is insufficient to reach a definitive conclusion regarding the association of garlic consumption with <i>H. pylori</i> infection.	[36]
β -Carotene, ascorbic acid	852 Adult patients with confirmed histologic diagnoses of multifocal nonmetaplastic atrophy, intestinal metaplasia, or dysplasia	Ascorbic acid or β -carotene for 6 years	USA	Both ascorbic acid and β -carotene increase lesion regression rates. Combinations of triple therapy with ascorbic acid or β -carotene did not significantly increase the rates of lesion regression. By contrast, the combination treatment was worse than placebo.	[37]
β -Carotene, ascorbic acid	795 Adult patients with confirmed histologic diagnoses of multifocal nonmetaplastic atrophy, intestinal metaplasia, or dysplasia	Ascorbic acid or β -carotene for 6 years and follow-up for another 6 years	USA	Ascorbic acid and β -carotene were effective to increase the regression rates of lesions at the first 6 years. However, this effect disappeared during the 6 years of follow-up of no supplementation, and no residual effect was observed.	[19]

CI, confidence interval; UBT, urea breath test.

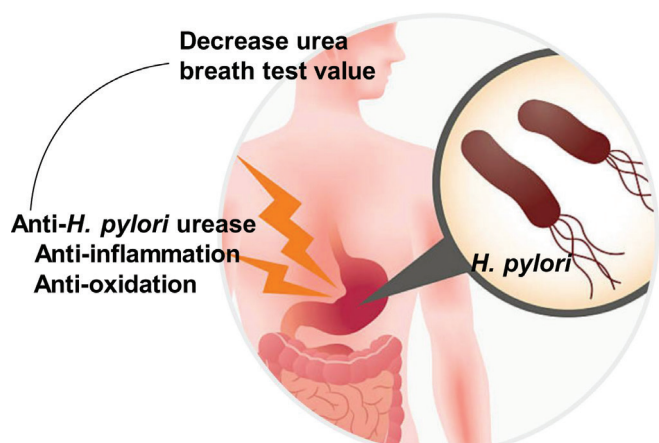


Figure 1 | Mechanisms by which natural compounds suppress *Helicobacter pylori* in humans. Adapted from <https://centrefordigestivediseases.com/for-patients/tests/urea-breath-test/>.

was increased, and phenolic levels were increased when compared with placebo. Endoscopic examination indicated that consumption of burdock complex considerably reduced the stomach ulcers.

4.2. Conifer Green Needle Complex

In an observational pilot study conducted in Russia, a tablet form of Conifer Green Needle Complex (CGNC) extracted from *Pinus sylvestris* and *Picea abies* (L) Karst was prescribed to 26 patients with precancerous gastric lesions for 6 months [24]. Another 24 patients received no treatment (control/no placebo). Compared with control patients, CGNC-treated patients showed total or partial regression of dyspeptic symptoms, a decrease in *H. pylori* infection, a decrease in endoscopic signs of gastritis, an increase of pepsinogen-pepsin in the gastric juice, and total regression or reduction in the degree of intestinal metaplasia and lymphoplasmacytic infiltration.

4.3. Bovine Lactoferrin

Bovine lactoferrin, a protein with antimicrobial activity, was evaluated for its efficacy to suppress *H. pylori* colonization in a randomized, double-blind, placebo-controlled study in Japan [25]. Fifty-nine healthy individuals (34 adults and 25 children, aged 4–13) positive for *H. pylori* infection were recruited and received bovine lactoferrin or placebo for 12 weeks. UBT was performed before, during, and at the end of administration, and again at 4 weeks after administration (week 16). Positive response was defined as a >50% decrease in the UBT value at the end of administration. At week 12, a significantly higher positive response was observed in 10 of 31 lactoferrin-treated individuals versus one of 28 placebo-treated cases, combining data from both adults and children. There was no significant difference in positive response rates between adults and children. However, at week 16, the responders in both adults and children had increased UBT, suggesting that continuous consumption of lactoferrin is required to eradicate *H. pylori*.

