

Clinical Trials Study

Centralized isolation of *Helicobacter pylori* from multiple centers and transport condition influences

Ya-Nan Gong, You-Ming Li, Ning-Min Yang, Hong-Zhang Li, Feng Guo, Lang Lin, Qun-Ying Wang, Jia-Kun Zhang, Zi-Zhong Ji, Ji-Bo Mao, Jun-Liang Mao, Zheng-Chao Shi, Wu-Heng Tang, Xin-Jian Zhu, Wei Shao, Xiao-Feng Zhang, Xing-Hua Wang, Yue-Feng Tong, Mi-Zu Jiang, Guang-Lan Chen, Zhi-Yong Wang, Hui-Min Tu, Guo-Fa Jiang, Jian-Sheng Wu, Xu-Peng Chen, Qiu-Long Ding, Hong Ouyang, Feng-Zhe Jin, Yan-Li Xu, Jian-Zhong Zhang

Ya-Nan Gong, Jian-Zhong Zhang, State Key Laboratory for Infectious Disease Prevention and Control, National Institute for Communicable Disease Control and Prevention, Chinese Center for Disease Control and Prevention, Beijing 102206, China
Ya-Nan Gong, Jian-Zhong Zhang, Collaborative Innovation Center for Diagnosis and Treatment of Infectious Diseases, Hangzhou 310003, Zhejiang Province, China
You-Ming Li, The First Affiliated Hospital, College of Medicine, Zhejiang University, Hangzhou 310003, Zhejiang Province, China
Ning-Min Yang, Zhiyuan Medical Inspection Institute Co., Ltd., Hangzhou 310021, Zhejiang Province, China
Hong-Zhang Li, Sanmen People's Hospital, Taizhou 317100, Zhejiang Province, China
Feng Guo, The First People's Hospital of Xiaoshan District, Hangzhou 311200, Zhejiang Province, China
Lang Lin, The First People's Hospital of Cangnan, Wenzhou 325800, Zhejiang Province, China
Qun-Ying Wang, Jinhua Municipal Central Hospital, Jinhua 321001, Zhejiang Province, China
Jia-Kun Zhang, The First People's Hospital of Pingyang, Wenzhou 325400, Zhejiang Province, China
Zi-Zhong Ji, The First Hospital of Jiaying, Jiaying 314001, Zhejiang Province, China
Ji-Bo Mao, Zhoushan Hospital, Zhoushan 316021, Zhejiang Province, China
Jun-Liang Mao, The First People's Hospital of Wenling, Wenling 317500, Zhejiang Province, China
Zheng-Chao Shi, Rui'an People's Hospital, Rui'an 325200, Zhejiang Province, China
Wu-Heng Tang, Maternal and Child Health Hospital of Zhoushan City, Zhoushan 316000, Zhejiang Province, China
Xin-Jian Zhu, Shangyu People's Hospital, Shaoxing 312300, Zhejiang Province, China
Wei Shao, People's Hospital of Putuo District, Zhoushan 316399, Zhejiang Province, China
Xiao-Feng Zhang, Hangzhou First People's Hospital, Hangzhou 310006, Zhejiang Province, China

Xing-Hua Wang, Qingtian People's Hospital, Lishui 323900, Zhejiang Province, China
Yue-Feng Tong, The First People's Hospital of Yongkang, Yongkang 321300, Zhejiang Province, China
Mi-Zu Jiang, The Children's Hospital Zhejiang University School of Medicine, Hangzhou 310003, Zhejiang Province, China
Guang-Lan Chen, Lishui People's Hospital, Lishui 323000, Zhejiang Province, China
Zhi-Yong Wang, The Affiliated Hospital of Hangzhou Normal University, Hangzhou 310015, Zhejiang Province, China
Hui-Min Tu, Wuxi No. 4 Hospital Affiliated to Suzhou University, Wuxi 214062, Zhejiang Province, China
Guo-Fa Jiang, Jinhua Wenrong Hospital, Jinhua 321001, Zhejiang Province, China
Jian-Sheng Wu, The First Affiliated Hospital of Wenzhou Medical College, Wenzhou 325000, Zhejiang Province, China
Xu-Peng Chen, Yueqing People's Hospital, Wenzhou 325600, Zhejiang Province, China
Qiu-Long Ding, Tiantai People's Hospital, Taizhou 317200, Zhejiang Province, China
Hong Ouyang, Lin'an people's Hospital, Lin'an 311300, Zhejiang Province, China
Feng-Zhe Jin, Rui'an City TCM Hospital, Rui'an 325200, Zhejiang Province, China
Yan-Li Xu, Medical College of Hebei University of Engineering, Handan 056038, Hebei Province, China
Author contributions: Gong YN and Xu YL analyzed data and drafted this paper; Li YM and Yang NM were involved in designing the protocol and providing results of this study; Zhang JZ had the idea for this study and final responsibility for the decision to submit for publication; Li HZ, Guo F, Lin L, Wang QY, Zhang JK, Ji ZZ, Mao JB, Mao JL, Shi ZC, Tang WH, Zhu XJ, Shao W, Zhang XF, Wang XH, Tong YF, Jiang MZ, Chen GL, Wang ZY, Tu HM, Jiang GF, Wu JS, Chen XP, Ding QL, Ouyang H and Jin FZ were responsible for sample collection, the order of authorship was based on specific contribution; all authors were involved in data interpretation and critical revisions of the

report, and approved the final version.

Supported by Grants from the Science and Technology Program of Zhejiang Province China, No. 2001C23140, National Technology RD Program in the 12th Five-Year Plan of China, No. 2012BAI06B02, the Major Technology Project as part of "Prevention and Control of Major Infectious Diseases including AIDS and Viral Hepatitis", No. 2013ZX10004216-002, the National Key Scientific Instrument and Equipment Development Project, No. 2012YQ180117, the Medical and Health Science and Technology Plan Project of Zhejiang Province, No. 2012KYB248, and the Science and Technology Project of Zhejiang province, No. 2011C23140.

