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Clinical Outcomes of Percutaneous Endoscopic Gastrostomy in the **Surgical Intensive Care Unit**

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Background/Aims: Percutaneous endoscopic gastrostomy (PEG) is usually performed on patients with chronic underlying diseases in the general ward (GW). This study evaluated the clinical outcomes of PEG performed on patients in the surgical intensive care unit (SICU) compared with those of PEG performed in the GW.

Methods: The medical records of 27 patients in the SICU and 263 in the GW, who underwent PEG between January 2013 and July 2017, were retrospectively reviewed.

Results: The median age of the 27 SICU patients was 66 years, and their median body mass index was 21.1 kg/m2. In the SICU group, the median baseline Sequential Organ Failure Assessment (SOFA) score was 4, and the median Acute Physiology and Chronic Health Evaluation II (APACHE II) score was 16. The median interval between surgery and PEG in SICU patients was 30 days, with a PEG failure rate of 3.7%. Acute complications in SICU patients included bleeding (7.4%) and ileus (11.1%), while chronic complications included aspiration pneumonia (7.4%) and tube obstruction (3.7%). The rates of acute and chronic complications did not differ significantly between the SICU and GW groups. The 30-day mortality rate was 14.8% in SICU patients and 5.3% in GW patients (p=0.073).

Conclusions: PEG is a safe and feasible method of enteral feeding for critically ill patients who require ICU care after surgery. Clin Endosc 2020;53:705-712

Key Words: Endoscopy; Gastrostomy; Intensive care unit

INTRODUCTION

Percutaneous endoscopic gastrostomy (PEG) is usually performed on patients with chronic underlying disease in the general ward (GW). PEG has been reported to be indicated for several chronic neurological diseases such as cerebrovascular disease, Parkinson's disease, dementia, motor neuron disease,

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and specific malignancies. PEG is the most common method of supplying enteral nutrition to patients expected to require supplementary enteral feeding for longer than 2-3 weeks.² Although it is safe and minimally invasive, few reports have assessed the clinical outcomes of PEG administered to critically ill patients or compared these outcomes in patients who had undergone previous abdominal surgery and those with an intact abdomen.3-8 In addition, no study to date has compared the safety and feasibility of PEG in patients in the surgical intensive care unit (SICU) and GW. Accordingly, this study assessed the clinical outcomes of PEG in patients in the SICU who had undergone surgical intervention or were scheduled for surgery, and compared these outcomes with those of patients who underwent PEG in the GW.



MATERIALS AND METHODS

Patients

Of the 324 patients who underwent PEG at the Asan Medical Center (Seoul, Korea) between January 2013 and July 2017, 27 underwent PEG in the SICU and 263 underwent the procedure in the GW. The remaining 34 patients, consisting of 32 who underwent PEG in the medical ICU and two who underwent PEG in the emergency room, were excluded from this study (Fig. 1). Both patients in the emergency room underwent PEG due to cancer progression and were not indicated for surgery. The medical records of all included patients were retrospectively reviewed, and clinical outcomes were compared between SICU and GW patients. Demographic and clinical characteristics were evaluated, including underlying diseases, body mass index (BMI), indications for PEG, complications after PEG, comorbidities, and baseline laboratory values. In addition, baseline Sequential Organ Failure Assessment (SOFA) scores and Acute Physiology and Chronic Health Evaluation (APACHE) II scores for SICU patients were evaluated. Weight and serum albumin concentrations were compared before and after PEG. If post-PEG weight and albumin level were assessed multiple times between 2 weeks and 1 month after PEG, the most recent measurements were used. PEG was performed on patients who had normal gastrointestinal function and were expected to require enteral feeding for longer than 2–3 weeks. The Institutional Review Board (IRB) of Asan Medical Center (Seoul, Korea) approved the study (IRB number, 2018-0928). The requirement for patient informed consent was waived due to the retrospective design of the study.