The efficacy of a triple antibiotic therapy regimen in combination with bovine lactoferrin for the eradication of *H. pylori* infection was investigated in an open, randomized, single-center study in

Italy [26]. This trial included 74 *H. pylori*-positive patients with symptomatic dyspepsia, with or without active ulcer. There were three groups: 24 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) + lactoferrin for 7 days (group A); 26 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) for 7 days (group B); and 24 patients received triple therapy (rabeprazole, clarithromycin, and tinidazole) for 10 days (group C). Eight weeks after the end of treatment, the *H. pylori* status was assessed using UBT or *H. pylori* stool antigen test. The results showed that the eradication rates in groups A, B, and C were 100% (24/24), 76.9% (20/26; 95% CI, 61–93%) and 70.8% (17/24; 95% CI, 53–89%), respectively. Significant differences were found between groups A and B and groups A and C, but not between groups B and C. These data suggest that combination of lactoferrin and triple therapy is more effective in *H. pylori* eradication. One limitation of this trial was that resistance to antibiotics was not determined. Although clarithromycin resistance is low in Italy (<2%), it is about 10% in the United States and metronidazole resistance is about 37% in the United States [15]. Therefore, it will be important to determine the ability of lactoferrin to effectively eradicate clarithromycin resistance in countries such as the United States where resistance is higher than in Italy [27]. Of importance in this study is that the incidence of adverse effects was not increased by the addition of lactoferrin to the antibiotic therapy. However, the study was too small to assess whether lactoferrin diminishes antibiotic-related side effects.

Results from the aforementioned study were confirmed by another study conducted by the same group of scientists in Italy [28]. In a multicentered prospective study [28], 402 *H. pylori*-positive adult patients were assigned to one of three following regimens: 7 days of triple therapy (esomeprazole, clarithromycin, and tinidazole; group A); 7 days of lactoferrin followed by 7 days of triple therapy (esomeprazole, clarithromycin, and tinidazole; group B); 7 days of triple therapy (esomeprazole, clarithromycin, and tinidazole) followed by 7 days of lactoferrin (group C). The results showed that the eradication rate was 77% (105/136) in group A, 73% (97/132) in group B, and 90% (120/134) in group C. The incidence of side effects was 9.5%, 9%, and 8.2% in groups A, B, and C, respectively. This study demonstrated that bovine lactoferrin is an effective addition to the triple therapy (7 days) for eradication of *H. pylori* infection. However, the adverse effects were not significantly decreased by adding lactoferrin [28].

4.4. Cranberries

The potential of cranberries to inhibit *H. pylori* infection in humans has been summarized in a recent review [29]. The effects of regular consumption of cranberry juice and the probiotic *Lactobacillus johnsonii* La1 (La1), either alone or in combination, on inhibition of *H. pylori* colonization in children were investigated in a study in Chile [30]. The study was a randomized, controlled, double-blind multicentric trial carried out in 295 asymptomatic children (aged 6–16) positive for *H. pylori*, as determined using UBT [30]. There were four groups: cranberry juice + La1 (CB/La1), placebo juice + La1 (La1), cranberry juice + heat-killed La1 (CB), and placebo juice + heat-killed La1 (control). The children consumed cranberry juice and La1 product for 3 weeks and then a second UBT was carried out. After a 1-month washout, a third UBT was performed in children who were negative in the second UBT. *H. pylori* eradication

rates were 1.5% in the control and 14.9%, 16.9%, and 22.9% in the La1, CB, and CB/La1 groups, respectively. The increase in UBT in the CB/La1 group was not significant compared with the La1 and CB groups. In the third UBT, which was performed in 19 of the 38 children who were negative in the second UBT, *H. pylori* was positive in 80% of the children. These results suggest that regular and/or frequent intake of cranberry juice or La1 could be a useful management in asymptomatic children with *H. pylori* infection. The results also suggest that there are no apparent synergistic inhibitory effects on *H. pylori* infection when patients consumed both cranberry juice and the probiotic *L. johnsonii* La1 (La1) together.

A trial conducted in China reported similar results. In Linqu County of Shandong Province in China, a prospective, randomized, double-blind, placebo-controlled trial suggested that cranberry juice suppresses *H. pylori* infection [31]. A total of 189 adults with positive *H. pylori* were randomly assigned into two groups to receive either cranberry juice ($n = 97$) or placebo juice ($n = 92$) for 90 days. *H. pylori* infection was determined using UBT. The study concluded that 14 of the 97 adults in the cranberry juice treatment group versus five of the 92 in the placebo group yielded negative results in UBT.

