A Comparative Study on the Efficacy of Covered Metal Stent and Plastic Stent in Unresectable Malignant Biliary Obstruction

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Background/Aims: The placement of self expandable metal stent (SEMS) is one of the palliative therapeutic options for patients with unresectable malignant biliary obstruction. The aim of this study was to compare the effectiveness of a covered SEMS versus the conventional plastic stent.

Methods: We retrospectively evaluated 44 patients with unresectable malignant biliary obstruction who were treated with a covered SEMS (21 patients) or a plastic stent (10 Fr, 23 patients). We analyzed the technical success rate, functional success rate, early complications, late complications, stent patency and survival rate.

Results: There was one case in the covered SEMS group that had failed technically, but was corrected successfully using lasso. Functional success rates were 90.5% in the covered SEMS group and 91.3% in the plastic stent group. There was no difference in early complications between the two groups. Median patency of the stent was significantly prolonged in patients who had a covered SEMS (233.6 days) compared with those who had a plastic stent (94.6 days) (p=0.006). During the follow-up period, stent occlusion occurred in 11 patients of the covered SEMS group. Mean survival showed no significant difference between the two groups (covered SEMS group, 236.9 days; plastic stent group, 222.3 days; p=0.182).

Conclusions: The patency of the covered SEMS was longer than that of the plastic stent and the lasso of the covered SEMS was available for repositioning of the stent.

Key Words: Malignant biliary obstruction; Self-expandable metal stent; Plastic stent

INTRODUCTION

Biliary stent insertion is performed in patients with unresectable malignant biliary obstruction as a palliative therapy to prevent early death due to deterioration of liver function or suppurative cholangitis.¹³ A plastic stent insertion had been the standard of the endoscopic biliary drainage, but this method was limited by the fact that it needs to be replaced every 3 to 4 months due to the stent occlusion or migration.⁴ Self expandable metal stent (SEMS) with longer patency than the conventional plastic stent was developed to address this limitation and has been in use since 1989.⁵,⁶ An uncovered metal stent is advantageous in its low frequency of migration after the mesh was embedded with the tumor and the availability of using it in the intrahepatic bile duct; it is, however, associated with 20% to 40% incidence of early occlusion mainly due to the tumor ingrowth,⁷⁸ and is not available for replacement or removal once it was placed inside the bile duct.⁹,¹⁰,¹¹

Covered SEMS was developed to address these limitations and to prolong the patency, but it also has limitations, such as higher frequency of migration compared with the uncovered metal stent¹² and the difficulty of stent repositioning.
once inserted inside the bile duct. We performed this multi-center study to evaluate the clinical efficacy of a lasso-attached covered SEMS insertion to prevent tumor ingrowth in patients with malignant biliary obstruction and to compare the efficacy with the conventional 10 Fr plastic stent.

MATERIALS AND METHODS

Patients
This study was performed in patients who were diagnosed of having an extrahepatic biliary stricture due to a malignant tumor and received the endoscopic retrograde biliary drainage (ERBD) at 7 university hospitals in South Korea from January 2005 to July 2006. These patients were unresectable or refused a surgery. A retrospective analysis was performed in 21 patients who received the covered SEMS insertion and 23 patients who received the 10 Fr plastic stent insertion among those available for long-term follow-up.

Malignant biliary stricture was diagnosed based on ultrasoundography, abdominal computed tomography (CT), magnetic resonance cholangiopancreatography, endoscopic retrograde cholangiopancreatography (ERCP), and histological findings. Patients with hilar cholangiocarcinoma, history of biliary stent insertion, concomitant duodenal obstruction, or past history of gastrointestinal operation were excluded from the study.

Methods
A 10-mm covered SEMS (Hanaro stent; M.I. Tech, Seoul, Korea), with lasso for location adjustment and silicone membrane for the prevention of tumor ingrowth, was used for the metal stent group (Fig. 1). A 10 Fr polyethylene plastic stent (Wilson-Cook Medical, Winston-Salem, NC, USA) was used for the plastic stent group.

Conventional side viewing duodenoscopes (JF 240V, TJF 240; Olympus Medical System Co., Tokyo, Japan) were used for ERCP in every patient. Endoscopic sphincterotomy (EST) was performed using a pull-type papillotome, and along the papillotome, a guide wire was inserted in the bile duct through the stricture site. The location and length of the stricture was assessed with the fluoroscopy, and then the stent was inserted along the guidewire. The Hanaro stent (M.I. Tech) was inserted across the stricture site into the extrahepatic bile duct, along the guidewire, until the distal end was exposed about 1 cm from the major papilla to the duodenum. For the plastic stent, a guidewire was first inserted into the bile duct across the stricture site for the following guiding catheter insertion into the bile duct, and then the plastic stent was pushed across the stricture site using the pusher catheter.

