An overview of percutaneous endoscopic gastrostomy tube placement in the intensive care unit

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Abstract: Critically ill patients are at increased risk for malnutrition as they often have underlying acute and chronic illness, stress related catabolism, decreased appetite, trauma and ongoing inflammation. Malnutrition is recognized as a leading cause of adverse outcomes, higher mortality, and increased hospital costs. Percutaneous endoscopic gastrostomy (PEG) tubes provide a safe and effective route to provide supplemental enteral nutrition to these patients. PEG placement has essentially replaced surgical gastrostomy as the modality of choice for longer term feeding in patients. This is a highly prevalent procedure with 160,000 to 200,000 PEG procedures performed each year in the United States. The purpose of this review is to provide an overview of current knowledge and practice standards with regards to placement of PEG tube in the Intensive Care Unit (ICU). When a patient is considered for a PEG tube, it is important to evaluate the treatment alternatives and identify the best option for each patient. In this review, we provide the advantages and disadvantages of various feeding modalities and devices. We review the indications and contraindications for PEG tube placement as well as the risks of this procedure. We then describe in detail the per-oral pull, per-oral push, and direct percutaneous techniques for PEG tube placement. Additionally, we review the feasibility of having interventional pulmonologists place PEG tubes in the ICU.

Keywords: Percutaneous endoscopic gastrostomy (PEG) placement; interventional pulmonology; Intensive Care Unit (ICU)

Introduction

Critically ill patients are at increased risk for malnutrition, particularly given their underlying acute and chronic illness, stress related catabolism, decreased appetite, trauma and ongoing inflammation (1). Malnutrition is recognized as a leading cause of adverse outcomes, higher mortality, and increased hospital costs (2).

Percutaneous endoscopic gastrostomy (PEG) tubes provide a safe and effective method to provide supplemental enteral nutrition to these patients. PEG placement was a technique first described by Gauderer *et al.* in 1980 (3). PEG placement has essentially replaced surgical gastrostomy as the procedure of choice for a longer term feeding in patients. PEG tubes allow for enteral feeding when oral feeding is not possible or insufficient. This is a highly prevalent procedure with 160,000 to 200,000 PEG procedures occurring each year in the United States.

The purpose of this review is to provide an overview of current knowledge and practice standards with regards to placement of PEG tubes in the Intensive Care Unit (ICU). Additionally, we consider the feasibility of having

interventional pulmonologists place PEG tubes in the ICU.

Feeding modalities

When a patient is considered for a PEG tube, it is important to evaluate the treatment alternatives and identify the best option for each individual patient. Below we consider the advantages and disadvantages of various feeding modalities and devices.

Parenteral vs. enteral

Parenteral as compared to enteral nutrition has been associated with higher costs of care, increased risk of both bacterial and fungal infections, particularly central line associated infections, hyperglycemia, as well as longer ICU length of stays. However, there is no difference in the mortality (4-9).

A recent multicenter randomized control trial found no difference between patients receiving parenteral and enteral feeding with regards to 30-day mortality, but did find an increased risk of hypoglycemia and vomiting with enteral feeding (10).

Also, one recent review found no difference between parenteral and enteral feeding in mortality, ventilator free days, aspiration, or pneumonia. It did however find that enteral nutrition may be associated with lower risk of sepsis as compared to parenteral nutrition (RR 0.59, 95% CI, 0.37–0.95) (11).

PEG vs. nasogastric/nasoduodenal/nasojejunal feeding

Temporary enteral access can be achieved with nasogastric (NG), oral gastric (OG), nasojejunal (NJ), or oral jejunal (OJ) feeding tube. These options are recommended for short-term use, when feeding is required for a few days or up to one month.

Bedside enteric tube placements are commonly used in the hospital and long term care environments. NG, OG, NJ, OJ can be placed blindly with confirmation of appropriate positioning by auscultation and radiography (12).

A variety of novel systems for navigational guidance during nasogastric tube (NGT) placement are currently being studied. Rivera *et al.* evaluated the use an electromagnetic tube placement device whereby the tip of a nasogastric feeding tube emitted electromagnetic pulses, which were picked up by sensors on a receiver placed on the epigastric region. He demonstrated successful post-pyloric

placement of the NGT with this device and showed that localization with this device correlated with that of plain film radiography (13).

Li *et al.* studied the feasibility of deploying a flexible uteroscope inside a gastric tube to provide visual guidance for placement of the gastric tube. With this method, there was an increase in rate of successful tube placement and reduced time for tube placement. This study also noted an overall operator preference for the visual system over the conventional method (14). Sun *et al.* has studied the use of electromagnetic tracking system to identify the location of the NGT during the insertion process. With this technique, a permanent magnet is embedded within the tip of the NGT and a wearable device with embedded sensors is placed at the neck, which senses the passive magnetic field of the NGT as it traverses the esophagus into the stomach (15).

These navigational systems have yet to be adopted into standard clinical practice. However, in the future we may see increased adoption of these systems given the risk of aspiration, pneumonia, pneumothorax, hemothorax, esophageal perforation, or in some cases even death with incorrect placement of NGTs (16,17).

Nasoenteric tube placement is a known risk factor for nosocomial sinusitis. These tubes can cause obstruction of the nasal ostia and provide a conduit for colonization of the nasopharynx by gastric microorganisms. One prospective epidemiologic study performed in ICU patients noted that feeding through a NGT was associated with increased risk of sinusitis (OR 14.1, P=0.015) (18). A subsequent epidemiologic study also found that while there was an overall low incidence (0.15%) of nosocomial sinusitis, the combination of nasoenteric and endotracheal intubation was associated with increased odds of sinusitis compared with endotracheal intubation alone (19).

Furthermore, prolonged use of NGTs can lead to nasal and esophageal ulcerations, aspiration pneumonia, and gastroesophageal reflux disease. These tubes can also cause trauma to the stomach, peptic esophagitis, stricture formation, and even tracheoesophageal fistula formation (20).

These complications are less of a concern with PEG tubes, which enter the stomach directly, bypassing the nasopharynx and esophagus. PEG tubes also have lower rates of intervention failure and offer patients an improved quality of life. As such they are preferred for longer term nutrition.

One systemic review evaluated rates of intervention failure (e.g., feeding interruption, blockage or leakage of the tube, treatment non-adherence) and concluded failure rates

in the PEG tube population were significantly lower than failure rates in the NGT population at 9.22% and 39.11%, respectively (RR 0.18, 95% CI, 0.05–0.59, P=0.005). However, the review failed to demonstrate differences in mortality, complications of the intervention, aspiration rates, and nutritional status (21).