Percutaneous endoscopic gastrostomy methods

Of 290 PEG procedures, 289 were performed using the introducer technique and one was performed using the pull-type technique. The PEG procedure was performed by experi-

enced gastroenterologists. The introducer (Russell) technique has been the main approach since November 2011, while pull-type PEG is usually performed for the purpose of drainage. The introducer technique was performed using a transilluminated endoscope, and the gastropexy device and trocar were used to penetrate the abdominal wall. A 15-Fr Cliny PEG tube (Create Medic Co., Ltd., Yokohama, Japan) was subsequently inserted directly through the abdominal wall and fixed by filling the balloon of the tube with 5 cc of saline (Fig. 2).

Of 290 patients, 279 (96.2%) were administered prophylactic antibiotic 30 min before PEG, with most receiving 2 g of cefazolin. If a patient was already receiving an antibiotic, an altered regimen was not required unless the current regimen was inappropriate for gram-positive cocci. For sedation, intravenous midazolam (0.05 mg/kg) and pethidine (25 or 50 mg) were administered before the procedure.

Definitions

Acute complications were defined as complications that developed within the first 7 days of PEG insertion, and chronic complications were those that occurred after 7 days. Acute complications included bleeding, ileus, pneumoperitoneum, aspiration pneumonia, wound infection, and Mallory–Weiss tear. Chronic complications included tube obstruction, tube leakage, spontaneous tube removal, wound infection, recurrent aspiration pneumonia, and buried bumper syndrome.

Bleeding was defined as a hemorrhagic event requiring endoscopic intervention or embolization for hemostasis and reduction of >2 g/dL of hemoglobin concentration. Aspiration pneumonia was defined as newly developed symptoms, such as cough, purulent sputum, or fever after PEG, in addition to changes in chest X-ray. Diagnosis of ileus was confirmed using X-ray or computed tomography imaging indicative of ileus in addition to decreased bowel sounds with or without symptoms.

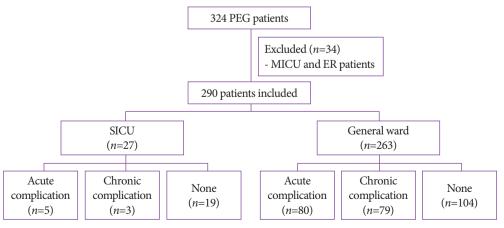
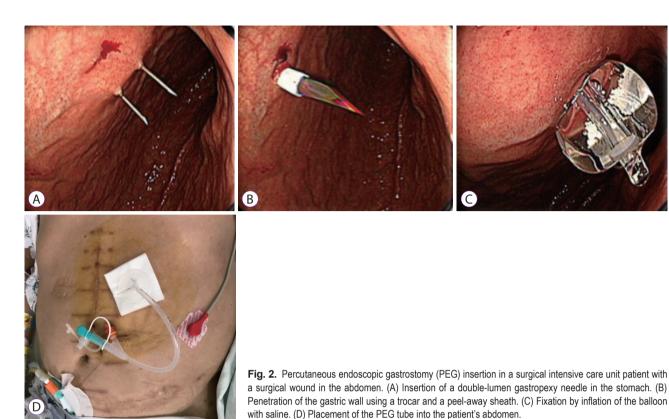


Fig. 1. Study flowchart. ER, emergency room; MICU, medical intensive care unit; PEG, percutaneous endoscopic gastrostomy; SICU, surgical intensive care unit.



Post-percutaneous endoscopic gastrostomy management

Patients were kept nil per os on the day of the PEG procedure with natural tube drainage. The next day, the PEG tube of patients with a soft abdomen and those with no abdominal discomfort was flushed with 100 mL of clear water. After the water flush, patients without abdominal pain, vomiting, or tube leakage started tube feeding with 100–200 mL of fluid.

