A fucoidan plant drink reduces *Helicobacter pylori* load in the stomach: a real-world study

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**Background:** *Helicobacter pylori* (Hp) infection is highly prevalent globally and is predominantly managed by antibiotics. Recently, the anti-adhesive, antioxidant, antitoxin, immunomodulatory, anti-coagulant, and anti-infective activities of fucoidan, a polysaccharide extracted from brown seaweeds, have been widely studied, and the results showed promise. Fucoidan has the potential to be utilized in Hp eradication therapy. Our present clinical study was designed to evaluate the efficiency of Lewuyou⁶, a fucoidan plant drink (FPD) in eradicating Hp in humans.

**Methods:** This multi-center, clinical study was conducted between October 2020 and July 2021. Hp infection was confirmed by urea breath test (UBT). A total of 122 patients with confirmed Hp infection were enrolled; after exclusion of incomplete data, 85 eligible patients (37 males and 48 females aged 20–81 years) were included in the final analysis. FPD (50 mL per vial) was orally administered twice daily for a 4-week cycle, and 41 patients completed an 8-week cycle.

**Results:** No adverse event (AE) was reported in all 122 participants who had consumed FPD. The Hp eradication rate and clearance rate were 77.6% (66/85) and 20.0% (17/85), respectively, after 4 weeks of FPD consumption and 80.5% (33/41) and 26.8 (11/41) , respectively, after 8 weeks of consumption.

**Conclusions:** The 4- and 8-week protocols of FPD consumption were safe and effective at reducing Hp load on the gastric mucosa, with Hp eradicated in the majority of participants.

**Keywords:** Helicobacter pylori (Hp); eradication; fucoidan; treatment; real world

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Introduction

According to epidemiological research data, the global prevalence of *Helicobacter pylori* (Hp) infection among adults is approximately over 50% with infection rates higher in developing countries (1). The infection rate among children is also considerably high (2). Hp colonizes the gastric mucosa during childhood and persists without successful eradication therapy. Chronic infection can lead to serious complications such as peptic ulcer disease, gastric adenocarcinoma and mucosa-associated lymphoid tissue (MALT) lymphoma (3). Hp infection can lead to changes to the delicate balance of the gastric microbiota which has implications in human health and disease (4). Hp infection can induce insulin resistance and predispose individuals to the development of type 2 diabetes mellitus (5). Recent studies have suggested an increased risk of diabetes mellitus and hypertension (6,7). Hp infection is widely recognized as an infectious disease that warrants antimicrobial therapy, with bactericidal eradication being an immediate treatment goal (8). In East Asian countries, Hp eradication therapy has reduced gastric cancer incidence in healthy individuals and patients with gastric tumors. It even appears to lower the mortality associated with gastric cancer (9). However, almost all the currently available treatments for Hp, including the standard triple therapy [STT; a proton pump inhibitor (PPI) plus 2 antibiotics, usually clarithromycin and amoxicillin] and bismuth-free or bismuth-containing quadruple therapy (BQT), are based on a combination of antibiotics and adjuvants. BQT usually consists of a PPI, bismuth, metronidazole and antibiotics (e.g., tetracycline). The recommended STT for eradicating Hp traditionally combines PPI with clarithromycin, amoxicillin, or metronidazole. Failure rate can be up to 20%, predominately due to the increasing antibiotic resistance worldwide (10). In the Asia-Pacific region, the prevalence of Hp resistant to clarithromycin, levofloxacin, and metronidazole has increased to 21%, 27%, and 45%, respectively (11). Patient compliance with STT and BQT is less than satisfactory, with the classical BQT being the least well tolerated: approximately 37% of overall cases had adverse events (AEs), with taste disturbance (7%), diarrhea (7%), nausea (6%), and abdominal pain the most common (12). When used as first-line empirical therapy, the 10-day BQT had a significantly higher eradication rate but lower compliance rate (23.1% vs. 9.1%) than 7-day PPI-clarithromycin containing triple therapy (13). Therefore, Hp eradication remains a significant clinical challenge. The development of effective, less toxic eradication therapies remains a key research topic (14), such as scutellaria baicalensis (15). Recently, the anti-adhesive, antioxidant, antitoxin, immunomodulatory, anti-coagulant, and anti-infective activities of *Ascophyllum nodosum* (knotted wrack; a type of brown seaweed that contains fucoidan) extracts were extensively studied. It was proposed that fucoidan may be classified as a new drug that could be included in therapeutic regimens for Hp eradication (16). This novel treatment could avoid antibiotic associated side effects and resistance to eradicate HP infection and avoid Hp associated complications (17).