Open-Access: This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Correspondence to: Jian-Zhong Zhang, MD, PhD, State Key Laboratory for Infectious Disease Prevention and Control, National Institute for Communicable Disease Control and Prevention, Chinese Center for Disease Control and Prevention, 155 Changbai Road, Changping District, Beijing 102206, China. zhangjianzhong@icdc.cn

Telephone: +86-10-58900707

Fax: +86-10-58900700

Received: June 16, 2014

Peer-review started: June 16, 2014

First decision: July 21, 2014

Revised: August 11, 2014

Accepted: September 18, 2014

Article in press: September 19, 2014

Published online: January 21, 2015

Abstract

AIM: To evaluate the efficacy of centralized culture and possible influencing factors.

METHODS: From January 2010 to July 2012, 66452 patients with suspected *Helicobacter pylori* (*H. pylori*) infection from 26 hospitals in Zhejiang and Jiangsu Provinces in China underwent gastrointestinal endoscopy. Gastric mucosal biopsies were taken from the antrum for culture. These biopsies were transported under natural environmental temperature to the central laboratory in Hangzhou city and divided into three groups based on their transport time: 5, 24 and 48 h. The culture results were reported after 72 h and the positive culture rates were analyzed by a χ^2 test. An additional 5736 biopsies from *H. pylori*-positive patients (5646 rapid urease test-positive and 90 ¹⁴C-urease breath test-positive) were also cultured for quality control in the central laboratory setting.

RESULTS: The positive culture rate was 31.66% (21036/66452) for the patient samples and 71.72% (4114/5736) for the *H. pylori*-positive quality control specimens. In the 5 h transport group, the positive

culture rate was 30.99% (3865/12471), and 32.84% (14960/45553) in the 24 h transport group. In contrast, the positive culture rate declined significantly in the 48 h transport group (26.25%; $P < 0.001$). During transportation, the average natural temperature increased from 4.67 to 29.14 °C, while the positive culture rate declined from 36.67% (1462/3987) to 24.12% (1799/7459). When the temperature exceeded 24 °C, the positive culture rate decreased significantly, especially in the 48 h transport group (23.17%).

CONCLUSION: Transportation of specimens within 24 h and below 24 °C is reasonable and acceptable for centralized culture of multicenter *H. pylori* samples.

Key words: Centralized isolation; *Helicobacter pylori*; Influencing factor; Multiple centers; Personalized treatment

© The Author(s) 2015. Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: This is the first large-scale study on the centralized culture of *Helicobacter pylori* in a large number of clinical samples from multiple centers. The efficacy of centralized culture and possible influencing factors were evaluated. The results confirm the feasibility of establishing a culture center for individualized medical use. The findings of this study can be promisingly applied in clinical and public health practice.

Gong YN, Li YM, Yang NM, Li HZ, Guo F, Lin L, Wang QY, Zhang JK, Ji ZZ, Mao JB, Mao JL, Shi ZC, Tang WH, Zhu XJ, Shao W, Zhang XF, Wang XH, Tong YF, Jiang MZ, Chen GL, Wang ZY, Tu HM, Jiang GF, Wu JS, Chen XP, Ding QL, Ouyang H, Jin FZ, Xu YL, Zhang JZ. Centralized isolation of *Helicobacter pylori* from multiple centers and transport condition influences. *World J Gastroenterol* 2015; 21(3): 944-952 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v21/i3/944.htm> DOI: <http://dx.doi.org/10.3748/wjg.v21.i3.944>

INTRODUCTION

Helicobacter pylori (*H. pylori*), one of the most common human pathogens, can lead to gastric ulcers, gastritis, gastric cancer, and mucosa-associated lymphoid tumors^[1]. According to the Maastricht IV and Chinese Consensus Report, two antibiotics combined with one proton-pump inhibitor are recommended as standard first-line treatment to eradicate *H. pylori*^[2,3]. However, in recent years, the success rates have declined below 80% in most European and Asian countries^[4-6]. There are many reasons accounting for eradication failure, such as poor patient compliance, low gastric pH, and resistant bacteria. In fact, the main reason for the decline is the increasing *H. pylori* resistance to the antibiotics used^[7,8]. In China, the resistance to clarithromycin, a key antibiotic in the triple

therapy, has reached above 20% and should not be used in anti-*H. pylori* therapy without a susceptibility test^[2,9-11].

Accumulated evidence suggests that culture-susceptibility tests may improve the eradication rate of *H. pylori*^[12-17]. However, *H. pylori* is a rather fastidious bacterium at culture, especially when a low bacterial load is present^[18]. The ideal situation for culture-susceptibility test is not available in many clinical settings, since most endoscopic units do not have direct access to a microbiology laboratory. Therefore, gastric biopsy specimens should be transported to a central laboratory for *H. pylori* culture. During transportation, time and temperature could influence the survivability of *H. pylori* and these issues are still in debate^[19-23]. Some investigators emphasized the need for rapid transport at a low temperature^[21]. Others demonstrated that *H. pylori* could survive at room temperature for 24 h without loss of the ability to recover^[22]. The transport of samples at natural temperature is required for a routine clinical application system. In order to assess the factors, transportation^[24] and isolation tests of thousands of *H. pylori* strains were performed in our lab, but the complexity of large-scale clinical application could not be accurately represented. The feasibility of centralized isolation of *H. pylori* in a large number of clinical samples from multiple centers and influencing factors need to be tested for actual practice.

A central laboratory for *H. pylori* isolation was set up in Zhiyuan Medical Inspection Institute Co., Ltd., in Hangzhou city, which has provided a personalized treatment strategy for *H. pylori* eradication in recent years. In order to evaluate the efficacy of centralized culture and possible influencing factors, we conducted research in the central laboratory to analyze positive culture rates of a large number of clinical samples collected from 26 hospitals in nine cities.