Technical success was defined when the stent was placed across the stricture and allowed the passing of the contrast media or the bile; while functional success was defined when the level of bilirubin decreased by 30% or more compared with the baseline level within a week or to a normal level within a month. Early or late complication was defined as any stent-related complication occurring within 30 days of or 30 days after the insertion, and the procedure-related death was defined as deaths due to complications associated with the EST or stent insertion. Functional failure of the stent was defined as signs of acute cholangitis, such as abdominal pain, jaundice or fever; increase of bilirubin level; or radiologic (such as abdominal CT or US) evidence of stent occlusion.

Stent patency was defined as the period between the date of stent insertion and the date of functional failure or death by disease progression. We also determined the cause and period of stent functional failure as well as the cause and time to death.

Statistical analysis
Chi-square test was performed for comparison of categorical variables, and Student’s t-test for continuous variables. Stent patency and patient survival were calculated according to the Kaplan-Meier analysis, and comparison of stent patency and survival between the two groups were analyzed by the log-rank test. A p-value <0.05 was considered significant. Statistical analysis was performed using SPSS version 12.0 (SPSS Inc., Chicago, IL, USA).

RESULTS
Among a total of 44 patients, 21 (10 males) were assigned to the Hanaro covered SEMS group and 23 (10 males) to the plastic stent group, and their mean age were 73.1 ± 7.4 years (range, 60 to 88 years) and 65.7 ± 13.2 years (range, 51 to 85 years), respectively. Biliary obstruction was caused by chol-
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angiocarcinoma in 12 patients, pancreatic cancer in 6 patients, gallbladder cancer in 2 patients and ampullary cancer in 1 patient of the covered SEMS group; and by cholangiocarcinoma in 9 patients, pancreatic cancer in 7 patients, gallbladder cancer in 4 patients and ampullary cancer in 3 patients of the plastic stent group, without a significant difference between the two groups (Table 1). The stricture site in the common bile duct was proximal in 2 patients, mid in 10 patients and distal in 9 patients of the covered SEMS group; proximal in 5 patients, mid in 4 patients and distal in 14 patients of the plastic stent group.

Technical success rates were 95.2% (20/21) in the covered SEMS group and 100% (23/23) in the plastic stent group, without any significant difference between the two groups (p=0.290), except 1 case of technical failure due to the stent misplacement in the covered SEMS group, which was corrected using forceps to catch the lasso and achieved the functional success. Functional success rates were 90.5% (19/21) in the covered SEMS group and 91.3% (21/23) in the plastic stent group.

The mean follow-up periods were 329.52 ±246.53 days (range, 4 to 908 days) in the covered SEMS group and 336.48 ±271.52 days (range, 88 to 1,189 days) in the plastic stent group. Early complications occurred in 3 patients from each group. In the covered SEMS group, mild pancreatitis occurred in 1 patient, cholangitis in 1 patient and cholecystitis in 1, and all of them were recovered with conservative treatment. In the plastic stent group, mild pancreatitis occurred in 1 patient and the patient was recovered with conservative treatment, but 1 patient of cholangitis required covered SEMS insertion and another patient of the cholangitis was treated with percutaneous transhepatic biliary drainage (PTBD) (Table 2).

The mean patency duration was significantly longer in the covered SEMS group (233.6 days; range, 21 to 660 days) compared with the plastic stent group (94.6 days; range, 4 to 549 days) (p=0.006) (Table 2, Fig. 2).

The causes of stent occlusion in the covered SEMS group were bile sludge in 4 patients, tumor ingrowth in 4 patients, tumor overgrowth in 1 patient, stent migration in 1 patient and unknown origin in 1 patient. Among them, 9 patients were treated with plastic stent insertion inside the SEMS (n=4), metal stent insertion (n=4) and PTBD (n=1). In the plastic stent group, the causes of stent occlusion were bile sludge in 6 patients, tumor overgrowth in 2 patients, late stent migration in 2 patients and unknown origin in 1 patient, which were treated with metal stent insertion after re-

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Table 1. Baseline Characteristics of the Patients with Unresectable Malignant Biliary Obstruction

<table>
<thead>
<tr>
<th></th>
<th>Hanaro stent (n=21)</th>
<th>Plastic stent (n=23)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr, range)</td>
<td>73.1±7.4 (60-88)</td>
<td>65.7±13.2 (51-85)</td>
</tr>
<tr>
<td>Male/Female</td>
<td>10/11</td>
<td>10/13</td>
</tr>
<tr>
<td>Diagnosis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bile duct cancer</td>
<td>12</td>
<td>9</td>
</tr>
<tr>
<td>Pancreatic cancer</td>
<td>6</td>
<td>7</td>
</tr>
<tr>
<td>Ampullary cancer</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>Gallbladder cancer</td>
<td>2</td>
<td>4</td>
</tr>
</tbody>
</table>

Values are presented as mean±SD.