We conducted a real-world clinical study to explore the clinical efficacy of fucoidan (a compounded product) for the clearance of Hp in the stomach. This study was aimed at the assessment whether a drink containing fucoidan was clinically valuable for eradicating Hp and lowering Hp load. We present this article in accordance with the TREND reporting checklist (available at https://tgh.amegroups.com/article/view/10.21037/tgh-23-63/rc).

**Methods**

**Compound product**

The compounded product was a fucoidan plant drink (FPD) (Lewuyou®). It is categorized as a “food product” and containing the following main ingredients: water, radish seed compounded plant beverage (including water,
isomalto-oligosaccharide, glucose, yam, radish seed, and hawthorn), apple juice concentrate, broccoli powder (including maltodextrin and broccoli), fucoidan (200 mg), *Lycium ruthenicum* Murray powder, instant green tea powder, erythritol, pectin, citric acid, DL-malic acid, sucralose, stevioside, and edible flavors and fragrances. The compound was used due to the following principles: high affinity, the bio-targeted polysaccharide interferes with Hp's recognition of gastric mucosal epithelial cells and removes Hp by adhesion; potent antioxidants to reduce gastric inflammation and repair gastric mucosa and to improve gastric function.

In China, there is currently no requirement for ethical approval for the use of “food” in research. So, this study did not undergo ethics committee review. All research subjects will be informed of the intervention drink before enrollment. All participants agreed to take the intervention beverage daily as per the study requirements verbally.

**Experimental drink**

The FPD was presented as a liquid beverage in brown glass vials (50 mL in each vial). The product requires storage at room temperature and protection from light.

**Patients and study design**

**Patient registration**

Patients who volunteered to receive the anti-Hp FPD were registered as participants. All participants were treated with FPD. Medical histories and demographic data were collected, including age, gender, medical history, history of smoking, history of alcohol consumption, and history of coffee or tea drinking. Smoking was defined as smoking 1 pack of cigarettes or more per week.

**Enrollment**

This study is a multicenter real-world investigation. A total of 122 consecutive adult Hp-infected outpatients and health check-up patients aged 20–81 (41.47±12.23) years were enrolled in 7 centers in China (Taian, Taishan, Shanghai, Jinan, Dongying, Xian, Liaocheng) between October 2020 and July 2021. A diagnosis of Hp infection was made according to the following criteria: (I) a recent (within 7 days) positive result of 13C- or 14C-urea breath test (UBT); and/or (II) endoscopy, histologically confirmed Hp infection.

**Exclusion criteria**

(I) Withdrawal from the study within 1 month (4 weeks); (II) inability to respond to the test request and/or failure to provide UBT results within 5 weeks; (III) methods inconsistent between pre- and post-testing.

**Grouping**

**Real-world design**

13C- or 14C-positive Hp carriers were enrolled from large general hospitals or health check-up centers. The changes in 13C or 14C values were compared before and after FPD consumption. The participant activities (including diets) were not restricted during the study period.

**FPD consumption**

FPD (1 vial each time) was consumed twice daily (after waking and before bedtime) on an empty stomach for 8 consecutive weeks. The drink was shaken well before consumption.

Before and after the therapeutic intervention for Hp eradication, patients (carriers) were asked to return their hospitals at week 2 to receive an assessment of medication adherence and AEs, as specified in the study protocol. UBT was performed at 4 weeks (28±3 days) and 8 weeks (56±3 days) to learn the Hp status and thus assess the therapeutic effectiveness. Effectiveness data: 14C and 13C values were recorded to determine the changes in Hp status. Hp clearance was defined as a negative DOB (delta over baseline) value of UBT after the intervention. The intervention was judged as effective when there was a reduction in the 14C or 13C values.

**Primary endpoint**

Hp eradication and clearance rates at weeks 4 and 8. The AEs during the treatment period were also analyzed.