Gastric vs. post-pyloric enteral feeding

While PEG is often the preferred method of providing long term enteral nutrition, the concern with gastric feeding is the risk of aspiration, particularly in critically ill patients who may have delayed gastric emptying and altered mentation. Post-pyloric feeding may be preferred in patients with gastroparesis, gastric outlet obstruction, or history of gastrectomy. There are different methods to achieve post-pyloric feeding. The percutaneous endoscopic gastrojejunostomy (PEG-J) is commonly used and is achieved by advancing a smaller tube through a PEG tube with placement of the distal end of the smaller tube in the jejunum. An alternative approach is achieved through direct percutaneous endoscopic jejunostomy (PEJ), whereby a large bore feeding tube is directly placed in the jejunum under endoscopic guidance (22). PEG-J and PEJ tubes are more technically challenging to place and require specialized equipment and training, but may offer an advantage over PEG by minimizing aspiration risk.

One meta-analysis found that PEJ was able to reduce gastric residual volumes and was more likely to meet the energy requirements of a patient, but did not provide significant benefit over PEG tubes (23). Another study in trauma patients also found that patients who utilized PEG-J for post-pyloric feeding and gastric decompression were able to reach their goal rates faster than patients with PEG. However, there was no difference in ventilator days, pneumonia, sepsis, or hospital length of stay (24).

Several recent meta-analyses found evidence demonstrating lower rates of pneumonia with post-pyloric feeding compared with gastric feeding. Nonetheless, there was no difference in ventilator days or mortality between the two groups in these studies (25-27).

Timing of feeds

Some studies have suggested that early enteral nutrition reduces the risk of mortality in critically ill patients as compared with delayed initiation of enteral nutrition (28). Early enteral nutrition for stroke patients with dysphagia

also showed improvement in mortality (29).

Current European Society for Clinical Nutrition and Metabolism (ESPEN) and American Society of Parenteral and Enteral Nutrition (ASPEN) guidelines recommend consideration of starting enteral nutrition after 24–48 hours in the ICU. Enteral nutrition is recommended over parenteral nutrition when clinically appropriate (30,31). However, the majority of mechanically ventilated patients who receive enteral nutrition do not receive sufficient caloric intake to meet their daily energy requirements (32).

While the use of enteral nutrition alone often does not meet the caloric needs, early initiation of parenteral nutrition to supplement enteral nutrition has not been proven to be beneficial. One randomized control trial demonstrated that delayed parenteral nutrition in these patients was associated with shorter ICU length of stay, shorter duration of mechanical ventilation, shorter hospital length of stay, and reduced hospital costs (33).

While early feeding is recommended in the critical care setting, there has been conflicting evidence regarding whether high calorie feeding or trophic feeding offers greater benefit. A recent study has illustrated the feasibility of early trophic feeding in patients with septic shock as a potential approach to improve enteric immune function during the hyper-acute phase of sepsis (34).

However, one randomized control trial evaluated patients with acute lung injury and found that initial trophic feeding for 6 days as compared with early full caloric feeding did not improve ventilator free days, 60-day mortality, or incidence of pneumonia, but did lower the risk of gastrointestinal complications (35). A subsequent study also demonstrated that permissive underfeeding was associated with no difference in feeding intolerance (e.g., vomiting, abdominal distention, or gastric residual over 200 cc), diarrhea, ICU-associated infections, or mortality when compared to full caloric feeding (36).

Patient selection

PEG is indicated for patients who require long-term nutritional support (>30 days) and have a functional gastrointestinal tract, but without sufficient oral intake (37).

Neurologic dysphagia is one of the most common reasons for PEG tube insertion. Neurologic indications include stroke, dementia, amyotrophic lateral sclerosis (ALS), Guillain-Barré Syndrome, Parkinson's disease, cerebral palsy, multiple sclerosis (MS), myasthenia gravis, Huntington's disease, encephalitis, meningitis, and

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polymyositis (38).

PEG tube insertion is also common in patients with physical obstruction of the upper gastrointestinal tract as seen with head and neck cancers, esophageal cancers, or trauma. PEG tube placement can also be indicated for decompression, such as in gastric outlet obstruction.

Cerebrovascular disease/stroke

Dysphagia can occur in up to 65% of patients with stroke. George *et al.* found that 8.5% of patients hospitalized for stroke received tracheostomy in addition to PEG tube placement. Receiving a tracheostomy during an admission was the strongest predictor for subsequently receiving a PEG on that same admission (adjusted odds ratio 0.27, 95% CI, 0.24–0.29). Early PEG tube placement (<7 days) during a hospitalization for stroke was associated with shorter length of stay, greater odds of discharge to home or acute rehabilitation, and lower odds of discharge to skilled nursing or long-term care (39).

Patients with higher stroke severity, worse dysphagia severity, diabetes, history of prior intracerebral hemorrhage, increased age, and health insurance coverage have a significantly increased likelihood of PEG placement (40).

Motor neuron disease/ALS

ALS is a rapidly progressive neurologic disease that leads to quadriplegia, dysarthria, and dysphagia. PEG placement in ALS has been shown to be safe even in patients with pulmonary dysfunction and may improve quality of life (41-43).

Dementia

In the advanced stages of dementia, patients often develop feeding issues that can arise from difficulty with physically feeding themselves, controlling a food bolus, and swallowing. They may have difficulty recognizing food and have malfunction with the limbic and hypothalamic systems that regulate hunger and satiety. PEG tube placement in patients with dementia is very common and the decision to place a PEG tube is often influenced by family member or caregiver concerns that the patient may starve. However, in patients with dementia, enteral feeding via PEG tube has not been shown to improve the risk of aspiration, pressure ulcers, pneumonia, or mortality as compared with oral feeding (44,45). As such, the current American Geriatric

Society (AGS) guideline recommend oral feeding over PEG tube placement in patients with dementia (46).

Nevertheless, about one third of nursing home patients with dementia have PEG tubes (47). Hospital characteristics associated with higher rates of feeding tube placement include larger sized hospitals, for-profit hospitals, and hospitals with more ICU days documented for chronically ill patients within the last 6 months of life.

Black and Hispanic patients with dementia also have higher rates of PEG tube placement (48). It is unknown at this time if this is a result of cultural differences, disparities in health care access, or discrepancies in physician communication. Physician misperceptions regarding the utility of PEG tubes are also often a contributing factor to the overuse of PEG tubes in patients with dementia despite evidence to the contrary (49,50).

Head and neck malignancy

Head and neck cancer is a common indication for PEG tube placement. Patients with head and neck cancers can experience difficulty swallowing for a variety of reasons including pain, obstruction by the tumor, or chemo-radiation side effects including mucositis. In fact, dysphagia occurs in up to 54% of patients with advanced head and neck cancer treated with chemo-radiation (51).

One review showed that of all head and neck cancer patients studied, 30.5% had a gastrostomy tube placed at some time. Of all patients who receive treatment, 35.1% had a gastrostomy tube placed. Of those who had tubes placed, 16.9% had their gastrostomy tubes placed before treatment and 83.1% were placed after treatment (52).

Given that many patients with head and neck cancer often have a long term need for enteral nutrition, some studies have evaluated the use of prophylactic PEG tube placement for patient with head and neck cancers and have found this practice to be safe and effective without increasing the risk of long term dysphagia (53).