MATERIALS AND METHODS

Sample collection

The sampling was conducted in Zhejiang Province and Jiangsu Province, China. Between January 2010 and July 2012, consecutive participants with suspected *H. pylori* infection underwent gastrointestinal endoscopy. The gastric mucosal biopsies were taken from the greater curvature of gastric antrum using sterile disposable biopsy forceps. Additionally, in order to assess the positive rate of *H. pylori* culture, specimens from *H. pylori*-positive patients (5646 rapid urease test-positive and 90 ¹⁴C-urease breath test-positive) were also collected as a control. The *H. pylori*-positive samples were all collected from Wenzhou city, including The First Affiliated Hospital of Wenzhou Medical College and The First People's Hospital of Pingyang.

This study was approved by the Ethics Committee of National Institute for Communicable Disease Control and Prevention, Chinese Center for Disease Control and Prevention and the Ethics Committee of Zhiyuan

Medical Inspection Institute Co., Ltd., and written informed consent was obtained from all patients.

Transport and culture procedure

The gastric mucosal sample from each patient was stored in sterile tube that contained 1 mL brain-heart infusion with 20% glycerol and kept under 4 °C before transportation. All the fresh biopsy specimens were transported at external temperature to Zhiyuan Medical Inspection Institute Co., Ltd., for centralized isolation. The 66452 specimens were divided into three independent groups based on transport time: 5, 24 and 48 h. The first group (5 h; *n* = 12471) was collected from Hangzhou city between 9 am and 12 pm and cultured at 2 pm on the same day. The second group (24 h; *n* = 45553) was collected from other cities hundreds of kilometers away from Hangzhou between 9 am and 12 pm and sent to Hangzhou for culture at 9 am the next day. The third group (48 h; *n* = 8428) were collected from Jinhua city and transported to Hangzhou for culture 48 h later because of the limitations of local hospital conditions. The *H. pylori*-positive control samples were transported the same as the 24 h group.

The samples were ground/broken and cultivated on Columbia agar plates (Oxoid of Thermo Fisher Scientific Inc., Waltham, MA, United States) supplemented with 5% defibrinated sheep blood, 3 µg/mL synergist, 2.5 µg/mL vancomycin, 2 µg/mL amphotericin B, and 2 µg/mL bacillosporin B under microaerophilic conditions (5% O₂, 10% CO₂, 85% N₂) for 72 h. Translucent colonies (0.5-2.0 mm) from the original agar plates were selected for gram staining and urease, oxidase and catalase tests. Colonies with curved gram-negative rods resembling *Helicobacter* spp and positive in three enzyme tests were identified as *H. pylori*.

Meteorological data

The information concerning daily measured maximum, minimum and average temperatures in each city related with this study in Zhejiang Province from January 1, 2010 to December 31, 2012 was supplied by the Meteorological Bureau of Zhejiang Province. The average daily temperature was the average value of temperatures at four time points each day (2 am, 8 am, 2 pm and 8 pm). In our investigation, we used the regional level meteorological data from six cities (Hangzhou, Jiaying, Jinhua, Taizhou, Wenzhou, and Zhoushan) to calculate the monthly maximum, minimum and average temperatures and draw the temperature change curves.

Statistical analysis

The results of isolation and identification of *H. pylori* for all specimens were reported 72 h after culture and all calculations were performed using SPSS 18.0 software (SPSS Inc., Chicago, IL, United States). The positive culture rates were assessed with a χ^2 test. A *P* ≤ 0.05 was considered as statistically significant.

RESULTS

Sample collection

The random 66452 samples were collected from 24 hospitals in nine cities: Hangzhou ($n = 6$), Jiaxing ($n = 1$), Jinhua ($n = 3$), Shaoxing ($n = 1$), Taizhou ($n = 3$), Wenzhou ($n = 4$), Zhoushan ($n = 3$), Lishui ($n = 2$) and Wuxi ($n = 1$). The 5736 *H. pylori*-positive specimens were collected from another two hospitals in Wenzhou city. Eight of the 26 sampling hospitals are Class 2A hospitals, which are primarily county-level hospitals providing medical service to the communities. The other hospitals belong to Class 3A, which are regional hospitals and possess the highest medical level in Chinese hospital grading. The distribution and numbers of samples in each year and city are shown in Figure 1A.

Culture results

Of the 66452 specimens, 31.66% (21036/66452) were *H. pylori*-positive. The positive culture rates of specimens in 2010, 2011 and 2012 were 28.64% (4897/17098), 30.28% (6513/21509) and 34.57% (9626/27845), respectively. The differences showed statistical significance ($P < 0.001$). The positive culture rate for Class 2A hospitals was 31.96% (8103/25350) and that of Class 3A hospitals was 31.47% (12933/41102).

The positive culture rates of samples from nine cities varied from 26.59% (2354/8852) in Jinhua city to 42.51% (332/781) in Lishui city (Figure 1B). The positive culture rate of Hangzhou city, which was the culture center, was 30.99% (3866/12473).

Even in the same city, the positive culture rates of different sampling hospitals were different. The culture positive rates from four hospitals in Hangzhou city were 29.97% (3316/11064), 38.95% (298/705), 25.73% (79/307) and 47.66% (61/128) (Figure 2).

Culture results of different transport groups

The positive culture rates from the 5, 24, and 48 h transport groups were 30.99% (3865/12471), 32.84% (14960/45553) and 26.25% (2211/8428), respectively. The positive culture rate of the 48 h transport group was significantly lower than the other two groups ($P < 0.001$). The positive culture rate from the 48 h group was significantly lower than the 24 h group in samples from the same city, (26.25% *vs* 33.73%, $P < 0.001$).

