Original article

To evaluate the effect of adding licorice to the standard treatment regimen of Helicobacter pylori

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\begin{abstract}
Objective: The aim of this study was to evaluate the effect of licorice in H. pylori eradication in patients suffering from dyspepsia either with peptic ulcer disease (PUD) or non-ulcer dyspepsia (NUD) in comparison to the clarithromycin-based standard triple regimen.  
Methods: In this randomized controlled clinical trial, 120 patients who had positive rapid urease test were included and assigned to two treatment groups: control group that received a clarithromycin-based triple regimen, and study group that received licorice in addition to the clarithromycin-based regimen for two weeks. H. pylori eradication was assessed six weeks after therapy. Data was analyzed by chi-square and t-test with SPSS 16 software.  
Results: Mean ages and SD were 38.8 ± 10.9 and 40.1 ± 10.4 for the study and control groups, respectively, statistically similar. Peptic ulcer was found in 30% of both groups. Response to treatment was 83.3% and 62.5% in the study and control groups, respectively. This difference was statistically significant.  
Conclusion: Addition of licorice to the triple clarithromycin-based regimen increases H. pylori eradication, especially in the presence of peptic ulcer disease.

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\end{abstract}

Introduction

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\textit{Helicobacter pylori} (\textit{H. pylori}) is a Gram negative S shaped flagellated bacteria infecting half of mankind with a very variable prevalence depending on different factors such as race, age, and socioeconomic status ranging from a low prevalence in developed countries to a prevalence as high as 80% in developing countries like Iran.\textsuperscript{1} \textit{H. pylori} is a successful pathogen which can persistently survive in the stomach of infected persons throughout his/her life and if it results in chronic inflammation serious digestive diseases may ensue such as chronic gastritis, peptic ulcer disease (PUD), gastric cancer, and gastric MALT-oma (mucosa associated lymphoid tissue lymphoma).\textsuperscript{2}

Peptic ulcer disease is one of the consequences of \textit{H. pylori} infection and is one of the most common treatable diseases worldwide. The commonest cause of peptic ulcer is \textit{H. pylori}}
and if not eradicated it will result in 50–80% relapse in 6–12 months after treatment interruption, but if successfully treated its recurrence will drop to 6–20%, as reported by two separate meta-analyses.\(^5\,\^6\)

Relation between dyspepsia and \textit{H. pylori} is unclear, but eradication leads to symptoms improvement in 7.1% of patients.\(^6\)

American College of Gastroenterology (ACG) recommends the treatment of \textit{H. pylori} in all infected patients irrespective of the presence of ulcer. Recommended first line standard regimen by ACG for the treating \textit{H. pylori} is a triple regimen containing clarithromycin, amoxicillin or metronidazole, and proton pump inhibitor (PPI) for 10–14 days. Quadruple regimen is recommended in areas where the prevalence and resistance to clarithromycin or metronidazole is more than 20%, or if the first choice has been tried recently or repeatedly.\(^7\)

Antibiotic resistance to \textit{H. pylori} has increased in recent past years. Compared to 2010 the resistance rate to clarithromycin and amoxicillin in 2012 has raised significantly.\(^8\) The effectiveness of the present therapeutic regimen is still not satisfactory and despite several studies the best treatment regimen for treating \textit{H. pylori} remains a challenging clinical dilemma.\(^9\) Antibiotic resistance and adverse effects in addition to poor patient compliance have limited the efficacy of \textit{H. pylori} treatment. Therefore, the role of herbal medicine has been evaluated in the treatment of \textit{H. pylori}, including licorice (liquorice).\(^10\)

Licorice herb (\textit{Glycyrrhiza glabra} - \textit{G. glabra}) has routinely been used for centuries in traditional Chinese.\(^11\) Roots and rhizomes of this herb has been reported to be antioxidant, antimicrobial, and antiviral.\(^12\,\^13\) Furthermore, \textit{G. glabra} has anti-inflammatory, anticancer, and anti-ulcer activity.\(^14\,\^16\) Licorice is extracted from the root of \textit{G. glabra} herb and one of its active ingredient is glycyrrhizin acid, which due to its affinity to mineralocorticoid receptors may result in edema and hypertension. Therefore, it should be used with caution in patients suffering from cardiovascular disease and/or hypertension.\(^17\) (Fig. 1).

