A Matched Case-Control Study of a Novel Acid-Pump Antagonist and Proton-Pump Inhibitor for the Treatment of Iatrogenic Ulcers Caused by Endoscopic Submucosal Dissection

Yong Gil Kim, Byung-Ik Jang, and Tae Nyeun Kim
Department of Internal Medicine, Yeungnam University College of Medicine, Daegu, Korea

Background/Aims: Revaprazan, a novel acid-pump antagonist, and proton-pump inhibitors (PPIs) have pH-independent effects on ulcer healing. The addition of a PPI promotes the cell restitution rate as well as vessel regeneration and maturation for ulcer repair. Revaprazan is known to protect the mucosa by increasing the prostaglandin concentration.

Methods: We reviewed the medical records of patients who underwent endoscopic submucosal dissection (ESD) for gastric neoplasia at Yeungnam University Hospital between January 2008 and May 2009. We conducted a matched case-control study to compare the healing rates effected by revaprazan and rabeprazole.

Results: Each group consisted of 30 patients. The baseline characteristics did not differ significantly between the two groups. Stage S1 disease was observed in 97% and 100% of patients after 8 weeks of treatment in the revaprazan and rabeprazole groups, respectively. In the revaprazan group, only one patient had stage H2 disease: a 54-year-old man with a 5.5-cm lesion after ESD of the ulcer, type IIa early gastric cancer, and adenocarcinoma. No serious adverse effects occurred during the treatment period in either group.

Conclusions: The safety and efficacy profiles of revaprazan and rabeprazole are similar for the treatment of ESD-induced ulcers. (Gut Liver 2010; 4:25-30)

Key Words: Revaprazan; Rabeprazole; Proton pump inhibitors; Acid pump antagonists; Endoscopic submucosal dissection

INTRODUCTION

Proton pump inhibitors (PPIs), which are specific for H⁺/K⁺-ATPase, impede the function of the proton pump responsible for the terminal step in gastric acid secretion; they are currently the most widely used drugs for acid-related diseases including peptic ulcer disease, gastroesophageal reflux disease and Zollinger-Ellison syndrome. Reversible acid pump antagonists (APAs), another class of proton pump inhibitors, act by K⁺-competitive and reversible binding to the gastric proton pump.

APAs and PPIs have pH-independent effects on ulcer healing. The addition of PPIs to treatment promotes the cell restitution rate as well as vessel regeneration and maturation during wound healing. Rabeprazole demonstrates the significant protection of the gastric-mucosa of PPIs by increasing the prostaglandin E₂ production and reducing the leukotriene B₄ production. Revaprazan, a new drug, protects the mucosa by inhibiting secretion of pepsinogen and increasing the prostaglandin concentration. In previous study that compared revaprazan with a PPI for the treatment of gastric ulcers, the ulcer healing rates did not differ.

After the introduction of endoscopic submucosal dissection (ESD) in the late 1990s, ESD has become widely used as an alternative to surgical resection in patients with early-stage gastric cancer or for adenomas. However, EST causes artificial ulcers. It is unknown whether the healing of ESD induced ulcers differ from peptic ulcers with regard to healing rate and the factors that affect
healing. Furthermore, the efficacy of APA for ESD-induced ulcers has not been previously studied. Therefore, the aim of this study was to compare the efficacy of revaprazan with rabeprazole for treatment of ESD-induced ulcers.

MATERIALS AND METHODS

1. Endoscopic submucosal dissection

All ESD procedures were performed by a single endoscopist. ESD was performed by the endoscopic submucosal dissection method using a flex knife for en bloc resection (Fig. 1). If bleeding or visible vessels were found during the ESD, thermal coagulation or endoscopic hemoclipping was performed as required (Fig. 1).

2. Study design

We reviewed the medical records of patients that underwent ESD for gastric neoplasm at Yeungnam University hospital from January 2008 to May 2009. Demographic and clinical characteristics, including age and gender, were investigated. Location, size, and histopathological type of lesions were also investigated. Location was divided by antrum, proximal body, mid-body, distal body, angle and pre-pylorus. Histopathological types included adenomas (with low grade dysplasia, moderate dysplasia and severe dysplasia) and adenocarcinomas (well differentiated, moderately and poorly differentiated). Patients were evaluated for Helicobacter pylori infection by rapid urease test (CLO test) and endoscopic biopsy specimens by Giemsa staining.