4.5. Probiotics

A meta-analysis from 33 randomized controlled trials involving a total of 4459 patients suggested that supplementation of specific strains of probiotics (*Lactobacillus acidophilus*, *Lactobacillus casei* DN-114001, *Lactobacillus gasseri*, and *Bifidobacterium infantis* 2036) with *H. pylori* eradication therapy may be considered as an option for increasing eradication rates, particularly when antibiotic therapies are relatively ineffective [32].

4.6. Vitamins and Garlic

In China, a randomized trial was set out to test the eradication effects of one-time antibiotic treatment for *H. pylori* infection followed by a long-term vitamin or garlic supplementation in decreasing gastric lesion prevalence [33]. A total of 3365 eligible individuals selected from 13 villages in Linqu County, Shandong Province, China who were positive for *H. pylori* were randomly assigned to three interventions or placebo in a factorial design [33]. These groups were *H. pylori* treatment (amoxicillin and omeprazole) for 2 weeks; vitamin supplement (vitamin C, vitamin E, and selenium) for 7.3 years; and garlic supplement (aged garlic extract and steam-distilled garlic oil) for 7.3 years. Patients were treated for *H. pylori* eradication and with vitamin or garlic supplement, either alone or in combination. Histopathologic examination was performed on biopsies collected by endoscopies to determine the prevalence of precancerous gastric lesions. Long-term vitamin or garlic supplementation alone or in combination with eradication therapy did not have beneficial effects on either the precancerous gastric lesion prevalence or gastric cancer incidence [33] despite the fact that antioxidants were shown to reduce the risk of gastric cancer in nutritional intervention trials in China [34]. The lack of response in this patient group might have been due to their consumption of adequate amounts of vitamin and garlic before treatment; therefore, there were no added benefits of supplementation of these nutrients [35]. Individuals at the highest risk should be monitored endoscopically to detect dysplasia and early cancer.

The association of garlic with *H. pylori* infection and gastric cancer risk has been investigated in multiple human clinical trials around the world. A meta-analysis of 18 studies (142,921 patients) demonstrated that high garlic consumption (comparing the highest category with the lowest) was associated with a reduced gastric cancer risk [Odds Ratio (OR), 0.51; 95% CI, 0.44–0.57] [36]. However, this association became nonsignificant if prospective studies were included in the analysis (OR, 0.95; 95% CI, 0.66–1.24).

Pooled evidence from epidemiologic studies, mainly case-control studies, suggested a significant inverse association of garlic intake with gastric cancer risk. However, when analyzing data from all epidemiologic studies, the evidence is insufficient to reach a definitive conclusion concerning the association between garlic consumption and *H. pylori* infection.

4.7. Beta-Carotene and Ascorbic Acid

A total of 852 adult patients with confirmed histologic diagnoses of multifocal nonmetaplastic gastric atrophy ($n = 194$), intestinal metaplasia ($n = 579$), or dysplasia ($n = 79$) were recruited in a randomized, controlled chemoprevention trial conducted in the United States [37]. Patients were assigned to receive anti-*H. pylori* triple antibiotic therapy and/or dietary supplementation with ascorbic acid or β -carotene [37]. Gastric biopsy specimens were collected at baseline and 6 years for histopathological examination. The results showed that the rates of regression were significantly increased by all three basic interventions. For anti-*H. pylori* treatment, relative risk was 4.8 (95% CI, 1.6–14.2); for β -carotene treatment, the relative risk was 5.1 (95% CI, 1.7–15.0); and for ascorbic acid treatment, the relative risk was 5.0 (95% CI, 1.7–14.4) in patients with atrophy. Among patients with intestinal metaplasia, relative risks of regressions were 3.1 (95% CI, 1.0–9.3), 3.4 (95% CI, 1.1–9.8), and 3.3 (95% CI, 1.1–9.5) for anti-*H. pylori*, β -carotene, and ascorbic acid treatment, respectively. However, the regression rates were not increased by combination treatments. Importantly, 74% of the treated cases had negative *H. pylori* infection and they also had significantly decreased precursor lesions; relative risks for patients with atrophy were 8.7 (95% CI, 2.7–28.2) and for those with intestinal metaplasia were 5.4 (95% CI, 1.7–17.6). Accordingly, this study suggested that in the very high-risk population, anti-*H. pylori* antibiotic treatment and antioxidant micronutrient dietary supplementation may delay the precancer lesion progression. This approach may be useful to prevent gastric cancer.