Culture results from natural transport temperatures

The average natural temperature in Zhejiang province varied within one year from 4.67 to 29.14 °C. The average temperatures in January, May, June and July were 4.66, 20.85, 24.24 and 28.85 °C, respectively. The change curves of average temperatures were similar in six cities, and the monthly average temperature, average maximum and minimum temperatures of the six cities during this study are shown in Figure 3A.

As the temperature increased, the total positive culture rate declined from 36.67% (1462/3987) in December

to 24.12% (1799/7459) in July. Although the average temperature elevated significantly from January to May, the positive culture rates were similar. When the average temperature exceeded 24 °C in June and July, the positive culture rates declined significantly ($P < 0.001$), with the lowest positive culture rate in July (24.12%; 1799/7459). This phenomenon was particularly evident in Wenzhou city whose positive culture rates changed from 38.44% (316/822) in January to 25.38% (389/1533) in July. The positive culture rates in the other months changed slightly even when the temperatures changed significantly (Figure 3B). Positive culture rates of six cities in Zhejiang Province in January, May, June and July are shown in Figure 4.

Culture results of *H. pylori*-positive specimens

Of the *H. pylori*-positive specimens, the isolation of *H. pylori* was successful in 71.72% (4114/5736) of cases, demonstrating the positive predictive value of *H. pylori* isolation in the central cultural platform.

DISCUSSION

Isolation of all samples was conducted in the same laboratory to eliminate the intrinsic variability from multiple laboratories. The advantage of our study is that it contains largest sample size to date, involving a wide range of demographic characteristics, geographic areas and hospitals from various medical levels (nearly all the gastrointestinal endoscopic tests were done in Class 2A and 3A hospitals in China). Compared with a previous study^[25], the limitations of small sample size and collection of data from different laboratories were certainly avoided in this study, and more reliable information for the centralized culture of *H. pylori* was provided.

Previous studies reported *H. pylori*-positive isolation rates ranging from 75% to 94%^[25,26]. Considering the differences between the *H. pylori* cultures used for clinical study and those for individual medical service, as well as the amount of samples and the time-efficient characteristics (a limited time of about seven days for clinical report was usually set in the latter), the positive culture rate of 71.72% found in this study is acceptable.

In China, the total infectious rate of *H. pylori* was relatively high, reaching 40%-60% in adults. In this study, the positive culture rate of the suspected *H. pylori* infection group was 31.66%. Combined with the culture rate of positive control samples, a 44% infectious rate was calculated, in agreement with the reported data, indicating the positive culture rate of random samples was reasonable. In addition, the cost of culture and sensitivity tests for six antibiotics is approximately \$18, whereas the urease breath test costs \$29 in Zhejiang province. The random samples were collected without any *H. pylori* detection, leading to a lower cost (51.25% reduced) and economic burden in this population compared with the detection strategy.