Patients in the GW were started on tube feeding with 100 cc of formulated nutrition, with the volume incrementally increased by 50–100 cc for every meal, as tolerated until the target volume was reached. Patients in the SICU were started on continuous feeding with 20 cc/hr, with the amount increased by 10–20 cc/hr every 12–24 hr until the target volume was achieved.

Statistical analysis

Descriptive data are expressed as mean \pm standard deviation for normally distributed continuous variables or as median and interquartile range (IQR). Continuous variables were compared between groups using the t test or Mann–Whitey U test, and categorical variables were compared using the chi-square or Fisher's exact test, as applicable. Multivariate logistic regression models were constructed to evaluate risk factors for complications and 30-day mortality. SPSS version 21.0 (IBM

Co., Armonk, NY, USA) was used for all statistical analyses; differences with a p-value < 0.05 were considered to be statistically significant.

RESULTS

Baseline characteristics

The baseline demographic and clinical characteristics of the patients in the GW and SICU groups, including age, sex, BMI, indications for PEG insertion, comorbidities, and baseline laboratory values, are presented in Table 1. The median age of the 27 SICU patients was 66 years (IQR, 58-77 years), and their median BMI was $21.1 \, \text{kg/m}^2$ (IQR, $17.8-24.0 \, \text{kg/m}^2$).

The median baseline SOFA score of the SICU patients was 4 (IQR, 3–9), and their median APACHE II score was 16 (IQR, 12–22) (Table 2). Indications for surgery in the SICU group included nine (33.3%) patients with traumatic fracture with brain hemorrhage and four (14.8%) with malignancy including mouth floor cancer, distal common bile duct cancer, hepatocellular carcinoma, and sigmoid colon cancer. Four (14.8%) patients had panperitonitis. The mean interval from surgery for panperitonitis to the PEG procedure was 36.2 days. Patients exhibited improving clinical course of panperitonitis but presented other postoperative complications, such as ven-



Table 1. Baseline Characteristics of Patients Who Underwent Percutaneous Endoscopic Gastrostomy Insertion

	General ward (n=263)	SICU (<i>n</i> =27)	p-value
Clinical characteristics			
Age, median (IQR)	70 (61–79)	66 (58–77)	0.268
Sex, male	174 (66.2%)	20 (74.1%)	0.405
BMI, median (IQR)	18.9 (16.8–21.1)	21.1 (17.8–24.0)	0.003
Indications for PEG, n (%)			0.020
Neurologic disease	154 (58.6%)	14 (51.9%)	
Malignancy	73 (27.8%)	4 (14.8%)	
Other	36 (13.7%)	9 (33.3%)	
Comorbidities, n (%)			
Hypertension	109 (41.4%)	13 (48.1%)	0.502
Diabetes mellitus	70 (26.6%)	6 (22.2%)	0.621
Chronic kidney disease	13 (4.9%)	3 (11.1%)	0.177
Antibiotic prophylaxis, n (%)	254 (96.6%)	25 (92.6%)	0.273
Concurrent medication, n (%)			
Aspirin	38 (14.4%)	5 (18.5%)	0.571
Clopidogrel	25 (9.5%)	5 (18.5%)	0.176
Warfarin	10 (3.8%)	1 (3.7%)	1.000
Laboratory data			
WBC, μL	7,200 (5,800–10,000)	7,600 (5,400–9,900)	0.825
Hemoglobin, g/dL	11.0 (9.6–12.4)	8.4 (7.6–9.1)	< 0.001
Platelet, 10 ³ /μL	251 (191–319)	261 (143–338)	0.888
Albumin, g/dL	2.8 (2.4–3.3)	2.3 (1.9–2.6)	< 0.001
C-reactive protein, mg/dL	1.4 (0.4–4.2)	1.8 (1.2–6.0)	0.015
Procedure time, min	18.6±7.2	17.8±7.3	0.332

BMI, body mass index; IQR, interquartile range; PEG, percutaneous endoscopic gastrostomy; SICU, surgical intensive care unit; WBC, white blood cell.