**Diagnosis of Hp infection and efficacy**

**Blinding**

The staff performing the 13C- or 14C-UBT were blind to the details of this study.

**UBT**

UBT was performed before enrollment and at weeks 4 and 8 during FPD consumption. If possible, the test must be completed in the same institution using the same method.
If the test was performed using a different method before and after FPD consumption and the data were >0, the data were used for reference only and were not regarded as valid data; these patients were not included in the final analysis.

**Evaluation of efficacy**

The difference between the baseline and post-intervention values was regarded as the eradication volume of Hp in the stomach after FPD consumption, and the ratio between the eradication volume and baseline value was the eradication rate. The FPD intervention was regarded as effective if the eradication rate was >0, and the Hp status was defined as “cleared” if the eradication rate was 1.

**AEs and adherence during the follow-up period**

**Safety data (if any)**

The type, severity, and duration of AEs, serious adverse events (SAEs) and, in particular, adverse drug reactions (ADRs) were recorded in detail at each visit. AEs that affected the patient quality of life including abdominal pain, diarrhea, constipation, dizziness, dysgeusia, headache, anorexia, nausea, vomiting and rash were also recorded.

**Adherence**

“Adherence” was defined according to whether FPD was consumed in keeping with the trial protocol and when and why it was discontinued. Poor adherence was defined if the FPD was consumed less than 80% of the set amount or if more than 50% of doses were missed.

**Statistical analysis**

Treatment outcomes and main factors were compared using the software SPSS 10.1 (IBM Corp., Chicago, IL, USA). The chi-square test, with or without Yates’ continuity correction and Fisher’s exact test, was performed when appropriate.

**Results**

**Participant characteristics**

A total of 122 patients with Hp infection were enrolled in this study. All participants were asked to consume FPD, and the effects of the drink on 13C- or 14C-UBT results were analyzed. A total of 85 patients (37 males and 48 females aged 20–81 years (mean: 41.5±12.2 years)) completed the first phase of the study (17 in the 13C group and 68 in the 14C group). Among the 85 patients, 41 patients (5 in the 13C group and 36 in the 14C group) completed the second phase of the study and were considered eligible cases. Among the 41 patients who completed the 8-week intervention, the data of 7 patients were collected 2 weeks after cessation of the intervention.

**Hp eradication rate**

After 4 weeks of FPD consumption, the Hp eradication rate was 77.6% (66/85) and the clearance (negative conversion) rate was 20.0% (17/85); after 8 weeks of administration, the Hp eradication rate was 80.5% (33/41) and the clearance rate was 26.8 (11/41).

The actual 13C/14C values were compared before (at day 0) and after (4 weeks later, n=85; at 8 weeks later, n=41) FPD consumption. A line graph visualizing the overall decreasing trend and the number of cases reaching point 0 is displayed in Figure 1. We used scatter plots to illustrate the declining trend of Hp infection in patients at baseline, 4 weeks, and 8 weeks post-intervention. We set UBT value =4 as the cutoff for negative/positive classification (13C); and UBT value =99 as the cutoff for negative/positive classification (14C) (Figure 2).

**Symptom improvement**

Symptoms, including abdominal distension, nausea, belching, and acid reflux were improved in some patients. In a post-bone marrow transplant, for a patient with intestinal rejection combined with cytomegalovirus (CMV) infection who had been producing 30 bloody, watery stools per day, the frequency of defeation decreased to 4 times/day after FPD consumption.

**AEs and complications**

AEs were investigated among all participants. No AE occurred in all 122 participants from the first day of administration to week 8. The medication compliance rate was 69.7% in the first month and 33.6% in the second.

**Discussion**

We conducted a real-world clinical study to explore the clinical efficacy of fucoidan (a compounded product) for
Figure 1 Changes in the values of 13C- or 14C-UBT before and after fucoidan plant drink consumption. (A) UBT value of 13C at week 4 (vs. before consumption (day 0)); (B) UBT value of 14C at week 4 (vs. day 0); (C) UBT value of 13C at week 8 (vs. day 0); (D) UBT value of 14C at week 8 (vs. day 0). UBT, urea breath test.
the clearance of Hp in the stomach. It was found in this small-sample clinical observation study that fucoidan was clinically valuable for eradicating Hp in the stomach and lowering Hp load, as validated by the DOB values of 13C- or 14C-UBT.