Prophylactic PEG tube placement in patients with advanced head and neck cancer may also minimize weight loss during chemo-radiotherapy as compared with reactive PEG tube placement. Delayed PEG placement may be associated with increased risk of aspiration and stricture. However, timing of PEG tube placement does not improve tumor control or overall survival (54-56).

Alternatively, NGT remains a viable option to PEG tube placement in these patients. One recent study in patients with head and neck cancer found that while PEG provided
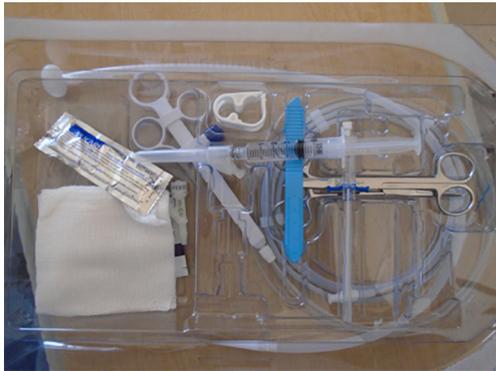
Figure 1 Boston Scientific EndoVive™ 20 French Safety PEG kit, Pull technique.

better weight management, patients with NGT placement had the device in place for a significantly shorter period of time. The PEG tube was often considered by patients to be more cosmetically appealing and more convenient. There were no quality of life differences between the two study groups. NGT was associated with increased risk of dislodgement, but even accounting for this difference, PEG tubes were costlier (57,58).

Gastric decompression

PEG tube placement for gastric decompression has also been described as a feasible palliative option for patients with malignant bowel obstruction, but is often associated with higher risk of complications and poor prognosis (38,59,60).

Miscellaneous

PEG tube placement has been described in patients with Crohn's disease, cystic fibrosis, severe trauma, and altered level of consciousness (61).

Procedure

There are three main techniques for PEG tube placement: the per-oral push technique, the per-oral pull technique, and the direct percutaneous technique.

Per-oral pull technique

The most commonly used technique is the pull method as originally described by Gauderer *et al.* in 1980 (3). At our institution, we use the Boston Scientific EndoVive™ 20 French Safety PEG kit (*Figure 1*) using the per-oral pull technique described below (*Table 1*).

The endoscope is passed into the stomach, which is then insufflated to achieve maximal apposition of the gastric and abdominal walls. A full esophagogastroduodenoscopy (EGD) is done to ensure that there is no obstruction or malignancy prior to PEG tube placement (*Figure 2*). This is critical to rule out any post-pyloric obstruction. Any abnormality should be documented. The gastrostomy site is located with a combination of trans-illumination and finger indentation of the abdominal wall (*Figure 3A,3B*). The ideal PEG tube location should be at the gastric antrum. Local anesthetic (lidocaine 1% with epinephrine) is infiltrated subcutaneously, and while maintaining a 90-degree angle with the abdominal wall, the local anesthetic is injected into the tract (*Figure 3C*).

A small skin incision is made to allow insertion of a needle-catheter combination also called trocar catheter (*Figure 3D-3F*). The trocar catheter is then inserted along the designated tract into the stomach. There is a risk of colonic interposition between the gastric and abdominal walls during this procedure. As such, in addition to trans-illumination and finger indentation, direct visualization with aspiration of the needle syringe provides additional reassurance that there is no colonic interposition. If the needle is not directly visualized entering the stomach wall, but reflux of air is noted in the syringe, this may indicate inadvertent puncture of the transverse colon.

Once within the stomach, the needle is removed and the small wire loop at the end of the guidewire is passed through the catheter (*Figure 3G*). During this phase of the procedure, it is critical not to damage the posterior wall of the stomach during needle insertion. A snare system is passed through the endoscope and secures the wire loop (*Figure 3H*). The wire loop is then pulled up through the esophagus and out of the mouth (*Figure 4A*).

In the Ponsky technique (per-oral pull), the wire exiting the mouth is attached to the feeding tube. The wire and the feeding tube are then pulled down through the esophagus, into the stomach, and out through the anterior abdominal wall. It is important to lubricate the tube for smooth placement (*Figure 4B*). A skin disc (external bolster) is then placed over the feeding tube (*Figure 5A*). While visualizing the internal bumper with the endoscope, the external bolster is adjusted until the internal bumper is flush against the anterior stomach wall.

The external bolster is kept between 2–5 cm as indicated

Table 1 The Per-oral pull technique for percutaneous endoscopic gastrostomy tube placement

Steps	Description	Reference Image
1	Gather all necessary equipment	*Figure 1*
2	Perform an esophagogastroduodenoscopy	*Figure 2*
3	Confirm gastrostomy location externally with trans-illumination and internally with finger indentation	*Figure 3A,3B*
4	Infiltrate the region with local anesthetic including the intended tract of the gastrostomy tube	*Figure 3C*
5	Make a small skin incision with a scalpel	*Figure 3D*
6	Insert the needle-catheter combination (trocar) through the incision and into the stromach	*Figure 3E*
7	Visualize passage of the needle-catheter combination through the anterior gastric wall with the endoscope	*Figure 3F*
8	Remove the needle, leaving the catheter in place Pass the wire loop through the catheter into the stomach	*Figure 3G*
9	Using the endoscope, snare the wire loop	*Figure 3H*
10	With the endoscope, pull the loop up through the esophagus and out of the mouth	*Figure 4A*
11	Apply lubricant to the gastrostomy tube	*Figure 4B*
12	Thread the wire loop through the tip of the gastrostomy tube	*Figure 4C*
13	Loop the head of the gastrostomy tube through the wire loop, securing the two pieces together with a knot	*Figure 4D*
14	Apply firm pressure while pulling the wire and gastrostomy tube through the anterior abdominal wall	*Figure 4E*
15	Continue pulling until the internal bumper sits flush against the anterior gastric wall	*Figure 4F*
16	Cut the gastrostomy tube and place the external bolster onto the feeding tube	*Figure 5A*
17	Push the external bolster so that it gently secures the tube in place, making note of the markings on the tube	*Figure 5B*
18	Apply antibiotic ointment between the external bolster and skin surface	*Figure 5C*
19	Thread the C-clamp onto the gastrostomy tube	*Figure 5D*
20	Insert gastrostomy tube adapter to the end of the tube	*Figure 5E*

by the markings on the tube. This may vary depending on the patient's body habitus. Care should be made to ensure that the bolster is neither too tight nor too loose to prevent buried bumper syndrome or leakage, respectively (*Figure 5B*). Antibiotic ointment is applied between the bumper and the skin (*Figure 5C*). A clamp and an external port are added to the PEG tube to complete the procedure (*Figure 5D-5F*). The external bolster should be evaluated the day after the procedure to ensure appropriate positioning. If the bolster is too tight, it may need adjustment or loosening.

Per-oral push technique

In the Sacks-Vine technique (per-oral push), the feeding tube is passed over the wire and pushed down through the stomach until the tapered end emerges through the skin. The guide wire is then withdrawn and the endoscope is reinserted to ensure proper positioning of the feeding tube.

A skin disc is secured over the feeding tube to keep it in place (61,62).