Table 2. Outcome for Patients with Stent Placement

<table>
<thead>
<tr>
<th></th>
<th>Hanaro stent</th>
<th>Plastic stent</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Technical success rate (%)</td>
<td>20/21 (95.2)</td>
<td>23/23 (100.0)</td>
<td>0.290</td>
</tr>
<tr>
<td>Functional success rate (%)</td>
<td>19/21 (90.5)</td>
<td>21/23 (91.3)</td>
<td>0.924</td>
</tr>
<tr>
<td>Early complication</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pancreatitis</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Cholangitis</td>
<td>1</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Cholecystitis</td>
<td>1</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Mean stent patency (range), day</td>
<td>233.6 (21-660)</td>
<td>94.6 (4-549)</td>
<td>0.006</td>
</tr>
<tr>
<td>Cause of obstruction</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bile clogging</td>
<td>4</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>Tumor ingrowth</td>
<td>4</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Tumor overgrowth</td>
<td>1</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Stent migration</td>
<td>1</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>Unknown</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Mean survival (range), day</td>
<td>236.9 (4-673)</td>
<td>222.3 (12-647)</td>
<td>0.182</td>
</tr>
</tbody>
</table>

One case that had failed technically, was repositioned successfully using lasso.

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Stent has significantly longer patency than plastic stent \((p=0.006)\). Moving the plastic stent \((n=6)\), re-insertion of the plastic stent \((n=3)\) and PTBD \((n=2)\) (Table 2).

Mean survival was 236.9 days (range, 4 to 673 days) in the Hanaro stent group and 222.3 days (range, 12 to 647 days) in the plastic stent group. There was no significant difference in the mean survival between the two groups \((p=0.182)\) (Fig. 3).

**DISCUSSION**

ERBD is effective in the prevention and treatment of cholangitis and enables physiologic drainage of the bile, making it a suitable palliative therapy for patients with unresectable malignant biliary obstruction.\(^2,3\) The indications of ERBD include malignant biliary obstruction due to pancreatic cancer or cholangiocarcinoma, suppurrative cholangitis, common bile duct stone impaction, bile duct stricture after surgery, and obstructive jaundice concomitant with chronic pancreatitis. In malignant biliary obstruction, the placement of biliary plastic stent rapidly improves the jaundice and allows semi-permanent biliary drainage in unresectable cases. However, the plastic stent is not permanent and requires replacement or additional stent insertion in several months when the occlusion occurs. Efforts have been made to reduce the stent occlusion, such as by decreasing bacterial adhesion inside the stent or by changing the material or structure-size and shape, that is of the stent,\(^4\) without an apparent success yet.

Plastic stent is not expensive and replaceable, but requires frequent replacement due to the stent migration or frequent occlusion leading to the recurrence of obstructive jaundice or cholangitis.\(^5,10\) Metal stent was developed in 1989 to overcome these limitations and has been in use since then.\(^5,6\) SEMS is associated with less occlusion and cholangitis and shorter duration of hospital stay compared with the plastic stent.\(^7,8\) The plastic stent induces occlusion mostly by the formation of biliary sludge. The metal stent, however causes occlusion mostly by the tumor ingrowth, where the tumor grows through the gaps of the stent mesh; by the tumor overgrowth, where the tumor grows to the upper and lower space of the stent; and by epithelial hyperplasia inside the stent.\(^16,17\) The rate of metal stent occlusion due to the tumor ingrowth or overgrowth after some time, has been reported variously from 7% to 42%.\(^5,10,18-20\)

According to a prospective trial by Davids et al.,\(^7\) the early efficacy of drainage by the plastic stent and the metal stent are not significant different at 95% and 96%, respectively. This study also revealed no difference of the early efficacy by the materials of the stents, with the polyethylene plastic stent and the Hanaro covered SEMS respectively showing 91.3% (21/23) and 90.5% (19/21) of functional success rates. Given the fact that a stent of 7 Fr or more is physiologically sufficient to induce biliary drainage,\(^21\) this lack of difference in the early efficacy of drainage between the plastic stent group and the Hanaro covered SEMS group in our study may come from using 10 Fr plastic stents. The time to biliary stent occlusion of a 10 Fr plastic stent has been reported average 134 to 164 days.\(^22,23\) Davids et al.\(^7\) reported stent patency of 273 days for metal stent (Wallstent; Schneider, Minneapolis, MN, USA) and 126 days for polyethylene plastic stent in distal malignant biliary obstruction. Our findings also showed 233.6 days and 94.6 days of mean stent patency for the Hanaro covered SEMS group and the plastic group, respectively.