The infection rate of Hp is extremely high worldwide, with the highest prevalence in developing countries (1). Current guidelines recommend using triple or quadruple therapy as the treatment of choice for Hp infection, but the failure rate exceeds 20%. The rapid rise of antibiotic resistance, poor compliance and disruption of the gut microbiota has stimulated interest in alternative therapies (18). Among them, marine biologics are particularly promising. For example, epinecidin-1, a multifunctional antimicrobial peptide produced by Epinephelus coioides, was shown to have the potential to replace antibiotics with its significant efficacy in fighting against *Staphylococcus aureus* (*S. aureus*) and Hp (19).

Epinecidin-1 (Epi-1) was found to have potent bactericidal activity against Hp *in vitro* and modulated Hp-induced host immune responses in a mouse model (20).

Fucoidan (also known as fucoidin or sulfated fucan), first extracted from seaweed in 1913, is a negatively charged, highly hygroscopic polysaccharide soluble in water and acid solutions (21). It contains L-fucose and sulfate groups and is mainly derived from brown algae, red algae, and some marine invertebrates. It is well known amongst the food and pharmaceutical industry, due to its good therapeutic effects on some specific diseases. Fucoidan is not a single-structural compound, but contains xylose, mannose, galactose, arabinose, and glucuronic acid. Its biological functions are ascribed to its unique biological structure. Classical bioactivities associated with fucoidan include antioxidant, anti-tumor, anti-coagulant, anti-thrombotic, immunoregulatory, anti-viral and anti-inflammatory...
effects (22). A study assessing the bioactive antimicrobial capability of fucoidan ("Generally Recognized as Safe" approval—European Commission December 2017) from different species of Phaeophyceae algae (Fucus vesiculosus, Undaria pinnatifida, Macrocystis pyrifera) against Hp, all the studied fucoids showed bacteriostatic and bactericidal effects at the studied concentrations (5–100 μg/mL) and exposure time (0–7 days) (23).

The exact mechanism by which fucoidan eradicates Hp remains unclear. Fucoidan is believed to interfere with the linkage/attachment of Hp to gastric mucosal cells. Shibata et al. found that fucoidan could block the attachment of Hp to gastric cells (24). An in vitro experiment also confirmed that Cladosiphon fucoidan inhibited the Hp attachment to porcine gastric mucin at pH 2.0 and 4.0 (17). Based on this property, genipin-crosslinked low molecular weight fucoidan/chitosan-N-arginine nanogels (FCSA) were prepared for targeted delivery of amoxicillin to the site of Hp-infected human gastric adenocarcinoma epithelial (AGS), cells and the negatively charged nanogels (n-FCSA) adhered to Hp and exhibited pH-responsive drug release property to reduce cytotoxic effects in Hp-infected AGS cells (25). In the study conducted by Chua et al., adding fucoidan polysaccharide formulation (1,000 μg/mL) to the culture medium did not inhibit the growth of Hp. However, adding different fucoidan salt formulations (1,000 μg/mL) to the tissue culture medium exhibited toxicity towards AGS cells, reducing the number of viable AGS cells. Furthermore, a fucoidan polysaccharide formulation at a concentration of 100 μg/mL significantly decreased the adhesion of HP, suggesting a potential mechanism of fucoidan treatment for HP infection involving AGS cell toxicity and reduction of HP adhesion (26).

Furthermore, it was demonstrated in Cai et al. research that a combination of fucoidan polysaccharides and evening primrose extract (FEMY-R7), exhibited complete inhibition of Hp in vitro at a concentration of 100 μg/mL. Animal experiments revealed that FEMY-R7 cleared gastric mucosal infection by direct killing the bacteria and preventing their adhesion and invasion (27). In a clinical study, humans confirmed to be infected with Hp were orally administered twice daily with a capsule containing 150 mg FEMY-R7 for 8 weeks. FEMY-R7 significantly decreased the value in UBT and the serum pepsinogens I and II levels. The results indicate that FEMY-R7 eliminates Hp from animal and human gastric mucosa and improves gastric function (28).