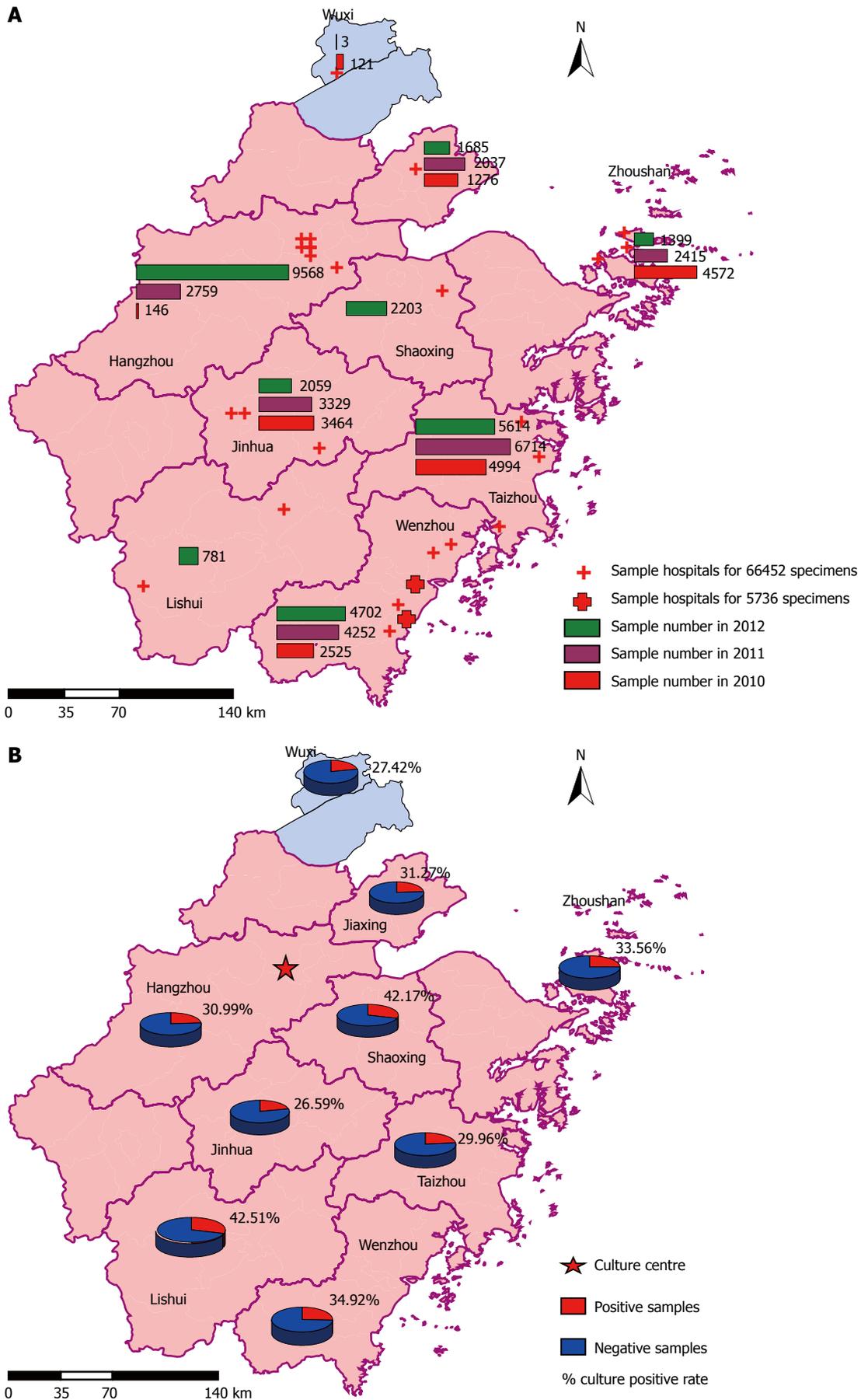


Figure 1 Distributions of samples and *Helicobacter pylori* positive culture rates. A: Distribution of the 26 hospitals and sample numbers from 2010 to 2012 in each city; B: Positive rates in nine cities in Zhejiang and Jiangsu Provinces. The map was drawn by ARCGIS 9.3. software.

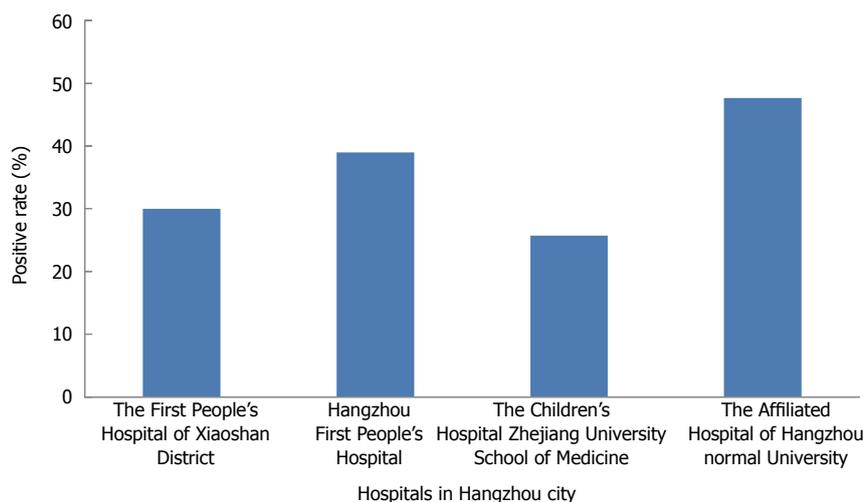


Figure 2 Positive culture rates in four different hospitals in Hangzhou.

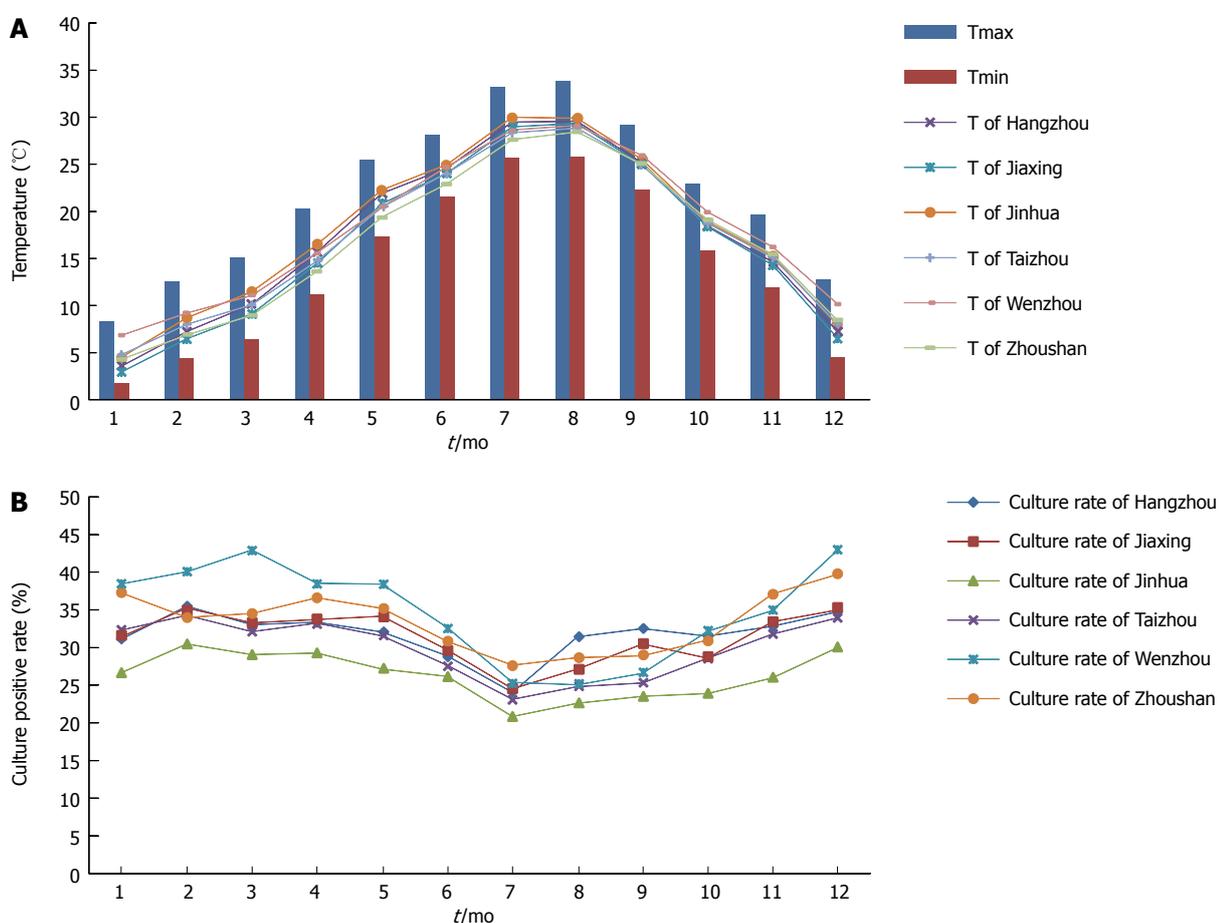


Figure 3 Temperatures and positive culture rate in six cities within three years. A: The maximum (T_{max}), minimum (T_{min}) and average (T) temperatures; B: Positive culture rates.

