RESEARCH ARTICLE

Effects of the CYP2C19 Genetic Polymorphism on Gastritis, Peptic Ulcer Disease, Peptic Ulcer Bleeding and Gastric Cancer

Wannapa Jainan¹, Ratha-Korn Vilaichone¹, ²

Abstract

Background: The CYP2C19 genotype has been found to be an important factor for peptic ulcer healing and H. pylori eradication, influencing the efficacy of proton pump inhibitors (PPIs) and the pathogenesis of gastric cancer. The aim of this study was to investigate clinical correlations of the CYP2C19 genotype in patients with gastritis, peptic ulcer disease (PUD), peptic ulcer bleeding (PUB) and gastric cancer in Thailand. Materials and Methods: Clinical information, endoscopic findings and H. pylori infection status of patients were assessed between May 2012 and November 2014 in Thammasat University Hospital, Thailand. Upper GI endoscopy was performed for all patients. Five milliliters of blood were collected for H. pylori serological diagnosis and CYP2C19 study. CYP2C19 genotypes were determined by polymerase chain reaction (PCR) and restriction fragment length polymorphism analysis (RFLP) and classified as rapid metabolizer (RM), intermediate metabolizer (IM) or poor metabolizer (PM). Results: A total of 202 patients were enrolled including 114 with gastritis, 36 with PUD, 50 with PUB and 2 with gastric cancer. Prevalence of CYP2C19 genotype was 82/202 (40.6%) in RM, 99/202 (49%) in IM and 21/202 (10.4%) in PM. Overall H. pylori infection was 138/202 patients (68.3%). H. pylori infection was demonstrated in 72% in RM genotype, 69.7% in IM genotype and 47.6% in PM genotype. Both gastric cancer patients had the IM genotype. In PUB patients, the prevalence of genotype RM (56%) was highest followed by IM (32%) and PM (12%). Furthermore, the prevalence of genotype RM in PUB was significantly greater than gastritis patients (56% vs 36%: p=0.016; OR=2.3, 95%CI=1.1-4.7). Conclusions: CYP2C19 genotype IM was the most common genotype whereas genotype RM was the most common in PUB patients. All gastric cancer patients had genotype IM. The CYP2C19 genotype RM might be play role in development of PUB and PUB. Further study in different population is necessary to verify clinical usefulness of CYP2C19 genotyping in development of these upper GI diseases.

Keywords: CYP2C19 genotype - gastritis - peptic ulcer disease - gastric cancer

Asian Pac J Cancer Prev, 15 (24), 10957-10960

Introduction

CYP2C19 genotype has been found to be an important factor in peptic ulcer healing rate, H. pylori eradication, influencing the therapeutic efficacy of proton pump inhibitors (PPI) and development of gastric cancer (Sugimoto et al., 2009; Furuta et al., 2010; Prasertpetmanee et al., 2013). CYP2C19 genotypes can be determined by polymerase chain reaction (PCR) and restriction fragment length polymorphism analysis (RFLP) analysis and divided into 3 groups in relation to PPI metabolism; rapid metabolizer (RM), intermediate metabolizer (IM) or poor metabolizer (PM). Genotype RM experience rapid clearance of PPI and generally has less effective than PM (Yamada et al., 2001; Furuta et al., 2001; Sugimoto et al., 2009; Furuta et al., 2010; McNicholl et al., 2012). Among phenotypic RM and IM, higher dose of PPI or frequent dosing of PPI may require increasing efficacy of gastroprotective effect. Prior study suggested that it was difficult to assess the effect of CYP2C19 genotype on PPI because of small number studies, inadequate CYP2C19 genotype testing and lack of control (Estany-Gestal et al, 2011). The aim of this study was to investigate status and clinical correlation of CYP2C19 genotype in patients with gastritis, peptic ulcer disease (PUD), peptic ulcer bleeding (PUB) and gastric cancer in Thailand.