tilator-associated pneumonia and cerebral infarction, which lengthened hospital stay. Seven (25.9%) patients underwent surgery for other conditions including abdominal aorta aneurysm, total hip replacement arthroplasty, infective endocarditis, kidney transplantation rejection, simultaneous pancreas kidney transplantation, and ulcerative colitis. Three patients in the SICU did not undergo any surgical intervention. One patient was admitted to the SICU due to a traffic accident and was diagnosed with brain death. A second patient was admitted to the SICU after falling and sustaining an intracerebral hemorrhage; after undergoing a tracheostomy and being treated for pneumonia, he was transferred to a local hospital. The third patient had undergone a kidney transplant and was admitted to the SICU for suspicion of acute rejection. She was treated medically for the rejection without surgical intervention.

Clinical outcomes

The median interval between surgery and PEG in the SICU group was 30 days (IQR, 19–51 days), and the median follow-up period was 73 days (IQR, 21–189 days). PEG insertion was successful in 26 of the 27 patients in the SICU. PEG insertion was unsuccessful in one (3.7%) patient due to the inability of the needle to puncture the gastric wall. Repeated surgical procedures in this patient may have led to the development of postoperative anatomic distortion. This patient had undergone a subtotal colectomy for an ischemic colitis induced perforation with panperitonitis and had underlying end-stage renal failure. After the colectomy, she underwent additional anastomosis revision and ileostomy due to anastomosis site leakage. Three weeks after surgery, respiratory arrest occurred due to aspiration. PEG in this patient failed.

Post-PEG complications are summarized in Table 3. Acute complications in SICU patients included bleeding (7.4%) and

Table 2. Clinical Characteristics of Surgical Intensive Care Unit Patients

APACHE II score (median)	16 (IQR, 12-22)
SOFA score (median)	4 (IQR, 3–8)
Vasopressor	3 (11.1%)
Indication for surgery	
Traumatic fracture with brain hemorrhage	9 (33.3%)
Traumatic fracture without brain hemorrhage	1 (3.7%)
Malignancy	4 (14.8%)
Panperitonitis	4 (14.8%)
Others (UC, THRA, AAA, IE, KT rejection, SPKT)	6 (22.2%)
No surgery	3 (11.1%)
Abdominal surgery	10 (37.0%)
Bowel resection (small bowel resection and colectomy)	4
Transplantation (KT and SPKT)	2
Others (lobectomy, primary repair of cystic duct stump, PPPD, AAA surgery)	4
Interval between surgery and PEG, days (median)	30 (IQR, 19-51)

AAA, abdominal aorta aneurysm; APACHE, Acute Physiology and Chronic Health Evaluation; IE, infective endocarditis; IQR, interquartile range; KT, kidney transplantation; PEG, percutaneous endoscopic gastrostomy; PPPD, pyloric preserving pancreaticoduodenectomy; SOFA, Sequential Organ Failure Assessment; SPKT, simultaneous pancreas and kidney transplantation; THRA, total hip replacement arthroplasty; UC, ulcerative colitis.

Table 3. Complications in General Ward and Surgical Intensive Care Unit Patients

	General ward (n=263)	SICU (n=27)	<i>p</i> -value
Acute complications			
Bleeding	23 (8.7%)	2 (7.4%)	1.000
Pneumoperitoneum	30 (11.4%)	0 (0.0%)	0.090
Ileus	27 (10.3%)	3 (11.1%)	0.749
Chronic complications			
Wound infection	13 (4.9%)	0 (0.0%)	0.618
Leakage	16 (6.1%)	0 (0.0%)	0.378
Tube obstruction	34 (12.9%)	1 (3.7%)	0.222
Spontaneous removal	12 (4.6%)	0 (0.0%)	0.611
Buried bumper syndrome	1 (0.4%)	0 (0.0%)	1.000
Aspiration pneumonia	3 (1.1%)	2 (7.4%)	0.070

SICU, surgical intensive care unit.

ileus (11.1%), and chronic complications included aspiration pneumonia (7.4%) and tube obstruction (3.7%). The incidence rates of acute and chronic complications did not differ significantly between the GW and SICU groups.