Loading dose of pantoprazole 80 mg was given intravenously 2 hours before the ESD, and 8 mg/hr of intravenous pantoprazole was given continuously for the first 48 hours as maintenance. Starting three days after the ESD, oral acid suppressing agents were administered for 8 weeks. In the revaprazan group, revaprazan 200 mg was administered orally for 8 weeks, and in the rabeprazole group, rabeprazole 20 mg was given orally for the same period.

Bleeding encountered during the ESD was defined as immediate, and bleeding after the ESD as delayed. After the ESD, the patients were observed for their vital signs and complete blood counts. Delayed bleeding was suspected in patients with one or more of the following bleeding signs: fresh hematemesis, hematochezia, instability of vital signs (tachycardia or hypotension), or a reduction of hemoglobin by more than 2 g/dL after the

Fig. 1. After initial incision of the mucosa with the needle knife to allow insertion of tip of insulation-tipped diathermic (IT) knife, complete incision is made by IT knife circumferentially. (D) En bloc dissection of the submucosal tissue was performed. (E) EGC was resected completely. (F) Hemostasis was achieved with hemoclips and cauterezation.
ESD. When delayed bleeding was suspected, immediate endoscopy was performed.

Follow-up endoscopy was performed the next day, 56 days and 5-months after the ESD. The ulcer size was measured the day after the ESD. Ulcer stages were assessed using a six-stage system as proposed by Sakita and Fukutomi (Table 1) at 56 days after the ESD.

### Table 1. Gastric Ulcer Stages Using a Six-Stage System

<table>
<thead>
<tr>
<th>Stage</th>
<th>Finding</th>
</tr>
</thead>
<tbody>
<tr>
<td>A1 (active stage 1)</td>
<td>Ulcer that contains mucus coating, with marginal elevation because of edema</td>
</tr>
<tr>
<td>A2 (active stage 2)</td>
<td>Mucus-coated ulcer with discrete margin and less edema than active stage 1</td>
</tr>
<tr>
<td>H1 (healing stage 1)</td>
<td>Unhealed ulcer covered by regenerating epithelium &lt;50%, with or without converging folds</td>
</tr>
<tr>
<td>H2 (healing stage 2)</td>
<td>Ulcer with a mucosal break but almost covered with regenerating epithelium</td>
</tr>
<tr>
<td>S1 (scar stage 1)</td>
<td>Red scar with rough epithelization without mucosal break</td>
</tr>
<tr>
<td>S2 (scar stage 2)</td>
<td>White scar with complete re-epithelization</td>
</tr>
</tbody>
</table>

English version of ulcer stages as classified by Sakita and Fukutomi.

3. Patients

This study included 193 patients that underwent ESD for a gastric neoplasm between January 2008 to May 2009. Among them, 54 patients had received revaprazan therapy and 139 patients were treated with rabeprazole. The indications for ESD were as follows: gastric adenoma and early gastric adenocarcinoma (well or moderately differentiated; no ulceration; and no lymph node involvement or metastasis by CT). Patients were excluded if they had a previous history of upper gastrointestinal surgery or vagotomy; known hypersensitivity to revaprazan or rabeprazole; current use of aspirin, non-steroidal anti-inflammatory drugs, corticosteroids, PPIs, prostaglandins, or mucoprotecting agents, or had surgery for vertical margin-positive adenocarcinoma in the resection specimen. The patients that had delayed bleeding or gastric perforation the day after the ESD and required rapid treatment with IV PPIs (duration of taking acid suppressing agents influences gastric ulcer healing) were also excluded.

4. Case control study

All matching were performed with the Statistical Analysis Systems software package (Release 9.1.3; SAS Institute, Cary, NC, USA). The matching variables were age, gender, histology and initial ulcer size. The patients were matched on gender and histology by exact match, and matched on age intervals (plus or minus 5 years). The matched patients were divided into three categories based on initial ulcer size according to the medical literature: ≤30 mm, between 31 mm and 40 mm, and >40 mm. Then, thirty patients were randomly selected for each group using SAS program.