Materials and Methods

Patients who underwent gastroscopic examination at Thammasat University Hospital, Thailand for dyspeptic symptoms and UGIB were recruited between May 2012 and November 2014. Entry criteria included age over 18 years and those who had never received H. pylori eradication and had indication for upper GI endoscopy. We excluded those who were receiving anticoagulants, who had previously undergone gastric surgery and those patients with significant systemic diseases eg. congestive
heart failure, chronic renal failure and AIDS, patients who abused drug or alcohol, and women who were breast feeding or had child-bearing potential without using effective contraception. Informed consent was obtained before participating in this study. The diagnosis of gastritis, PUD, PUB and gastric cancer was made according to symptomatic assessment and endoscopic findings. Dyspeptic patients with normal endoscopy and those with gastric inflammation without erosions or ulcer were considered of having gastritis. Five milliliters of blood was collected for *H. pylori* serological diagnosis (Anti-*H. pylori* ELISA IgG) and for CYP2C19 study. CYP2C19 genotype was determined by PCR and RFLP analysis and divided into 3 groups: rapid metabolizer (RM), intermediate metabolizer (IM) or poor metabolizer (PM) and was performed as described previously (Furuta et al., 2001). After upper GI endoscopy, gastritis, PUD and gastric cancer patients continued to receive standard treatment from Thammasat university hospital and all PUB patients were received intravenous pantoprazole and the dose and duration of intravenous pantoprazole was introduced to these patients as suggested from the International consensus recommendations on the management of patients with nonvariceal upper gastrointestinal bleeding (Barkun et al., 2010).

**Statistical analysis**

The demographic information and frequencies of adverse effects were compared using chi-squared and Fisher’s exact test. The P-values <0.05 were considered to be statistically significant. For each variable, the odds ratio (OR) and 95% confidence interval (CI) were calculated. The data analysis was performed using SPSS version 19 (SPSS Inc., Chicago, IL, USA). The study was conducted according to the good clinical practice guideline, as well as the Declaration of Helsinki and was approved by our local ethics committee. All subjects signed informed consent to participate in this study.

**Results**

Total of 202 patients were enrolled (109 males and 93 females, mean age of 53 years). There were 114 patients with gastritis, 36 patients with PUD, 50 patients with PUB and 2 patients with gastric cancer.

**The CYP2C19 genotype**

CYP2C19 genotype was performed in all patients. Overall, the prevalence of genotype IM (49%) was the most common genotype followed by PM (40.6%) and RM genotype (10.4%). All gastric cancer patients had genotype IM. In PUB patients, the prevalence of RM genotype (56%) was highest followed by IM (32%) and PM genotype (12%). Furthermore, the prevalence of *H. pylori* infection was 138/202 patients.

<table>
<thead>
<tr>
<th>CYP2C19 Genotype</th>
<th>Gastritis (N=114)</th>
<th>PUB (N=50)</th>
<th>PUD (N=36)</th>
<th>Gastric cancer (N=2)</th>
<th>Total (N=202)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rapid metabolizer (RM)</td>
<td>41 (36%)</td>
<td>28 (56%)*</td>
<td>13 (36.1%)</td>
<td>0</td>
<td>82(40.6%)</td>
</tr>
<tr>
<td>Intermediate metabolizer (IM)</td>
<td>60 (52.6%)</td>
<td>16 (32%)</td>
<td>21 (58.3%)</td>
<td>2 (100%)</td>
<td>99(49%)</td>
</tr>
<tr>
<td>Poor metabolizer (PM)</td>
<td>13 (11.4%)</td>
<td>6 (12%)</td>
<td>2 (5.6%)</td>
<td>0</td>
<td>21(10.4%)</td>
</tr>
</tbody>
</table>

*p=0.016 OR=2.3 (95%CI=1.1-4.7) when compare with gastritis; ** p=0.068 OR=2.3 (95%CI=0.9-6) when compare with PUB