Factors influencing the isolation rates during transportation have been discussed by previous studies^[18,23,27,28]. Use of dry ice and need for storage at a constant temperature of 4 °C were recommended, but they cannot be put into actual practice in the *H. pylori* isolation for individualized medical need in a large number of cases because dry ice is too costly and constant 4 °C preservation condition is hard to maintain. Despite the appearance as a necessary condition

for the optimal diagnosis of *Helicobacter* infections *via* bacteriologic methods in a small-sample study^[29], transport on ice was not recommended, as repeated freezing and thawing is more harmful for the successful isolation of *H. pylori*. Thus, in our study, transportation of biopsy specimens under ambient temperature was performed. Our results indicate that the samples were maintained without influencing the culture rates within 24 h, especially

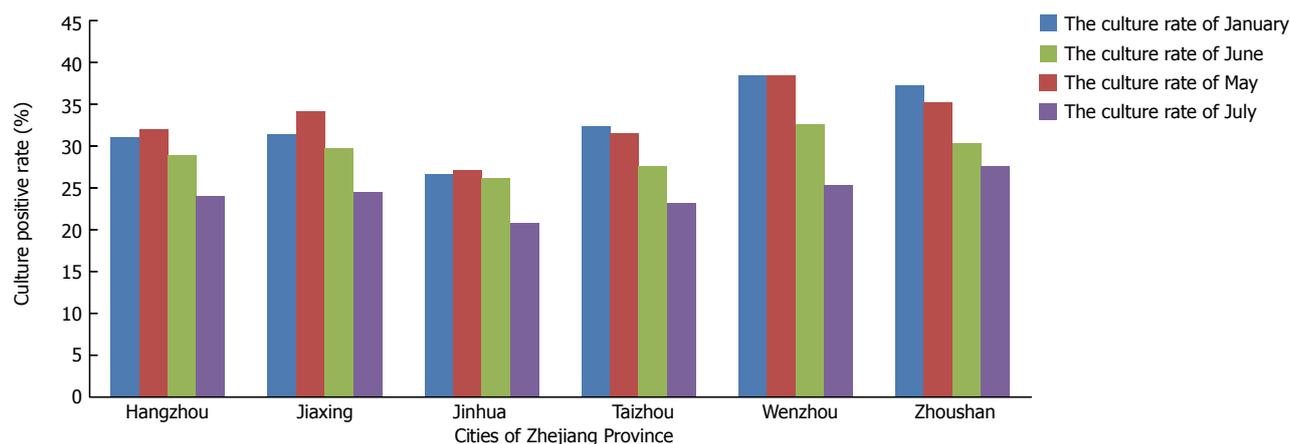


Figure 4 Positive culture rates of six cities in Zhejiang Province in January, May, June and July.

when the mean temperature was lower than 24 °C. A prolonged or delayed transport time and higher natural temperature (daily average temperature above 24 °C) could significantly influence the positive culture rates, probably due to contamination and overgrowth of other bacteria. Based on these factors, the proper position of the culture center should be taken into account, and measures to lower ambient temperature should be taken in summer.

Besides the two strong influencing factors, other factors such as sampling and operation levels should also be considered, since variability in the positive culture rates was observed among the four hospitals in Hangzhou city that shared the same sample population and transport conditions. In one study^[30], low isolation rates were obtained from failed eradication therapy patients. The patients from Class 2A and 3A hospitals might have different backgrounds of antibiotics use. In this study, a novel design was to compare the culture rates between Class 2A and Class 3A hospitals. The results show no obvious differences, indicating the medical levels of hospitals can be ignored. In addition, the positive culture rates significantly increased over three years, probably due to the improved sampling skills and experience of the laboratory personnel. This indicates that the positive culture rate may increase, even within an *H. pylori*-isolation platform for individualized medical need in a large number of cases.

There were also some limitations in this study. First, we only compared the positive culture rates of four hospitals in Hangzhou city. The other two hospitals were excluded, as the number of samples was small and the sampling was not consecutive. Second, we only selected six cities to assess the temperature effects on positive culture rates, because the samples collected in Lishui, Wuxi and Shaoxing were missed for some months, and the meteorological data was incomplete. Even with these exclusions, the selected data accounted for the majority and was sufficient to compare these differences. Third, although the *H. pylori*-positive group was available to assess the culture results, there was no direct comparison or control group in this study for all samples, such as a

non-culture based test, and the positive culture rates in this study were considered acceptable.

In general, this was the first large-scale study on centralized culture of *H. pylori*, confirming the feasibility of establishing a culture center for individualized medical use. We believe the findings of this study can be promisingly applied in clinical and public health practice. The resistance to clarithromycin often poses a challenge in clinical design. For example, as the eradication rate of two couplet therapeutics decreased ten years ago, triple therapy was recommended, thus doctors had to choose quadruple or sequential therapy in failed cases. However, this cycle is likely to continue, and the eradication rate of quadruple or sequential therapies will be reduced to unacceptable levels. Establishment of a personalized treatment strategy may potentially resolve the public health problem and reduce economic burden on the patients and the community, especially in those populations with high resistance.

COMMENTS

Background

The centralized culture of *Helicobacter pylori* (*H. pylori*) in a large number of clinical samples from multiple centers was established. The efficacy of centralized culture and possible influencing factors was evaluated.