The advantage of the SEMS is that the stent diameter can be increased remarkably to allow early sufficient drainage, the duration of patency is longer than the plastic stent, and the frequency of migration is low.\(^7\) But several disadvantages of the SEMS have been also pointed out, including the difficulty of insertion technique compared with that of the plastic stent,\(^24\) the possibility of injuries such as bleeding or perforation in the duodenum and the papilla,\(^25\) high cost, inability to remove once placed, and the possibility of stent occlusion due to the tumor ingrowth or overgrowth.\(^26\) Despite the development of covered SEMS, using a thin layer of polyurethane, silicon, polytetrafluoroethylene or Gore-Tex to cover
SEMS (30 Fr), to prevent functional failure due to the tumor ingrowth or overgrowth, tumor ingrowth is still reported to occur with the membrane covered metal stent. 27-30 Suh et al. 31 reported, in a study of 59 patients who were diagnosed with extrahepatic biliary stricture due to the malignant tumor, that the tumor ingrowth was the cause of the stent functional failure in 6.6% (2/29) of the covered SEMS group compared to 24% (7/30) of the uncovered SEMS group, suggesting that the covered SEMS may prevent the tumor ingrowth efficiently. The functional failure of the Hanaro covered SEMS in our study was due to stent occlusion in 9 patients and of migration in 1 patient (late stent migration); among these 9 patients of occlusion, 4 patients (44.4%) were associated with the tumor ingrowth, 1 patient with the tumor overgrowth, and the remaining 4 patients with the biliary sludge, reflecting that the covering membrane of the SEMS cannot prevent the tumor ingrowth completely. Tumor ingrowth in the covered SEMS may occur when the membrane was damaged during the stent insertion, by the bile or biliary stone, or by bacterial infection, but little data are available regarding the exact cause of tumor ingrowth.

Early complications occurring during the stent insertion or within 2 weeks after the insertion include cholangitis, cholecystitis, perforation, bleeding, and acute pancreatitis, at 6.4% to 26.3% of incidence, among which cholangitis occurring most frequently. 22 Davids et al. 27 reported that 11% of the plastic stent group and 12% of the metal stent group were associated with the early (within a week after the stent insertion) complications, which is similar to our results of 12.5% (3/23) for the plastic stent group and 14.3% (3/21) for the Hanaro covered SEMS group. Iyayama et al. 32 reported that the membrane attached to a stent often blocks the orifice of the cystic duct or the pancreatic duct causing cholecystitis or pancreatitis. In this study, there were 3 cases of early complications in the both groups, respectively. In the Hanaro covered SEMS group, there were pancreatitis in 1 patient, cholangitis in 1 patient and cholecystitis in 1 patient, all of them were recovered with conservative treatment. In the plastic stent group, pancreatitis occurred in 1 patient and cholangitis in 2 patients; the former was recovered with conservative treatment, while each of the patients with cholangitis required SEMS insertion and PTBD, respectively.

There is still a controversy in what type of stent should be used for ERBD in terms of cost-effectiveness. Prat et al. 33 reported 3.2 months of survival for ≥30 mm of tumor diameter and 6.6 months of survival for ≤30 mm of tumor diameter in 101 patients with unresectable malignant biliary obstruction. Therefore, they insisted that the metal stent may be more effective than the plastic stent when the tumor diameter is 30 mm or less in unresectable malignant biliary obstruction. 33 Schmassmann et al. 37 also reported that the metal stent was more effective than the plastic stent in patients with more than 6 to 9 months of life expectancy.

Cahen et al. 34 emphasized that modification of the SEMS design was necessary for convenient removal and repositioning of the stent. The lasso attached on the covered SEMS in our study allowed functional success in 1 case of technical failure during the insertion by enabling the stent repositioning at ease.

In conclusion, the placement of the covered SEMS with longer stent patency seemed to be more effective than that of the plastic stent in patients with unresectable malignant biliary obstruction who their life expectancy are estimated to be longer than 6 to 9 months based on their general condition, the presence of comorbidity and biliary stricture diameter. The lasso of the covered SEMS seemed to be helpful in repositioning of the stent to achieve functional success, although this issue needs to be further studied in prospective studies with more cases.

**Conflicts of Interest**

The authors have no financial conflicts of interest.

**REFERENCES**

179:603-605.