**Discussion**

CYP2C19 genotype has recently been found to have important roles on peptic ulcer healing and variable effectiveness of PPI (Yamada et al., 2001; Furuta et al., 2001). CYP2C19 genotype distributions vary among different ethnic groups. In Asian countries such as China, Japan and Vietnam, the prevalence of genotype IM was predominated (Yamada et al., 2001) whereas genotype RM was mainly found in European population (Xie et al., 2001). In our study, genotype IM was the most common genotype in Thailand same as other Asian countries. Because CYP2C19 is the main metabolizing enzyme of PPI, the variation of CYP2C19 genotype might be one of the explanations why PPI had different response between Asian and Western population.
The regimens for *H. pylori* eradication consisted of PPI and few antimalarial agents eg. amoxicillin, clarithromycin, metronidazole and fluoroquinolone. These antimalarial agents have uniquely pharmacologic actions. PPIs are metabolized by cytochrome P450 2C19 (CYP2C19), which is polymorphic. CYP2C19 genotypic differences might be influence the *H. pylori* eradication by variable response of PPI (pharmacokinetics and pharmacodynamics effect) in the regimens (Furuta et al., 2010). Eradication rates from PPI-based therapies might be interfering by different CYP2C19 genotype (Furuta et al., 2010). However, a recent study from Thailand demonstrated 7-day standard triple therapy plus bismuth and probiotic provided an excellent cure rate of *H. pylori* (100%) regardless of CYP2C19 genotype (Chanagune et al., 2014).

CYP2C19 genotype is also considered to be one of the factors that determine cancer susceptibility by different ability of carcinogen detoxification. CYP2C19*2 and CYP2C19*17 genotypes were associated with better survival in breast cancer patients using tamoxifen (Bai et al., 2014) and CYP2C19 polymorphisms might be significantly affect the treatment in lung cancer (Chen et al., 2014). It suggests that CYP2C19 might be participating and initiating procarcinogen and finally increase risk of these cancers. Gastric cancer is one of the important cancer in Thailand and Laos with grave prognosis (Varanarth et al., 2014; Vilaichone et al., 2014). Recent studies indicated that PM genotype of CYP2C19 had higher risk of gastric cancer (Shi et al., 2004; Sugimoto et al., 2005) and requires close monitoring future possible gastric cancer development (Sugimoto et al., 2005). Genotype PM had also appeared to be high incidence in esophagus cancer and lung cancer due to higher carcinogen level and potent cell toxicity from lower ability in carcinogenic detoxification. In our study, we demonstrated that genotype IM was found in all gastric cancer patients but only in a small number of patients. A larger group for CYP2C19 genotype in patients with gastric cancer is required for future analysis of the relevance to cancer development.

Our findings also indicated that CYP2C19 genotype RM might play an important role in the pathogenesis of PUB. Recent study from United Kingdom demonstrated that CYP2C19*17 was associated with PUB irrespective of etiology (Musumba et al., 2013). The effect of CYP2C enzymes on the metabolism of arachidonic acid, which is well known to be involved in the pathogenesis of PUB, might be the responsible mechanism (Musumba et al., 2009). Genotype RM was known to increase metabolic and clearance rate of PPI gastroprotection and finally enhancing risk of PUB. Genotype RM could also reduced mucosal gastroprotection from NSAIDS and *H. pylori* infection and might lead to develop PUB and PUB (Estany-Gestal et al., 2011; Musumba et al., 2013). Therefore, the direct effect of CYP2C19 genotype could contribute to the development of optimal therapy and pathophysiology in upper GI diseases patients. However, further large study in a different population is required to validate the usefulness of CYP2C19 genotyping in clinical practice.

**Acknowledgements**

This study was partially supported by Research Fund at Faculty of Medicine, Thammasat University Hospital, Thailand, Gastroenterology Association of Thailand (GAT) and the National Research University Project of Thailand Office of Higher Education Commission.

**References**