Of the four SICU patients who developed bleeding, none required intervention to stop the bleeding. By comparison, five patients in the GW group underwent endoscopic intervention, such as clipping, coagrasper, or embolization, to stop the bleeding. Ileus was detected in three patients in the SICU group, and all patients exhibited improvement following conservative management.

Of the 27 patients in the SICU group, 12 were discharged to the GW, seven were transferred to local hospitals, and eight died during SICU admission. All 12 patients discharged to the GW were fed through PEG tubes; their median length of stay in the SICU was 53 days (IQR, 25–75 days). The median length of stay in the SICU for all 27 patients was 57 days (IQR, 28–83 days). The median interval from surgery to PEG insertion in patients with intra-abdominal surgery was 46 days (IQR, 20–71 days).

Nine patients (33.3%) were able to have their PEG tube removed and resume oral feeding. The median interval of PEG use in these nine patients was 37 days (IQR, 25–75 days).

Prognosis

Clinical prognosis after PEG insertion, including early mortality rate and nutritional status, is summarized in Table 4. The 30-day mortality rate in the SICU group was 14.8%, which was considerably higher than the 5.3% in the GW group. However, none of the patients in the SICU group died of procedure-related causes. Four patients in the SICU group died within 30 days of the PEG procedure. One of these patients was admitted

Table 4. Prognosis of General Ward and Surgical Intensive Care Unit Patients after Percutaneous Endoscopic Gastrostomy Insertion

	General ward (n=263)	SICU (n=27)	<i>p</i> -value
Follow-up period, days	252±16	159±44	0.078
Nutrition status			
Mean weight change after PEG, kg	0.14±3.46	-0.59±3.66	0.472
Mean BMI change after PEG	0.04±1.35	-0.25±1.32	0.411
Mean albumin level change after PEG, g/dL	0.17±0.59	0.33±0.54	0.419
30 day mortality	14 (5.3%)	4 (14.8%)	0.073

BMI, body mass index; PEG, percutaneous endoscopic gastrostomy; SICU, surgical intensive care unit.



to the SICU due to a traffic accident and was diagnosed with brain death. The second patient underwent intra-abdominal surgery after being transferred to the SICU for panperitonitis and died 15 days after PEG from aggravated ischemic colitis. A third patient with rectal cancer and multiple lung metastases was admitted to the SICU for drug-induced lung injury; this patient died of aggravated pneumonia 3 days after PEG. The fourth patient, who had ischemic colitis and underwent a right hemicolectomy and cholecystectomy, developed pneumonia and ventricular tachycardia and died of cardiac arrest 8 days after PEG insertion.

Fourteen patients in the GW group died within 30 days after PEG, including one from PEG procedure-related causes. This patient died 12 days after PEG insertion of newly developed aspiration pneumonia. Four patients died of aspiration pneumonia not related to the PEG. Other causes of death included renal failure, acute myocardial infarction, traumatic subdural hemorrhage, and sepsis due to urinary tract infection. Five patients died of unknown causes, because they were lost to follow-up.

Mean changes in weight, BMI, and serum albumin concentration from before to after PEG insertion did not differ significantly in the GW and SICU groups.

DISCUSSION

PEG has been commonly administered as a method of enteral feeding for patients with chronic medical conditions. Nutritional support is important for both critically ill patients and postoperative patients, and recent systematic reviews of randomized trials found that 'nil by mouth' following gastrointestinal surgery yielded no obvious advantage, and recommended early enteral feeding. Appropriate nutritional support can benefit critically ill patients by improving wound healing and reducing complication rates and length of stay. 12

Currently, there are no standard guidelines or indications for PEG in patients in the GW and in critically ill patients. Although PEG is a safe and minimally invasive procedure, no risk factors have been consistently found to determine candidates for PEG procedures. Despite its introduction in the 1980s, PEG is administered to relatively few patients in the ICU, and clinical trials of PEG have included only small numbers of patients.