At the 12-year follow-up of the aforementioned trial [19], 795 adults with preneoplastic gastric lesions were randomized to receive anti-*H. pylori* antibiotic treatment and/or antioxidants. At baseline, and at 3, 6, and 12 years, gastric biopsies were collected for histopathological examination. At baseline and 12 years, 97% and 53% of patients were *H. pylori* positive, respectively. More rapid healing was observed in patients with years free of *H. pylori* infection and those with less advanced lesions. In other words, regression rate of preneoplastic gastric lesions is equal to the square of time that patients were *H. pylori* infection free. These findings suggest that patients who have preneoplastic gastric lesions and *H. pylori* infection should be treated by antibiotic therapy. However, patients treated with antioxidant supplements did not have an additional benefit compared with those treated with antibiotics only [19]. Although an antioxidant effect was observed at 6 years, it disappeared during the 6 years of no supplementation,

and no residual effect was observed. This study also showed that acute gastric inflammation disappeared soon after *H. pylori* treatment, and chronic inflammation responded at a slower pace but was still reduced significantly after 12 years [19]. Accordingly, gastric cancer chemoprevention via eradication of *H. pylori* infection is a viable option, but the greatest beneficial effects might not be evident in the first 3–6 years of observation [19]. It should be noted that analysis at 6 years showed that there was a significant negative interaction between *H. pylori* treatment and β -carotene supplementation; some patients who received antibiotic treatment for *H. pylori* showed improvement, but those who received *H. pylori* treatment and β -carotene supplementation did worse than those on placebo [37]. This finding may suggest that both interventions likely have a final common pathway—decreasing oxidative damage to the gastric epithelial cell.

5. NATURAL COMPOUNDS FIGHT AGAINST RESISTANT *H. PYLORI*

Antibiotic treatment combined with a PPI is effective in eradication of *H. pylori* [11]. However, the occurrence of drug-resistant *H. pylori* and the adverse effect of antibiotics are significant problems of this therapy [7]. The adverse effects include the disturbance of human gastrointestinal microflora, severe complications such as liver and kidney dysfunction, and rejection of or allergy to antibiotics [7]. Natural products that have been shown to suppress clinical isolates of resistant *H. pylori* strains are Epigallocatechin-3-Gallate (EGCG) and Epicatechin Gallate (ECG) [38], aqueous and ethanol extracts of propolis or *Zingiber officinale* [39], sulforaphane [40], and paeonol and benzoic acid identified in *Paeonia lactiflora* root [41]. Although they were tested in *in vitro* assays, together with several natural products that have been shown to suppress *H. pylori* in human clinical trials, it is now important to test these natural products in patients with resistant *H. pylori*.

6. FUTURE PERSPECTIVES

Many foodstuff such as burdock complex, CGNC, bovine lactoferrin, cranberries, probiotics have exhibited inhibitory activity against the growth of *H. pylori* in humans. It is encouraging that natural compounds such as EGCG and ECG from tea, the aqueous and ethanol extracts of *Propolis* or *Z. officinale*, sulforaphane, and paeonol and benzoic acid identified in *P. lactiflora* root suppressed clinical isolates of resistant *H. pylori* strains. The crucial next step is to test these promising natural products for their ability to suppress resistant *H. pylori* in humans. Although these natural products may reduce the bacterial load of *H. pylori*, the effects are usually transient and *H. pylori* is not totally eradicated in the majority of the patients. Considering the feasibility of consuming foodstuffs for controlling *H. pylori* infection in different countries around the world, it is important to find locally consumed foods that are anti-*H. pylori* as long term solutions to fight against this world-wide prevalent pathogen.

CONFLICTS OF INTEREST

The authors declare they have no conflicts of interest.

AUTHORS' CONTRIBUTION

LSW contributed in literature search, manuscript writing, editing, final approval. CEE, JY, YWH, JL, LM, PS, MY and KO contributed in concept approval. CKW contributed in editing, final approval. GS contributed in manuscript writing, editing, final approval.

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