The per-oral pull and per-oral push techniques, however, may not be possible or may be contraindicated in a variety of clinical scenarios. For example, patients with head and neck or esophageal cancer may have high-grade stenosis preventing the passage of the endoscope or the internal PEG tube bumper. Additionally, in patients with head and neck cancer, the passage of the tube through a primary site of malignancy may result in the implantation of cancer cells to the gastrostomy tube site resulting in abdominal wall metastasis (63).

Introducer PEG technique

For these patients, the Introducer PEG technique is an alternative to overcome the limitations of the traditional per-oral pull or push techniques (63). In the introducer
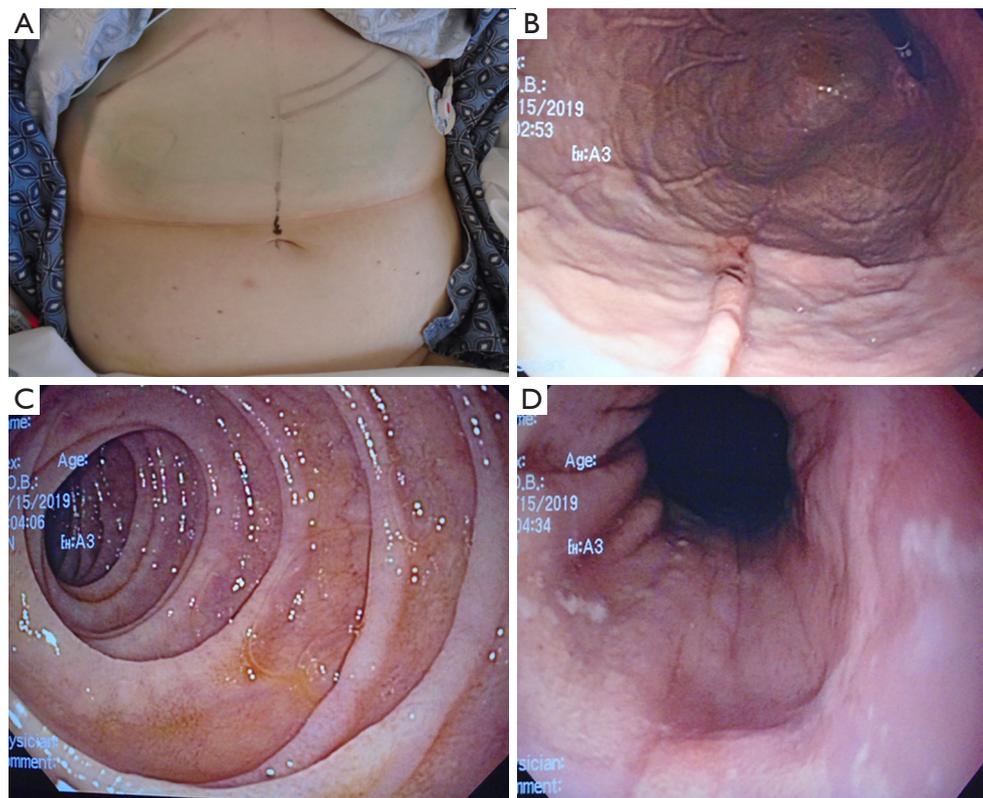

Figure 2 Pre-procedural evaluation for PEG tube placement. (A) Patient's abdomen marked with a surgical pen prior to the procedure; (B) endoscope retroflexed, evaluating the gastric cardia for lesions; (C) evaluating the duodenum for evidence of post-pyloric obstruction; (D) evaluating the esophagus for lesions.

PEG method, once the site of the gastrostomy is identified by trans-illumination and finger indentation, two to four sutures are placed under endoscopic guidance resulting in a gastropexy of the anterior gastric wall to the ventral abdominal wall. An incision is made between the sutures and a trocar with a peel-away sheath is introduced through the abdominal wall into the stomach. Alternatively, a guidewire can be introduced using the Seldinger technique with progressive dilation of the gastrostomy. The peel-away sheath is then introduced following dilation. A feeding tube is advanced through the sheath, which is then peeled off. An inflatable balloon at the tip of the feeding tube is filled with saline and serves as the internal bumper. A skin disc secures the tube externally. The gastropexy sutures are then removed. This technique can be used in patients with high-grade stenosis in the oropharynx or along the esophagus since it allows for use of an ultra-thin endoscope. Additionally, given that gastropexy is first achieved, this technique has been safely used in patients with ascites.

Another potential advantage of the introducer technique is that it avoids passage of the feeding tube through the oropharynx, thus preventing translocation of bacteria to the peristomal site. Several studies have suggested that there is a lower rate of peristomal skin infection associated with the introducer technique as compared with the per-oral pull technique for PEG tube placement (64,65). One randomized control trial has even suggested that with the introducer method, prophylactic antibiotics may no longer be needed (66).