While few studies have assessed outcomes of PEG insertion for patients in the ICU, PEG has been reported to be safe and effective for critically ill patients in an ICU, with low rates of procedure-related morbidity and mortality.³⁻⁵ However, to our knowledge, the present study was the first to compare clinical

outcomes of PEG insertion in SICU and GW patients.

The SICU patients had a median baseline SOFA score of 4 (IQR, 3–9) and a median APACHE II score of 16 (IQR, 12–22). A significant cutoff point for SOFA score has not yet been established, although SOFA score cutoffs in predicting mortality are reported to range from 2 to 7. ¹³⁻¹⁵ APACHE II score cutoffs of 10–24 are reported to have high sensitivity and specificity in predicting clinical outcomes. ¹⁶⁻¹⁸ Considering the roles of SOFA and APACHE II scores in determining disease severity, the SICU patients in the present study were regarded as having moderately severe disease.

The present study found that PEG was a successful nutritional method in patients who underwent abdominal surgery, including small bowel resection, colectomy, pyloric-preserving pancreaticoduodenectomy, liver lobectomy, and kidney transplantation, and that these patients maintained PEG without major complications. Moreover, the rates of acute and chronic complications related to PEG did not differ significantly between SICU and GW patients. A previous study comparing PEG in ICU patients with no previous abdominal surgery and those with recent or previous laparotomy found no significant differences in complication rates of bleeding, PEG dislodgment, and wound complication. Although critically ill patients in the ICU and SICU are vulnerable to infection, 19,20 the present study found that the rates of wound complications did not differ significantly between the SICU and GW groups. Antibiotic prophylaxis has been reported to be successful in preventing peristomal wound infection.^{21,22}

In addition, postoperative ileus is a frequent complication following gastrointestinal and other types of surgery and is associated with patients' morbidity and clinical outcomes.²³ In the present study, the rates of ileus after PEG were comparable between the GW (10.3%) and SICU (11.1%) groups. In a previous study, early enteral nutrition after gastrointestinal surgery was associated with significantly less ileus.²⁴ Although ileus is one of the potential complications of PEG, early enteral nutrition may provide some benefit, but should be selected after considering potential clinical benefits and harms.

Similar to a study demonstrating that no patients in the neurosurgical ICU experienced procedure-related complications, the rate of complications, including bleeding and pneumoperitoneum, was lower in SICU patients than in GW patients, although the difference was not statistically significant. Comparisons of the complication and mortality rates in the GW and SICU groups revealed that PEG was safe and effective in providing nutritional support.

The reported success rate of PEG in patients who previously underwent surgery range from 88% to 94.7%. ^{25,26} Anatomic deformity after surgery and intra-abdominal adhesions may

affect the success rate of PEG insertion. Eleftheriadis et al. emphasized the importance of transillumination and finger indentation observed during endoscopy for successful PEG.⁶ In addition, previous research reported complications of gastrocolic and colocutaneous fistulas.^{27,28} However, in the present study, no patient experienced fistulas penetrating other organs or the skin.

Limitations of the present investigation are mainly associated with its retrospective design. First, this study was conducted at a tertiary referral hospital, where disease severity is higher than at general hospitals. Second, the number of SICU patients was much smaller than that of the GW patients. However, this study is the first to report clinical outcomes of PEG in SICU patients and may inform further studies involving postoperative and critically ill patients. Large randomized controlled trials are required to confirm the efficacy of PEG for patients who undergo surgical interventions.

In conclusion, PEG was a safe and feasible method of enteral feeding for critically ill patients in the SICU. Before PEG insertion, anatomic deformities caused by surgical interventions should be considered. Transillumination and finger indentation during PEG insertion are important to its success.

Conflicts of Interest _

The authors have no financial conflicts of interest.

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