There are two types of action of licorice in PUD and against \textit{H. pylori}: (1) a repairing effect in PUD: protective mucosal effect by secreting a material named as secretin;\(^18\) and (2) antibacterial and anti-adhesive effect against \textit{H. pylori} by inhibiting DNA gyrase (a crucial enzyme for bacterial replication and transcription) and dihydrofolate reductase enzyme blockage.\(^19\)

In addition, the polysaccharide released from the root of licorice plays an inhibiting role in \textit{H. pylori} adhesion to gastric mucosa.\(^20\) Besides, treatment with licorice for patients suffering from dyspepsia improves their clinical complaints.\(^21\)

Owing to the rise of \textit{H. pylori} resistance against antibiotics and the effectiveness of licorice for treating \textit{H. pylori}, this study was designed to compare the efficacy of adding licorice to the standard clarithromycin-based triple regimen in the treatment of \textit{H. pylori}.

**Material and methods**

This randomized controlled clinical trial was conducted in Velayat teaching hospital in Qazvin (northwest of Iran). A total of 120 patients over 16 years of age referred for esophagogastroduodenoscopy (EGD) at the digestive disease outpatient department due to gastrointestinal complaints entered this study from May to December of 2015. Biopsy was performed when indicated by the endoscopist.

Patients with a positive rapid urease test were included in this study if none of the following conditions were present: pregnancy, use of proton pump inhibitor (PPI), bismuth or antibiotic in the past two weeks, gastrointestinal (GI) bleeding or any complication of PUD, giant ulcer, gastric ulcer, ethanol consumption or substance abuse, chronic underlying disease like diabetes mellitus, hypertension, cirrhosis, cerebrovascular attack, coronary artery disease, and gastric cancer. Patients who were reluctant to continue the treatment or complete the follow-up or suffering from a comorbidity needing non-steroidal anti-inflammatory drugs (NSAIDs) or antibiotic to be continued during the period of study were excluded. Informed consent was obtained from all eligible. Sixty patients were randomly assigned to receive one of two treatment regimens: triple regimen consisting of clarithromycin (500 mg BID) + Amoxicillin (1 gr qd) + 20 mg BID of Omeprazole (control group) or the same regimen supplemented by licorice [D-Regis (380 mg BID) made by Iran daruc Pharmacy Company] for two weeks (LR group). Both groups received at least four weeks of treatment with omeprazole 20 mg daily subsequently. Two weeks after completing the treatment, \textit{H. pylori} eradication was assessed with \textit{H. pylori} stool antigen (HPSA) test by Ag ACON Biotech (Hang Zhou) Co., Ltd.

This study is registered at Iran Registration Clinical Trial Center and was approved with the Irct registration number: IRCT2014061718124N1.

The outcome of this study was \textit{H. pylori} eradication. Data were stored at SPSS 16 software and analyzed by using of t-test, chi-square test. A \(p\)-value <0.05 was considered significant.

**Results**

Out of 120 eligible patients, 110 patients completed the study complying with the prescribed medications and follow-up;
54 patients were assigned to the licorice group (LR) and 54 patients to the control group (schematic presentation of study groups is illustrated in Fig. 2).

Demographic variables of both treatment groups were similar. Mean ages were 38.8 and 40.1 years for the LR and control groups, respectively (p = 0.56); 35 patients (51.0%) of LR group and 33 persons (48.1%) of the control group were female (p > 0.05). Patients in two groups were also similar regarding to their clinical characteristics (initial complaints and endoscopic findings). Frequency distribution of their clinical characteristics is shown in Table 1.

Furthermore, both groups had similar rates of endoscopic diagnosis of PUD or non-ulcer dyspepsia (NUD): 21 (35%) and 18 patients (30%) in the LR and control groups, respectively, had PUD (Table 2).

HPSA negative, the surrogate for H. pylori eradication and positive therapeutic response, was 83.3% in the LR group compared to 62.5% in the control group (p = 0.018). Response rate based on endoscopic findings was significantly higher for the LR group to treat PUD, but not for NUD (Table 3).