Percutaneous ultrasound gastrostomy (PUG) technique

PUG is an alternative to the standard PEG techniques. It utilizes magnetic coaptation and ultrasound guidance instead of endoscopic guidance. The Point-of-care Ultrasound Magnet-Aligned Gastrostomy (PUMA-G) by CoapTech™ is the first system that has been approved by the US Food and Drug Administration (FDA) for this
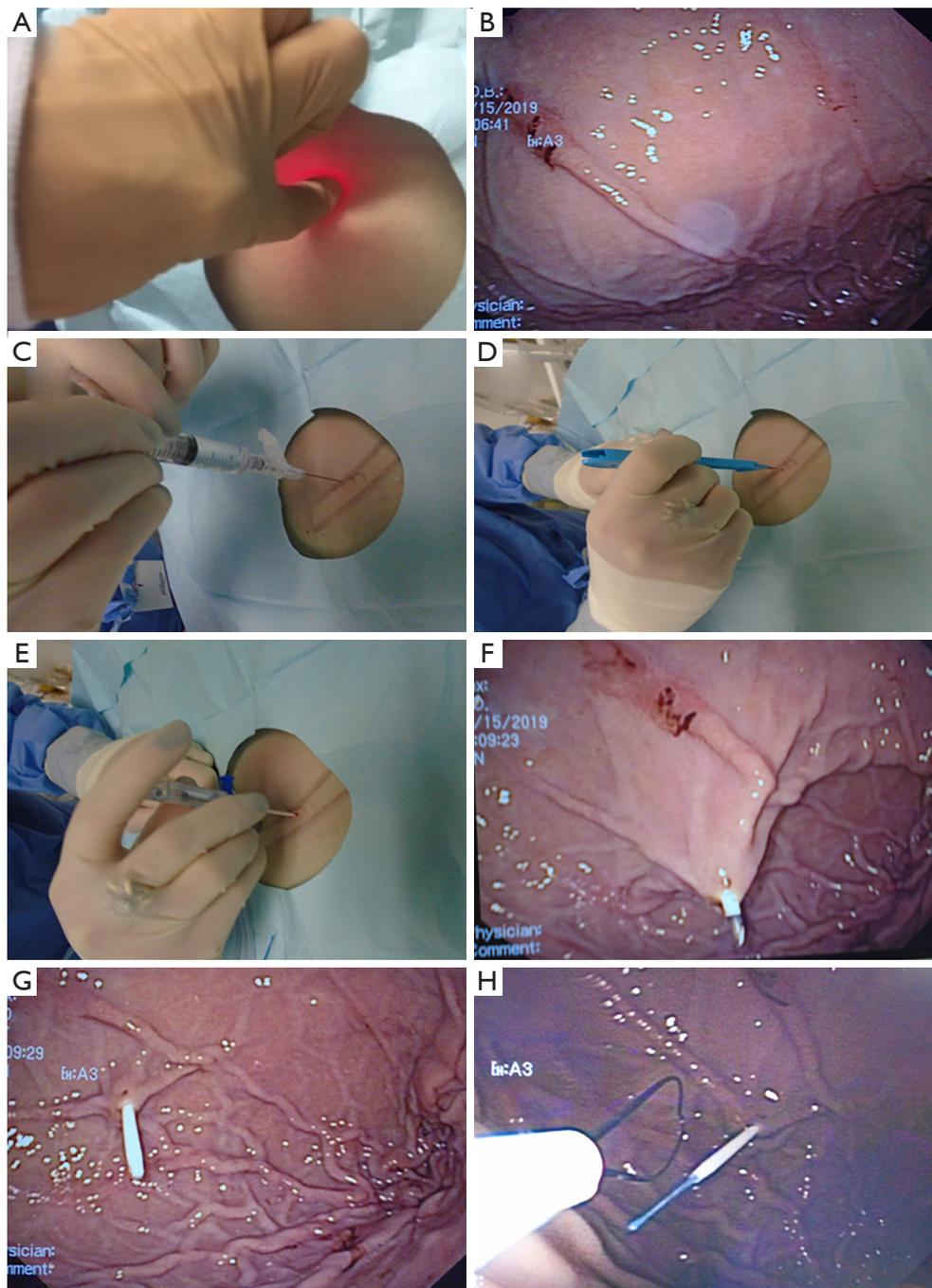
Figure 3 Insertion of the guidewire into the stomach. (A) Finger indentation at the site of trans-illumination; (B) indentation visualized from inside the stomach; (C) lidocaine with epinephrine administered at 90-degree angle to the surgical site; (D) small skin incision made with a scalpel at 90-degree angle; (E) insertion of the needle-catheter combination (trocar) through the incision at 90-degree angle; (F) visualization of the trocar from inside the stomach; (G) catheter positioned inside the stomach with the needle withdrawn; (H) snare system is passed through the endoscope and catches the wire loop.
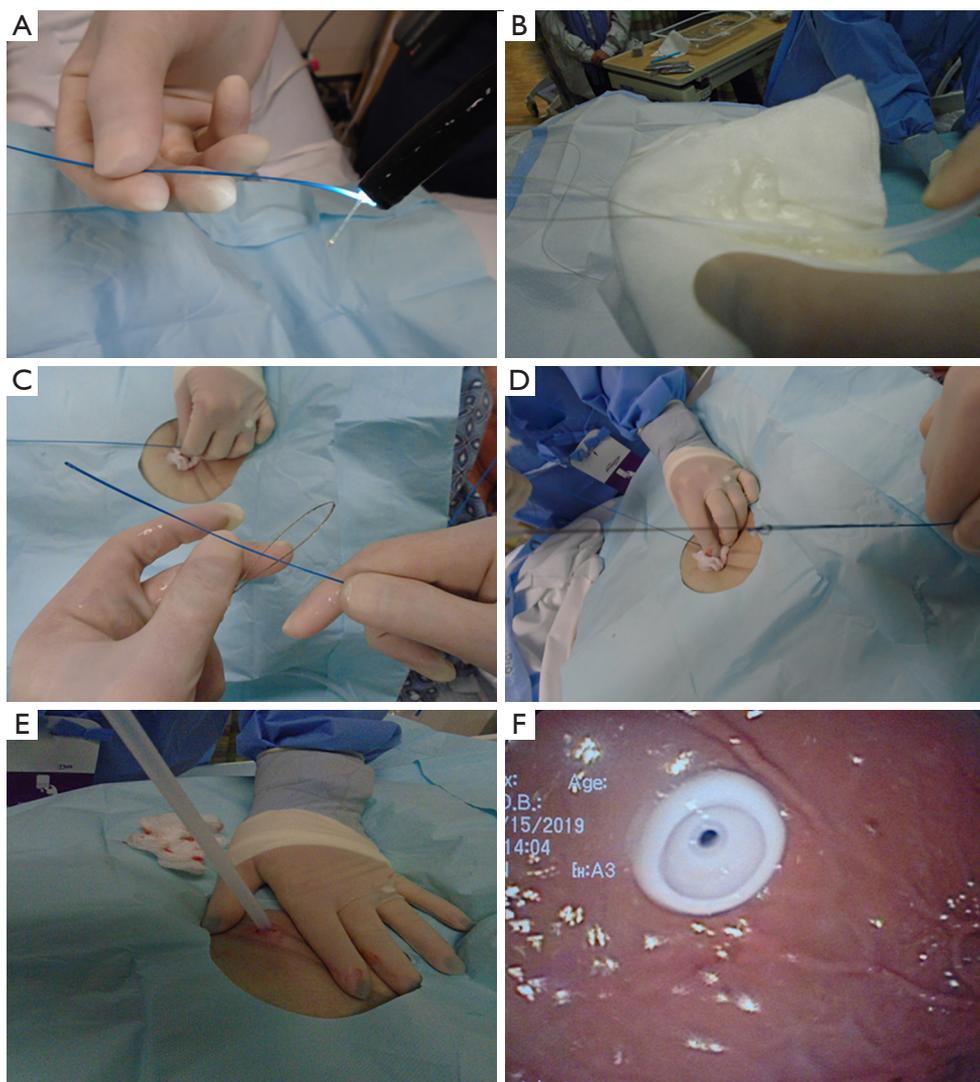
Figure 4 Placement of the PEG tube via the pull method. (A) Guidewire with wire loop pulled out of the mouth; (B) lubricating gel applied to the PEG tube prior to insertion; (C) PEG tube looped through the guidewire; (D) PEG tube attached to the guidewire via a knot; (E) firm pressure applied while pulling the PEG tube through the abdominal wall; (F) direct visualization of the PEG tube bumper from inside the stomach. PEG, percutaneous endoscopic gastrostomy.

technique (67).