Research frontiers

Antibiotic resistance is a worldwide problem that prevents *H. pylori* eradication. Personalized treatment strategy based on culture and antimicrobial susceptibility tests is one of the most promising ways to solve the problem.

Innovations and breakthroughs

Previous studies have discussed factors influencing *H. pylori* isolation rates during transportation. This is the first large-scale study on centralized culture of *H. pylori*, confirming the feasibility of establishing a culture center for individualized medical use.

Applications

The findings of this study can be promisingly applied in clinical and public health practice. Establishment of a personalized treatment strategy may potentially resolve the public health problem and reduce economic burden on the patients and the community, especially in those populations with high resistance.

Terminology

Clarithromycin is one of the core antibiotics of a triple regimen for *H. pylori* eradication. If the clarithromycin resistance rate of *H. pylori* increases to 15%-20%

in a population, this antibiotic should not be used without a susceptibility test.

Peer review

In this study, the authors evaluated the efficacy of centralized culture and discussed the possible influencing factors. It reveals that centralized culture of multicenter samples is feasible. The results of this study can be promisingly applied in individualized medical use and clinical *H. pylori* eradication.

REFERENCES

- 1 **Parsonnet J.** Helicobacter pylori: the size of the problem. *Gut* 1998; **43** Suppl 1: S6-S9 [PMID: 9764031 DOI: 10.1136/gut.43.2008.S6]
- 2 **Malfertheiner P,** Megraud F, O'Morain CA, Atherton J, Axon AT, Bazzoli F, Gensini GF, Gisbert JP, Graham DY, Rokkas T, El-Omar EM, Kuipers EJ. Management of Helicobacter pylori infection--the Maastricht IV/ Florence Consensus Report. *Gut* 2012; **61**: 646-664 [PMID: 22491499 DOI: 10.1136/gutjnl-2012-302084]
- 3 **Hu FL,** Hu PJ, Liu WZ, De Wang J, Lv NH, Xiao SD, Zhang WD, Cheng H, Xie Y. Third Chinese National Consensus Report on the management of Helicobacter pylori infection. *J Dig Dis* 2008; **9**: 178-184 [PMID: 18956598 DOI: 10.1111/j.1751-2980.2008.00342.x]
- 4 **Altintas E,** Sezgin O, Ulu O, Aydin O, Camdeviren H. Maastricht II treatment scheme and efficacy of different proton pump inhibitors in eradicating Helicobacter pylori. *World J Gastroenterol* 2004; **10**: 1656-1658 [PMID: 15162544]
- 5 **Gumurdulu Y,** Serin E, Ozer B, Kayaselcuk F, Ozsahin K, Cosar AM, Gursoy M, Gur G, Yilmaz U, Boyacioglu S. Low eradication rate of Helicobacter pylori with triple 7-14 days and quadruple therapy in Turkey. *World J Gastroenterol* 2004; **10**: 668-671 [PMID: 14991935]
- 6 **Suzuki H,** Nishizawa T, Hibi T. Helicobacter pylori eradication therapy. *Future Microbiol* 2010; **5**: 639-648 [PMID: 20353303 DOI: 10.2217/fmb.10.25]
- 7 **Mégraud F,** Lamouliatte H. Review article: the treatment of refractory Helicobacter pylori infection. *Aliment Pharmacol Ther* 2003; **17**: 1333-1343 [PMID: 12786627 DOI: 10.1046/j.1365-2036.2003.01592.x]
- 8 **Mégraud F.** H pylori antibiotic resistance: prevalence, importance, and advances in testing. *Gut* 2004; **53**: 1374-1384 [PMID: 15306603 DOI: 10.1136/gut.2003.022111]
- 9 **Gao W,** Cheng H, Hu F, Li J, Wang L, Yang G, Xu L, Zheng X. The evolution of Helicobacter pylori antibiotics resistance over 10 years in Beijing, China. *Helicobacter* 2010; **15**: 460-466 [PMID: 21083752 DOI: 10.1111/j.1523-5378.2010.00788.x]
- 10 **Su P,** Li Y, Li H, Zhang J, Lin L, Wang Q, Guo F, Ji Z, Mao J, Tang W, Shi Z, Shao W, Mao J, Zhu X, Zhang X, Tong Y, Tu H, Jiang M, Wang Z, Jin F, Yang N, Zhang J. Antibiotic resistance of Helicobacter pylori isolated in the Southeast Coastal Region of China. *Helicobacter* 2013; **18**: 274-279 [PMID: 23418857 DOI: 10.1111/hel.12046]
- 11 **Malfertheiner P,** Megraud F, O'Morain C, Bazzoli F, El-Omar E, Graham D, Hunt R, Rokkas T, Vakil N, Kuipers EJ. Current concepts in the management of Helicobacter pylori infection: the Maastricht III Consensus Report. *Gut* 2007; **56**: 772-781 [PMID: 17170018 DOI: 10.1136/gut.2006.101634]
- 12 **Dore MP,** Leandro G, Realdi G, Sepulveda AR, Graham DY. Effect of pretreatment antibiotic resistance to metronidazole and clarithromycin on outcome of Helicobacter pylori therapy: a meta-analytical approach. *Dig Dis Sci* 2000; **45**: 68-76 [PMID: 10695616]
- 13 **Realdi G,** Dore MP, Piana A, Atzei A, Carta M, Cugia L, Manca A, Are BM, Massarelli G, Mura I, Maida A, Graham DY. Pretreatment antibiotic resistance in Helicobacter pylori infection: results of three randomized controlled studies. *Helicobacter* 1999; **4**: 106-112 [PMID: 10382124 DOI: 10.1046/j.1523-5378.1999.99002.x]