5. Statistical analysis

Continuous data are summarized as the mean (95% confidence interval [CI]). The Student t-test was used to compare the mean values of continuous variables. The Pearson’s chi-square test with Yates’ correction for continuity and the Fisher’s exact test were used as appropriate for comparison of the categorical variables. The analysis was performed with the statistical software package (SPSS 17.0 version for Windows; SPSS, Chicago, IL, USA). A p value less than 0.05 was accepted as statistically significant.

RESULTS

Data regarding the clinical and endoscopic features of the patients are outlined in Table 2. There were no significant differences between the two groups with respect to age, gender, ulcer size, location of ulcer, tissue size, histopathology (included histopathology of subgroup) and positive H. pylori.

Ninety seven percent and 100% of the patients had S1 stage disease after eight weeks of treatment in the revaprazan and rabeprazole group, respectively. In the revaprazan group, only one patient had H2 stage disease (Fig. 2); a 54-year-old man with a 5.5 cm post ESD ulcer on the lesser curvature of the proximal body the day after the ESD, type IIa EGCA and well differentiated adenocarcinoma. The endoscopic findings revealed a 0.5 cm ulcer almost completely covered with regenerating epithelium, eight weeks after the ESD and a red scar with rough epithelialization without mucosal breaks at five months after the ESD (Fig. 2).

During follow-up, no significant side effects were associated with the medications in either treatment group. There were no cases of delayed gastric perforation or bleeding after discharge.
Table 2. Baseline Characteristics of Patients

<table>
<thead>
<tr>
<th></th>
<th>Revaprazan (n=30)</th>
<th>Rabeprazole (n=30)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, yr</td>
<td>61.0±9.5</td>
<td>63.1±10.1</td>
<td>0.414</td>
</tr>
<tr>
<td>Gender, M/F</td>
<td>21/9</td>
<td>21/9</td>
<td>1.0</td>
</tr>
<tr>
<td>Histology (Subgroup)</td>
<td></td>
<td></td>
<td>1.0 (0.494)</td>
</tr>
<tr>
<td>Adenoma</td>
<td>21</td>
<td>21</td>
<td></td>
</tr>
<tr>
<td>Adenocarcinoma</td>
<td>9</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Location</td>
<td></td>
<td></td>
<td>0.487</td>
</tr>
<tr>
<td>Antrum</td>
<td>16</td>
<td>19</td>
<td></td>
</tr>
<tr>
<td>Proximal body</td>
<td>0</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Mid-body</td>
<td>1</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Distal body</td>
<td>6</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>Angle</td>
<td>3</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Prepylorus</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>H. pylori (+)</td>
<td>4</td>
<td>7</td>
<td>0.333</td>
</tr>
<tr>
<td>Ulcer size</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean, mm</td>
<td>24.7±13.4</td>
<td>28.5±10.5</td>
<td>0.223</td>
</tr>
<tr>
<td>Median, mm</td>
<td>22 (10-81)</td>
<td>26.5 (11-49)</td>
<td></td>
</tr>
<tr>
<td>Tissue size</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean, mm</td>
<td>31.9±13.6</td>
<td>27.7±9.4</td>
<td>0.372</td>
</tr>
<tr>
<td>Median, mm</td>
<td>28 (12-75)</td>
<td>28 (12-63)</td>
<td></td>
</tr>
</tbody>
</table>

Continuous data are expressed as mean±standard deviation and median (minimum-maximum).

DISCUSSION

The use of endoscopic mucosal resection (EMR) procedures for early gastric cancer has been increasing. This is a preferred procedure because the patient quality of life after an EMR is superior to that of patients after a surgical gastrectomy. After the introduction of the endoscopic submucosal dissection (ESD), in the late 1990s, EMR has become more technically feasible for all tumors, regardless of size. Currently, ESD is most widely used as an alternative to surgical resection in patients with early-stage gastric cancer or adenomas.6

ESD induces the development of ulcers. It is unclear whether these ESD induced ulcers differ from peptic ulcers with regard to the healing rate or the factors that influence healing. EMR-induced ulcers have been reported to heal faster and to recur less often than non-iatrogenic gastric ulcers.10 Increased healing has been associated with preserved contractility of the proper muscle layer, which is usually damaged in patients with peptic ulcers,11 and the increased blood supply at the margin of EMR-in-
duced ulcers resulting from increased levels of nitric oxide, epidermal growth factor and endogenous prostaglandins. The healing process of ESD-induced ulcers may be similar to EMR-induced ulcers. A recent study comparing the healing rate of EMR-induced ulcers with regard to the duration of treatment with omeprazole suggested that one week of treatment may be sufficient. In addition, an observational study showed that gastric ulcers caused by ESD procedures healed within eight weeks regardless of the size and location. However, Oh et al. reported that the degree of healing of ESD-induced ulcers was dependent on the initial ulcer size, indicating that the duration of treatment with PPI should take into consideration the initial size of the ulcer.