In this technique, an existing orogastric or NGT in the stomach is used for insufflation. Then a specialized orogastric tube with a distal balloon, housing a bar magnet, is passed from the mouth into the stomach. A wire stylet is placed inside the tube to assist in the placement of the gastric tube balloon (GTB) within the stomach (*Figure 6*). Once the GTB is placed within the stomach, the wire stylet is removed and an external handheld magnet (EHM) is then placed over the GTB and used to move the GTB. The

magnetic force between the internal bar magnet and the EHM bring the gastric wall and abdominal wall together to achieve a magnetic gastropexy. The GTB is then filled with saline via a Luer lock at the proximal end of the gastric tube. A handheld ultrasound is used to localize the saline filled GTB. Next, an introducer needle is inserted under direct ultrasound guidance into the saline-filled GTB. Subsequently, a guidewire with a coiled tip is inserted through the needle. Upon exiting the needle, the guidewire expands into its coiled configuration. The GTB is deflated
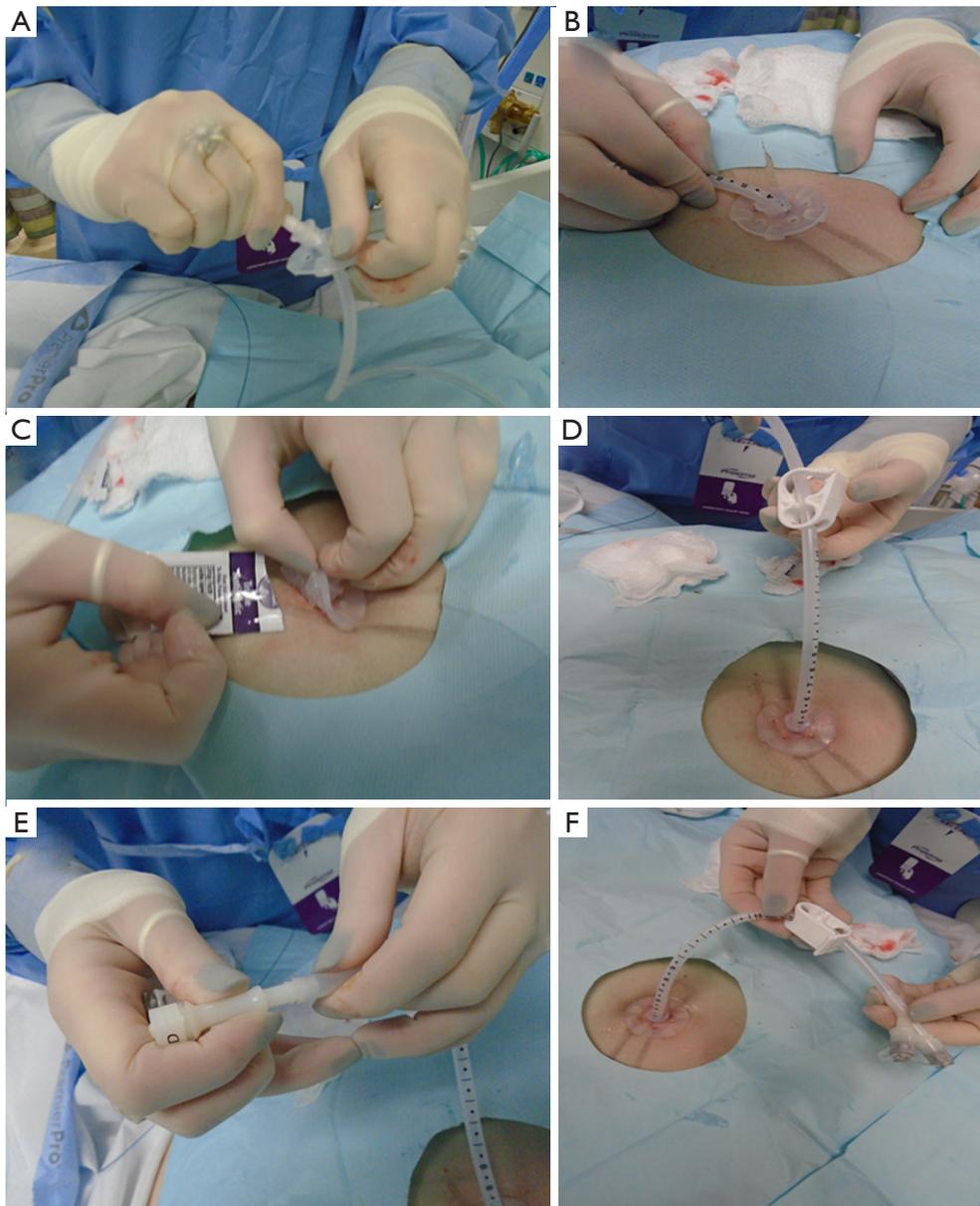
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Figure 5 Completion of the PEG tube procedure. (A) External bolster placed onto the PEG tube; (B) skin marking on the external bolster kept at 2–5 cm depending on the patient's body habitus, making sure that the bolster is neither too tight nor too loose; (C) application of antibiotic ointment at the surgical site; (D) clamp positioned onto the PEG tube; (E) external port attached to the PEG tube; (F) successful placement of the PEG tube. PEG, percutaneous endoscopic gastrostomy.

to capture the coiled end of the guidewire within the balloon. The GTB catheter and guidewire is then pulled out of the mouth and a gastrostomy tube is fed over the guidewire in similar fashion as the Sacks-Vine per-oral push technique.

The safety and efficacy of the PUMA-G technique has been demonstrated in case series of five patients. All patients had their gastrostomy tubes placed successfully, and none had complications in the 30 days following tube placement. In this early experience, the average time for gastrostomy tube placement with the PUMA-G technique compared with standard PEG tube placement techniques was longer. Additionally, fluoroscopy was required to localize the GTB within the stomach and to guide EHM placement
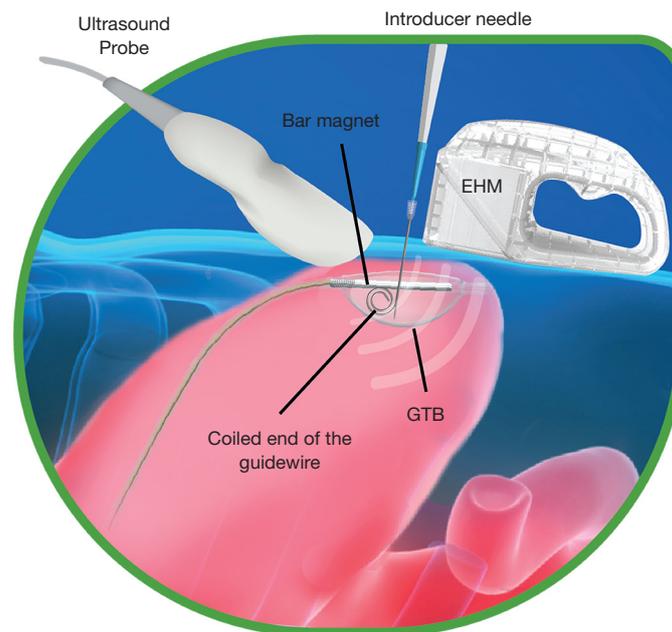
Figure 6 Illustration of the percutaneous ultrasound gastrostomy technique.

in 2 out of 5 patients, suggesting a learning curve for the operator (68). A second study (published in abstract form) also showed efficacy of the PUG with magnetic gastropexy technique, demonstrating success in 15 of 20 patients. In these patients that were evaluated, fluoroscopy was used to facilitate gastrostomy placement 95% of the time (69). However, both studies postulated that with continued operator experience PUG is feasible without fluoroscopy.