Additionally, the observations on healthy volunteers indicate that fucoidan can be absorbed and metabolized by the human body, with absorption being correlated to the frequency of Hp infection and seaweed product usage (29,30). The ingestion of fucoidan may also exert antimicrobial, antiviral, and even anti-tumor effects by improving the body's immune status.

A study on mice (Tomori et al.) revealed that when different doses of fucoidan were administered orally continuously for 6 weeks, immune cell proliferation, interleukin (IL)-2, macrophage phagocytes, and serum antibodies (IgM, -G, -A) increased significantly. Still, the levels of IL-4, -5, and IgE decreased significantly. These results indicated that fucoidan modulated cellular and humoral immunity (31). When cells were incubated with fucoidan in the presence of a viral mimic, fucoidan inhibited the production of some cytokines, chemokines, and prostaglandin E2. Additionally, fucoidan may help alleviate airway inflammation caused by viral infections (32). A recent study showed that fucoidan had a significant antiviral activity at 3.90–500 μg/mL concentrations, even for treating and preventing the coronavirus disease 2019 (COVID-19) (33). Ishikawa et al. investigated the anti-primary effusion lymphoma (PEL) effects of fucoidan obtained from Cladosiphon okamuranus Tokida, cultivated in Okinawa, Japan. Fucoidan dose-dependently inhibited the proliferation of KSHV-infected PEL cell lines, highlighting the anti-PEL actions of fucoidan, and the mechanism is believed to be associated with its immunomodulatory effects (34).

The compounded fucoidan product in our present study effectively reduced the Hp load in the stomach of participants (effective rate: 77.6% at 1 month and 80.5% at 2 months). It achieved Hp clearance in some carriers (clearance rate: 20.0% at 1 month and 26.8 % at 2 months), showing promising clinical efficacy in treating Hp.

Some studies implemented in China have investigated the clinical effectiveness of both STT and quadruple therapy. For the STT, the combination of pantoprazole, clarithromycin, and amoxicillin resulted in an eradication rate of 68.8% (55/80) (35) after 7 consecutive days and 80.8% (80/99) after 10 days (36); the combination of omeprazole, levofloxacin, and amoxicillin for 7 consecutive days, followed by 2 weeks of omeprazole, achieved an eradication rate of 80.2% or 83.5% when omeprazole was replaced by ecabet sodium (37). For the quadruple therapy, the combination of rabeprazole, colloidal bismuth, clarithromycin, and amoxicillin achieved an eradication rate of 92.00% (46/50); the eradication rate was 84% (42/50) when amoxicillin was replaced by metronidazole (38) or 83.67% (41/49) when rabeprazole was replaced by
omeprazole (39). In contrast, the clearance rate of Hp was slightly lower after 8 weeks of FPD consumption in our current study, and a variety of factors might explain. Participant diets were not restricted during the study period, and frequent re-infections were possible. Unlike antibiotic treatment, which has a half-life in the body and still has an antibacterial effect in the event of re-infection, FPD only has a physical adherence effect and/or an immunomodulatory activity. Since fucoidan is not an antibiotic, it is difficult to eliminate new infections between FPD consumption intervals.

As to other limitations of the study, the vast majority of the participants were asymptomatic, and some of them lacked knowledge and awareness of Hp infection, which increased the difficulty in project management and resulted in poor compliance. In addition, the custom of eating together among Chinese populations increases the risk of Hp persistence or infection. As a result, the dropout rate was high during the implementation of our study. No control group with standard practice was organized to fully compare intervention effectiveness. The follow up limited to 8 weeks.

Conclusions
In summary, FPD consumption was safe and effective at reducing Hp load on the gastric mucosa, with favorable eradication rates. More research is required to determine the long-term eradication rates, impact on Hp associated diseases and random controlled trials comparing with current standard therapy.

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Footnote

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://tgh.amegroups.com/article/view/10.21037/tgh-23-63/coif). All authors report that Blue Regale Clinical Nutrition Technology Co., Ltd. provides product support for this study. QHP is from Blue Regale Clinical Nutrition Technology Co., Ltd., Kunshan, China. The authors have no other conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. In China, there is currently no requirement for ethical approval for the use of “food” in research. So, this study did not undergo ethics committee review. All research subjects will be informed of the intervention drink before enrollment. All participants agreed to take the intervention beverage daily as per the study requirements verbally.

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