- 14 **Glupczynski Y,** Mégraud F, Lopez-Brea M, Andersen LP. European multicentre survey of in vitro antimicrobial resistance in Helicobacter pylori. *Eur J Clin Microbiol Infect Dis* 2001; **20**: 820-823 [PMID: 11783701 DOI: 10.1007/s100960100611]
- 15 **Wang G,** Zhao Q, Li S. Study of drug sensitivity test in Helicobacter pylori eradication therapy. *J Clin Intern Med* 2008; **25**: 474-477
- 16 **Romano M,** Iovene MR, Montella F, Vitale LM, De S : imone T, Del Vecchio Blanco C. Pretreatment antimicrobial-susceptibility testing in the eradication of H. pylori infection. *Am J Gastroenterol* 2000; **95**: 3317-3318 [PMID: 11095372 DOI: 10.1111/j.1572-0241.2000.03317.x]
- 17 **Romano M,** Marmo R, Cuomo A, De Simone T, Mucherino C, Iovene MR, Montella F, Tufano MA, Del Vecchio Blanco C, Nardone G. Pretreatment antimicrobial susceptibility testing is cost saving in the eradication of Helicobacter pylori. *Clin Gastroenterol Hepatol* 2003; **1**: 273-278 [PMID: 15017668 DOI: 10.1016/S1542-3565(03)00131-9]
- 18 **Soltész V,** Zeeberg B, Wadström T. Optimal survival of Helicobacter pylori under various transport conditions. *J Clin Microbiol* 1992; **30**: 1453-1456 [PMID: 1624562]
- 19 **Yuen B,** Zbinden R, Fried M, Bauerfeind P, Bernardi M. Cultural recovery and determination of antimicrobial susceptibility in Helicobacter pylori by using commercial transport and isolation media. *Infection* 2005; **33**: 77-81 [PMID: 15827875 DOI: 10.1007/s15010-005-4071-y]
- 20 **Xia HX,** Keane CT, O'Morain CA. Determination of the optimal transport system for Helicobacter pylori cultures. *J Med Microbiol* 1993; **39**: 334-337 [PMID: 8246249 DOI: 10.1099/00222615-39-5-334]
- 21 **Roosendaal R,** Kuipers EJ, Peña AS, de Graaff J. Recovery of Helicobacter pylori from gastric biopsy specimens is not dependent on the transport medium used. *J Clin Microbiol* 1995; **33**: 2798-2800 [PMID: 8567932]
- 22 **Heep M,** Scheibl K, Degrell A, Lehn N. Transport and storage of fresh and frozen gastric biopsy specimens for optimal recovery of Helicobacter pylori. *J Clin Microbiol* 1999; **37**: 3764-3766 [PMID: 10523597]
- 23 **Siu LK,** Leung WK, Cheng AF, Sung JY, Ling TK, Ling JM, Ng EK, Lau JY, Chung SC. Evaluation of a selective transport medium for gastric biopsy specimens to be cultured for Helicobacter pylori. *J Clin Microbiol* 1998; **36**: 3048-3050 [PMID: 9738066]
- 24 **Xia HX,** Keane CT, Chen J, Zhang J, Walsh EJ, Moran AP, Hua JS, Megraud F, O'Morain CA. Transportation of Helicobacter pylori cultures by optimal systems. *J Clin Microbiol* 1994; **32**: 3075-3077 [PMID: 7883907]
- 25 **Grove DI,** McLeay RA, Byron KE, Koutsouridis G. Isolation of Helicobacter pylori after transport from a regional laboratory of gastric biopsy specimens in saline, Portagerm pylori or cultured on chocolate agar. *Pathology* 2001; **33**: 362-364 [PMID: 11523941]
- 26 **Debonnie JC,** Delmee M, Mainguet P, Beyaert C, Haot J, Legros G. Cytology: a simple, rapid, sensitive method in the diagnosis of Helicobacter pylori. *Am J Gastroenterol* 1992; **87**: 20-23 [PMID: 1728119]
- 27 **van der Hulst RW,** Verheul SB, Weel JF, Gerrits Y, ten Kate FJ, Dankert J, Tytgat GN. Effect of specimen collection techniques, transport media, and incubation of cultures on the detection rate of Helicobacter pylori. *Eur J Clin Microbiol Infect Dis* 1996; **15**: 211-215 [PMID: 8740855 DOI: 10.1007/BF01591356]
- 28 **Veenendaal RA,** Lichtendahl-Bernards AT, Peña AS, Endtz HP, van Boven CP, Lamers CB. Effect of transport medium and transportation time on culture of Helicobacter pylori from gastric biopsy specimens. *J Clin Pathol* 1993; **46**: 561-563 [PMID: 8331182 DOI: 10.1136/jcp.46.6.561]
- 29 **Meunier O,** Walter P, Chamouard P, Piemont Y, Monteil H. [Isolation of Helicobacter pylori: necessity of control

- of transport conditions]. *Pathol Biol (Paris)* 1997; **45**: 82-85 [PMID: 9097852]
- 30 **Savarino V**, Zentilin P, Pivari M, Bisso G, Raffaella Mele M, Bilardi C, Borro P, Dulbecco P, Tessieri L, Mansi C,

Borgonovo G, De Salvo L, Vigneri S. The impact of antibiotic resistance on the efficacy of three 7-day regimens against *Helicobacter pylori*. *Aliment Pharmacol Ther* 2000; **14**: 893-900 [PMID: 10886045 DOI: 10.1046/j.1365-2036.2000.00780.x]

P- Reviewer: Buzas GM, Yula E **S- Editor:** Qi Y
L- Editor: AmEditor **E- Editor:** Zhang DN





Published by **Baishideng Publishing Group Inc**

8226 Regency Drive, Pleasanton, CA 94588, USA

Telephone: +1-925-223-8242

Fax: +1-925-223-8243

E-mail: bpgoffice@wjgnet.com

Help Desk: <http://www.wjgnet.com/esps/helpdesk.aspx>

<http://www.wjgnet.com>



ISSN 1007-9327