PUMA-G offers the advantage over traditional PEG tube techniques in that it can be done at the bedside with an ultrasound. The per-oral pull, per-oral push, and introducer techniques for PEG tube placement require specialized medical personnel and costly equipment. These may not be available in under-resourced settings and can be more expensive. However, further studies are needed to explore the safety and efficacy of the PUMA-G technique in comparison to standard PEG techniques. Another limitation of the PUMA-G system is that the EHM has only been FDA approved for gastropexy in patients with abdominal wall thickness ≤4.5 cm and maximal body mass index (BMI) between 30–35 kg/m^2 (67,68).

Prophylactic use of antibiotics

Patients who are candidates for PEG placement are often also at high risk of infection given malnutrition,

immunosuppression, and underlying medical comorbidities. As such, current guidelines recommend use of prophylactic antibiotics prior to PEG placement to minimize the risk of peristomal infection. In patients not allergic to penicillin (PCN), a cephalosporin or PCN antibiotic should be administered 30 minutes before the procedure. Patients should also be screened for methicillin-resistant Staphylococcus aureus (MRSA). Patients who test positive for MRSA should undergo decontamination and receive prophylactic vancomycin (70).

Contraindications

Contraindications to PEG tube placement include severe coagulopathy, colonic interposition, history of subtotal or total gastrectomy, hemodynamic instability, sepsis, severe ascites, peritonitis, abdominal wall infection at the insertion site, peritoneal carcinomatosis, gastric outlet obstruction, and history of gastric varices (71).

Obesity

Obesity, as defined by BMI >30 kg/m^2, can make trans-illumination more challenging and additional pressure is required on the abdominal wall to obtain successful trans-illumination. However, obesity is not a strict

contraindication to PEG tube placement. PEG tube placement in overweight and obese patients can be done without increased risk of complications compared to patients with normal BMI (18–25 kg/m^2) as long as there is good trans-illumination and indentation, and colonic interposition is not suspected (72,73).

Mackenzie *et al.* evaluated placement of direct percutaneous endoscopic jejunostomy (DPEJ) in obese and overweight patients (BMI >25 kg/m^2) and found decreased success with a trend towards increased complications as compared to patients with normal BMI (74). Safe and successful PEG tube placement has been documented in a case series of super-morbidly obese patients (BMI >60 kg/m^2) (75).

Pregnancy

PEG tubes are generally not used in pregnancy. This is due to the concern for uterine or fetal damage, the alteration of the intra-abdominal anatomy by the enlarging uterus, and the risk of infection. Nevertheless, there have been several case reports of successful PEG tube placement in pregnant patients with severe refractory hyperemesis gravidarum or who were comatose following trauma (76-79).

Ascites

Ascites is a relative contraindication for PEG tube placement, as it poses technical challenges for the procedure itself and increases risk of complications following the procedure. Ascitic fluid can make puncturing the anterior wall of the stomach more challenging. Ascites increases the intraperitoneal distance and can predispose to PEG tube dislodgement. Furthermore, ascitic fluid may prevent the maturation of a PEG tube tract thereby increasing risk of gastric fluid leakage and peritonitis.

However, Galaski *et al.* showed that in cases of malignant bowel obstruction, a palliative PEG tube for decompression can be performed successfully. These patients first had their ascites drained via placement of an intraperitoneal catheter, which was then followed by placement of a PEG tube. While these patients have a higher risk of complications, PEG tube placement is a feasible procedure (80,81). Furthermore, the use of gastropexy can be used to prevent ascitic leakage around the PEG tube (82,83).

Initiation of tube feeding after PEG placement

Enteral feeding is generally initiated 12–24 hours after PEG tube placement. Despite its popularity, this practice is not based on evidence from randomized controlled trials. Kirby *et al.* in 1986 published one of the earliest studies that documented the efficacy and safety of same day feeding (84). Early feeding following PEG tube placement has been demonstrated to be safe by a number of subsequent prospective studies (85-88). Early tube feeding allows for earlier achievement of goal nutritional rates, potentially reducing healthcare costs by shortening inpatient hospitalization.

One meta-analysis by Bechtold *et al.*, demonstrated that early (<4 hour) feeding is safe and well tolerated when compared to delayed or next day feeding following PEG tube placement (89). A more recent meta-analysis by Szary *et al.*, also showed that early (<3 hours) feeding is safe when compared to delayed feeding (90).

Complications

Acute complications

Patients who undergo PEG tube placement are at risk for procedural complications. While the risk is relatively low (0.1%), the procedure can be associated with significant morbidity and mortality. Possible complications from an endoscopy-guided procedure include aspiration, hemorrhage, and perforation. Sedation carries the risk of aspiration, hypotension, and hypoxia (12).

Bleeding

The post PEG-procedure bleeding risk is 2–2.5%. The current American Society for Gastrointestinal Endoscopy (ASGE) guidelines state that aspirin and low dose nonsteroidal anti-inflammatory drugs (NSAIDs) can be continued in patients at high-risk for thromboembolic disease. In patients with low thromboembolic risk, the decision to hold aspirin or NSAIDs is left to the discretion of the operator. In patients with low thromboembolic risk, thienopyridines (e.g., clopidogrel) should be held for 7–10 days prior to the procedure. In patients with high thromboembolic risk, PEG tube placement should be delayed until the thienopyridine can be safely held. However, some studies have suggested that clopidogrel can

be safely continued for PEG tube placement (91,92).

Interestingly, Richter *et al.* found that while the use of aspirin or clopidogrel was not associated with increased risk of bleeding, serotonin reuptake inhibitors have been linked to increased risk of bleeding (93).

Warfarin should be held for 4–7 days prior to the procedure. Warfarin should be bridged if patients are at higher risk for thromboembolism such as those with valvular atrial fibrillation, mechanical heart valves, left ventricular assist devices, or recent diagnosis of thromboembolism.

Bleeding can usually be controlled with pressure over the abdominal wound. In the case of delayed external bleeding at the incision site, we use topical tranexamic acid (5 mL) and oxidized regenerated cellulose (Surgicel™). If there is any concern for internal bleeding, endoscopic or surgical exploration of bleeding source may be required, and the use of endoscopic clips may be needed for hemostasis (38).

Peristomal wound infection

Peristomal wound infection is the most common complication of PEG tube placement with an incidence ranging from 4–30% (87). Three quarters of these infections are minor and resolve with antibiotic treatment (94). Several randomized control studies have shown that antibiotic prophylaxis reduces the risk of peristomal wound infection compared to placebo (95-97).

This has been confirmed by a meta-analysis, which showed that the use of prophylactic antibiotics prior to PEG placement was associated with a 64% relative risk reduction and a 15% absolute risk reduction in peristomal wound infections. The number needed to treat to prevent one wound infection was eight (98). One study showed that nasopharyngeal decolonization of MRSA significantly reduced peristomal infection rates among patient colonized by MRSA (99).

