Efficacy of one-day quadruple therapy for *H pylori* infection in Chinese patients

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**Abstract**

**AIM:** To compare the efficacies of one-day quadruple therapy and seven-day triple therapy in Chinese patients.

**METHODS:** Sixty consecutive patients with nonulcer dyspepsia and confirmed *H pylori* infection were randomized to receive either omeprazole 40 mg, amoxycillin 1 g, and furazolidone 100 mg, all twice a day for 7 d or omeprazole 20 mg (at breakfast and dinner), amoxicillin 1 g, furazolidone 200 mg, and colloidal bismuth subcarbonate 220 mg four times for only one day. *H pylori* status was determined before and at least 5 weeks after therapy by endoscopy with antral and corpus biopsies for rapid urease test and histology.

**RESULTS:** *H pylori* eradication was successful in 66.67% (20/30) patients in the 7-d group and in 36.67% (11/30) patients in the 1-d group (*P* = 0.037). Side effects were induced by the treatment in 13.3% (4/30) patients of each group, but these were all self-limiting, short-lasting, and did not require any specific treatment. **CONCLUSION:** The one-day quadruple therapy is less effective than the one-week regimen in curing *H pylori* infection in Chinese patients.
older than 75; (2) allergy to one of the drugs; (3) inability to attend follow-up; (4) previously failed eradication therapy; (5) treatment with antibiotics during the 4 weeks prior to the study; (6) previous ulcer surgery; and (7) liver or kidney disease, severe cardiac or pulmonary disease, alcoholism, drug abuse, or any other conditions associated with poor patient compliance. Eight patients were eventually excluded from the study after randomization. The remaining 60 patients (35 males and 25 female; mean age 54.3 years, range 19-72 years) were included in the study.

**Diagnosis of H pylori infection**
Four biopsy specimens, two from the antrum and two from the corpus, were obtained using gastroscopy. Biopsy specimens were stained to assess the presence of *H pylori*. Rapid urease test was considered positive if any color change occurred. Patients were considered as *H pylori*-positive when *H pylori* was positive on either histology and/or rapid urease test.

**Therapy**
Patients were randomized to receive a 7-d course of omeprazole (40 mg), amoxycillin (1 g), and furazolidone (100 mg), all twice daily with meals, or a 1-d course of omeprazole (20 mg, at breakfast and dinner), amoxycillin (1 g), furazolidone (200 mg), and colloidal bismuth subcitrate 220 mg four times a day. During the treatment period, the patients were asked to refer to their study doctor if they had side effects, or the side effects were recorded by means of a structured clinical interview immediately after the end of therapy.

**Assessment of H pylori eradication**
The efficacy of treatment was evaluated at least 5 weeks after therapy. Endoscopy was performed and a further four biopsy specimens (two from the antrum and two from the corpus) were obtained to assess the *H pylori* eradication status and histology. A patient was considered to be cured if all two tests were negative at the follow-up endoscopy. To evaluate possible recrudescence, a 14C-urea breath test was performed at least 6 mo after eradication therapy. The Medical Ethics Committee of the institution approved the study design. Informed consent was taken from all patients.

**Statistical analysis**
Proportions were compared using the chi-squared test. Calculations were performed using the SPSS (10.0) for Windows statistical package.

**RESULTS**
Sixty patients entered the study. Clinical characteristics of the patients are mentioned in Table 1. There were no significant differences between the 1-d quadruple therapy and 7-d triple therapy groups with respect to age, gender, tobacco and alcohol consumption.

**Eradication of H pylori**
As shown in Table 2, eradication rate in the 1-d group (36.7%) was significantly lower than that in the 7-d group (66.7%). The study was subsequently stopped. The upper gastrointestinal symptoms had improved in 93.3% (21/30) of patients in the 7-d group and in 90% (15/30) of patients in 1-d group.

**Side-effects and compliance**
As shown in Table 3, there was no difference between the incidences of side-effects in the two groups. Symptoms were almost always mild and not interfering with daily activities. Compliance was 100% in the 1-d therapy as well as in the 7-d therapy group.

**DISCUSSION**
The optimal anti-*H pylori* treatment should reach high eradication rates with only few and tolerable side-effects, short duration to assure good compliance and lower costs. Three European studies have first stated an effective one-day quadruple therapy for *H pylori* eradication with cure rates being comparable to one-week standard regimen[5-7]. Recently, Lara et al[8] showed more optimal eradication rates (95%) for the 1-d therapy group in the United States patients. One-day therapeutic protocols, to our knowledge, have not been studied in Chinese patients. An important consideration is the significantly lower cost of the 1-d regimen compared to the 7-d regimen of the standard regimen. The aim of our study was to investigate the potential advantages of the 1-d quadruple therapy by comparing to a standard 7-d triple therapy.

Unlike the previous reports[1-8], the eradication rate obtained with the 1-d therapy in our study was significantly lower than that obtained with the standard 7-d therapy. Side-effects were minor and did not interfere their daily activities.

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**Table 1** Characteristics of the treated patients

| Treatment     | n   | Male/Female | Mean age (range) | smokers (%) | alcohol users (%) |
|---------------|-----|-------------|------------------|-------------|------------------|
| 1-d therapy   | 30  | 16/14       | 38.4 (19-71)     | 6 (20)      | 7 (23.3)         |
| 7-d therapy   | 30  | 19/11       | 43.3 (18-73)     | 8 (27)      | 7 (23.3)         |

**Table 2** Evaluation of *H pylori* eradication regimens

|             | One-day therapy | Seven-day therapy | P   |
|-------------|-----------------|-------------------|-----|
| *H pylori* eradication | 36.7% (11/30) | 66.7% (28/30) | 0.037 |

**Table 3** Side effects associated with *H pylori* treatment

| Side effect (%) | One-day therapy | Seven-day therapy |
|-----------------|-----------------|-------------------|
| Dizziness       | 13.30%          | 13.30%            |
| Abdominal pain  | 1               | 1                 |
| Nausea          | 2               | 1                 |
| Skin rash       | 0               | 1                 |
| Side effect (%) | 13.30%          | 13.30%            |
activities in the both therapy groups. Compliance and antibiotic resistance are the major factors contributing to the failure of *H pylori* treatment. In our study, the compliance was reached 100% in the 1-d therapy group as well as in 7-d therapy group. However, culture was not performed in this study, so we do not know the status of *H pylori* susceptibility to metronidazole and/or furazolidone; therefore, we cannot exclude the possibility of an antibiotic resistance as the reason for our disappointing results. Nevertheless, to date, resistance to amoxycillin and furazolidone are lower as compared with clarithromycin and metronidazole. Smoking and drinking do not appear to be the factors in treatment failure with either one-day quadruple therapy or seven-day triple therapy in Chinese patients.

In conclusion, although a 1-d quadruple therapy is less expensive and has fewer side effects, due to the low efficacy obtained in this study, we do not suggest recommending this regimen to be used in Chinese patients for eradication therapy of *H pylori* infection.

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