### Discussion

This study was conducted with the aim of evaluating the effectiveness of licorice in H. pylori eradication of patients suffering from dyspepsia (both PUD and NUD) compared to the clarithromycin-based standard triple regimen. Results of this study showed that adding a low dose licorice (D-Reglis 380 mg BID) to the standard triple treatment results in a more effective H. pylori eradication, especially in patients with PUD. During the recent past decades many studies on H. pylori infection have been conducted to define the adequate therapy. As H. pylori is considered a major risk factor for upper digestive problems guidelines often recommend treating all infected persons irrespective of the clinical presentation. Besides the clinical impact, H. pylori eradication would also prevent its spread.22

In the present study the addition of licorice increased the effectiveness of H. pylori eradication compared to standard clarithromycin-based triple regimen, a finding similar to previous in vitro studies.20,23
Toshio Fukai investigated the anti-\textit{H. pylori} in vitro effect of various preparations of licorice and showed that the leaves and rhizomes of this herb prevented the multiplication of \textit{H. pylori}. In addition, he also proved that licorice had anti-\textit{H. pylori} effect against the species resistant to clarithromycin.\textsuperscript{24}

Ash MK reported that \textit{G. glabra} has antimicrobial effect against several Gram positive and Gram negative bacteria including \textit{H. pylori}. Additionally, licorice is also beneficial against \textit{H. pylori} due to its anti-adhesive property.\textsuperscript{25}

Dosage of licorice has varied in few clinical studies, such as in a study by Sreenivasulu Parum et al. in which a dose of 120 mg/day was used for \textit{H. pylori} eradication in patients with dyspepsia,\textsuperscript{21} or a dose of 250 mg TID in PUD with \textit{H. pylori} infection in a study by Rahnama et al.,\textsuperscript{26} or a tablet of D-Reglis 380 mg BID in a study by Momeni et al.\textsuperscript{27}

Rahnama et al. compared the effectiveness of quadruple treatment including licorice to the same regimen using bismuth in substitution of licorice in patients with PUD. The eradication rates were 45% and 75% with licorice and bismuth, respectively, showing higher effectiveness with the regimen including licorice.\textsuperscript{26} Momeni et al. compared the same two quadruple regimens and showed similar eradication rates and concluded that D-Reglis can be a substitute for bismuth in quadruple regimen.\textsuperscript{25}

In patients with PUD our results showed significantly higher eradication rate of \textit{H. pylori} in the LR group compared with the control group (\(p = 0.03\)), in line with the report by Rahnama et al. However, the eradication rate with licorice was not different from the control group in patients with dyspepsia (\(p = 0.18\)), as found by Momeni et al. The discrepancy between the results reported by Rahnama et al. and Momeni et al. is due to the of patients studied. Rahnama et al. evaluated the \textit{H. pylori} eradication in patients with PUD whereas the study by Momeni et al. entered patients with both PUD and NUD.

Another debatable issue regarding the effect of licorice on \textit{H. pylori} eradication is the duration of treatment for improving patients' symptoms of dyspepsia. In a clinical study Raveendra et al. evaluated the effect of licorice against placebo in functional dyspepsia and found that those who received licorice for one month had a lower Nepean Dyspepsia Index and therefore showed better improvement of symptoms compared to those who received placebo.\textsuperscript{28}

In a study by Sreenivasulu Parum et al. patients suffering for dyspepsia with \textit{H. pylori} infection were prescribed licorice at a dose of 120 mg a day for 60 days. The response rate was compared against the placebo using the HPSA test on days 30 and 60; showing that HPSA test on day 60 was negative in 56% of licorice group against 4% with placebo denoting a statistically significant difference.\textsuperscript{21}

Based on above two studies it seems that licorice in dyspepsia, besides its antimicrobial nature in \textit{H. pylori} infection, has also anti-inflammatory effect. Therefore, generally speaking prescribing licorice for \textit{H. pylori} eradication at a higher dose, and longer duration should also be considered.

### Conclusion

Because of the unsatisfactory rate of \textit{H. pylori} eradication by the standard triple regimen, especially in areas with high resistance rates to clarithromycin and amoxicillin, adding licorice to the triple regimen significantly increase \textit{H. pylori} eradication in patients with PUD.

Finally, it is recommended to use licorice in addition to the triple regimen as first line treatment in order to decrease drug adverse effects and its cost, and to increase patient compliance.

### Authors' contribution

i) Substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work: Ali Akbar Hajiaghamomhamadi, Sedigheh Reisian, Sonia Oveis, Rezaoul Samimi.

ii) Drafting the work or revising it critically for important intellectual content: Ali Akbar Hajiaghamomhamadi, Sedigheh Reisian, Sonia Oveis, Ali Zargar.

iii) Final approval of the version to be published; Ali Akbar Hajiaghamomhamadi, Sedigheh Reisian, Sonia Oveis, Ali Zargar and Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy
Conflicts of interest

The authors declare no conflicts of interest.

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