Recent double blind, randomized control studies evaluating the modified introducer technique have suggested that prophylactic antibiotics may not be needed in the modified introducer technique. This technique, as described earlier, does not require passage of the feeding tube through the mouth thus minimizing translocation of the oral flora to the stoma site, which occurs with the per-oral push and per-oral pull techniques (66,100).

Clogged feeding tube

The incidence of clogged feeding tube has been reported

as high as 23–35%. Risk factors for clogging include use of thick enteral feeding formulas, use of bulking agents, and use of smaller bore feeding tube (8–9 Fr). For this reason, we use a 20 Fr feeding tube at our institution for most cases.

One study showed that with increased acidity, particularly with pH <5.0, proteins become less soluble, increasing the risk of gastrostomy tube clogging (101). Prophylactic use of pancreatic enzymes with sodium bicarbonate mixture has shown to be effective in maintaining tube patency (102,103).

Buried bumper syndrome (BBS)

BBS is a severe complication of PEG tube placement which occurs when the internal bumper migrates along the stomal tract and is displaced outside of the gastric wall. This often occurs as a result of excessive compression between the internal bumper and external bolster. The incidence of BBS has been noted to be around 1% (0.3–2.4%). BBS can lead to gastrointestinal bleeding, perforation, peritonitis, or intra-abdominal abscess or phlegmon.

Patient specific risk factors for developing BBS include old age, immunosuppression, malnutrition, and treatment with steroids or chemotherapy. Internal bumper characteristics, PEG tube characteristics, and the quality of long term gastrostomy site care can affect the risk of developing BBS. Internal bumpers made of rigid or semi-rigid material or those with small contact area, sharp edges, or conical shape are at risk for developing BBS. Tubing with jejunal extension can cause the tubing to be positioned tangentially rather than perpendicularly, which also increases risk for developing BBS. It is recommended that as a preventative measure, the external bolster should be positioned about 10mm away from the skin. Permanent dressing between the external bolster and the skin or excessive tightening of the external bolster can increase the risk for BBS. The timing of relaxing the external bolster positioning to 10mm is controversial. Some operators prefer a tighter positioning of the external bolster in the first four days after PEG tube placement to prevent leakage at the stoma (104).

Tumor tract seeding

There is also a small but definite risk of tumor metastasis following PEG tube placement in patients with untreated head and neck cancers. One retrospective review of 304 patients with active head and neck cancer who underwent PEG placement noted a risk of 0.92% of developing

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abdominal wall metastasis (105). It has been suggested that the risk of tumor metastasis from head and neck cancer may be minimized with the use of the introducer technique for PEG placement (106).

Inadvertent tube removal

The rate of early PEG tube dislodgement has been reported as 0.6–4.0%. When followed longitudinally for the lifetime of a new PEG, PEG tube dislodgement rates were observed to be 12.8%. Dislodgement within 7 days of PEG tube placement, before the gastro-cutaneous tract has a chance to mature, results in an open gastrostomy. This allows for leakage of gastric contents and tube feeds into the intraperitoneal space, resulting in significant morbidity (107).

In patients who are disoriented or combative and at high risk of early tube dislodgement, various techniques have been described to minimize the risk of dislodgement, including endoscopic sutures, T-fasteners, and novel bumper designs (108-111).

Peristomal leakage/irritation

The reported incidence of peristomal leakage is around 1–2%, but the actual incidence is likely much higher. Risk factors for peristomal leakage include delayed wound healing, infection, gastric hypersecretion, excessive cleaning with hydrogen peroxide, BBS, and torsion or instability of the PEG tube. Patients should be optimized from a nutrition and glycemic control standpoint preceding the PEG procedure.

Skin protectants and barrier creams such as zinc oxide should be applied. Excessive torsion applied by the external bumper should be relieved. Proton pump inhibitors should be initiated to minimize gastric acid secretion.

It is not recommended to exchange the PEG tube to a larger bore tube as this would result in enlarging the gastro-cutaneous fistula and potentially exacerbate the problem. One strategy is to remove the PEG tube and allow the fistula to close before placing a new PEG tube. However, this can only be done once the gastro-cutaneous fistula tract has fully matured. As a last resort, the PEG tube can be removed and another PEG tube can be placed at a different site (112,113).

Gastro-colocutaneous fistulas

Gastro-colocutaneous fistula can arise from inadvertent puncture of overlying bowel or erosion over time into adjacent intestine (112,114). Risk factors include insufficient gastric insufflation, inadequate trans-illumination, or presence of adhesions which may result in trapped intestines. Patients may present acutely with peritonitis, fasciitis, or intestinal obstruction. However, more commonly the presentation is insidious, with leakage of fecal contents through the peristomal site or with watery diarrhea resembling tube feed formula. If a fistula is suspected, the patient should have a contrast study to confirm the anatomy (115). The fistulous tract can be managed conservatively by removal of the PEG tube and allowing for spontaneous fistula closure. In patients with large fistulas or delayed wound healing, the fistula can be closed surgically or endoscopically (116,117).

Interventional pulmonologists

Interventional pulmonologists have expertise with endoscopic and procedural interventions. They are comfortable placing tracheostomies, which often are required in patients who simultaneously require PEG tube placement (118,119). Therefore, the next logical evolution would be for interventional pulmonologists to perform both procedures at the same time when indicated. This would minimize the risks associated with sedation and paralytics as the two procedures can be merged into a single coordinated procedure performed by the same physician.

One small retrospective study demonstrated that interventional pulmonologists were able to place PEG tubes at bedside with 97.2% success rate (70 of 72 patients). Forty-one of the 70 PEG tubes placed (58%) were performed immediately after percutaneous tracheostomy. There were no complications associated with PEG tube placement. Mild cellulitis around the PEG tube site was noted in 1 patient (1.4%). This study found a 30-day mortality rate of 11.94% as compared with 25% in gastroenterology literature (120).

Another prospective study demonstrated similar efficacy and safety of PEG tube placement with bronchoscopic guidance in the ICU by interventional pulmonologists with a 97.6% success rate and 2.4% complication rate. Only minor complications including soft tissue infection and G-tube malfunction were noted. No major complications were noted. 30-day mortality in this study was 11.9% (121).

However, we prefer endoscopic rather than bronchoscopic assisted PEG placement since complete EGD is not possible with a bronchoscope. EGD is necessary

to rule out post-pyloric obstruction. Additionally, adequate insufflation and trans-illumination with examination of the gastric cardia by retroflexion is not always possible with a bronchoscope.

Conclusions

Gastroenteric feeding plays an important role in the critically ill patient who is unable to take sufficient oral nutrition. PEG tube placement is an increasingly common procedure and frequently coincides with tracheostomy placement. Interventional pulmonologists already have expertise in percutaneous tracheostomy placement, and they can also place PEG tube safely and effectively. We believe that interventional pulmonary trainees should be educated in PEG techniques, allowing for simultaneous percutaneous tracheotomy and PEG tube placement.

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