Gastric H⁺/K⁺-ATPase is a proton pump located at the apical membrane of the parietal cells; it transports H⁺ into the canaliculus of parietal cells in exchange for K⁺. Substituted benzimidazoles, such as omeprazole, lansoprazole and rabeprazole, inactivate the H⁺/K⁺-ATPase by covalent binding to the sulfohydryl group of H⁺/K⁺-ATPase, resulting in the long lasting inhibition of gastric acid secretion. PPIs also have pH-independent effects on ulcer healing. PPIs attenuate leukocyte-endothelial cell adhesion by inhibiting the expression of cellular adhesion molecules. PPIs have been reported to directly inhibit inflammation by suppressing neutrophil activity as well as by relaxing vascular smooth muscle. Thus, PPIs have multiple functions during gastric mucosal repair. In recent studies, PPIs were shown to promote the healing of injured gastric mucosal cells following injury by enhancing cell proliferation and migration; rabeprazole increased the expression of CXCR4 mRNA. Thus, it promoted vessel regeneration and maturation, facilitating ulcer healing.

Revaprazan, 5,6-dimethyl-2-(4-fluorophenylamino)-4-(1-methyl-1,2,3,4-tetrahydroisoquinoline-2-yl) pyrimidine hydrochloride, is a novel APA that is currently approved by the Korean Food and Drug Administration (KFDA) as a new drug for the treatment of gastric diseases including peptic ulcer disease. This APA has a reversible mode of action against H⁺/K⁺-ATPase, which is distinct from PPIs that perform irreversible inhibition. The drug causes inhibition of gastric acid secretion rapidly and effectively without sustained hypergastrinemia. This inhibitor is competitive with the luminal K⁺ ions, and has a highly selective affinity for H⁺/K⁺-ATPase, and low affinity for Na⁺/K⁺-ATPase. Interestingly, Revaprazan, like pantoprazole, plays a regulatory role in MAPK ERK1/2 signalling. MAPK is an executive critical enzyme that is active during inflammatory response signaling as well as cell proliferation. In addition, revaprazan has been shown to provide cyto-protection of the gastric epithelium. In the present study, the ulcer healing effect of revaprazan for ESD-induced ulcers was similar to that of rabeprazole. PPIs are generally considered to be extremely safe with little or no risk of serious adverse events. However, concerns continue to be expressed about existing and potential adverse effects of these drugs, especially with long-term use. In animal studies, in the K⁺ channel mutated mouse model, in beta subunit gastric H⁺/K⁺-ATPase deficient mice, and in the alpha subunit of the gastric H⁺/K⁺-ATPase deficient mice achlorhydria and vacuolization of the gastric parietal cells were described. Since the beta subunit of the gastric H⁺/K⁺-ATPase and gastrin double deficient mice did not exhibit such hyperplasia, it is speculated that such vacuolization might be attributed to secondary hypergastrinemia. Driman et al. reported enlarged parietal cells with luminal bulges of cytoplasm, expanded tubulovesicles, and poorly developed secretory canaliculi after omeprazole treatment for longer than 12 months in a human study. Bajaj et al. reported that PPI use during just more than 1 week was associated with a higher risk of SBP. These results suggest that PPI usage leads to hypergastrinemia, and hypergastrinemia leads to impaired intestinal permeability in humans. However, APA had a weaker effect on the serum gastrin concentration and antral gastrin content than PPIs. Therefore, APA would be beneficial for clinical use. This is the first study to demonstrate the efficacy of APA for the treatment of ESD-induced ulcers. In conclusion, Revaprazan showed a similar safety and efficacy profile to rabeprazole for the treatment of ESD-induced ulcers. Further prospective studies are needed to explore the pathophysiology of this novel acid pump antagonist and proton pump inhibitors for the treatment of ESD-induced ulcers.

REFERENCES

5. Yeo M, Kwak MS, Kim DK, et al. The novel acid pump an-