

REVIEW

Percutaneous endoscopic gastrostomy: An update on its indications, management, complications, and care

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ABSTRACT

Background: Numerous disorders impairing or diminishing a patient's ability to swallow may benefit from a PEG tube placement. This is considered the elective feeding technique if a functional digestive system is present.

Methods: A PubMed-based search restricted to the English literature from the last 20 years was conducted. References in the results were also reviewed to identify potential sources of information.

Results: PEG feeding has consistently demonstrated to be more effective and safe than nasogastric tube feeding, having also replaced surgical and radiological gastrostomy techniques for long term feeding. PEG is considered a minimally invasive procedure to ensure an adequate source for enteral nutrition in institutionalized and at home patients. Acute and chronic conditions associated with risk of malnutrition and dysphagia benefit from PEG placement: Beyond degenerative neuro-muscular disorders, an increasing body of evidence supports the advantages of PEG tubes in patients with head and neck cancer and in a wide range of situations in pediatric settings.

The safety of PEG placement under antithrombotic medication is discussed. While antibiotic prophylaxis reduces peristomal wound infection rates, co-trimoxazole solutions administered through a newly inserted catheter constitutes an alternative to intravenous antibiotics. Early feeding (3-6 hours) after PEG placement firmly supports on safety evidences, additionally resulting in reduced costs and hospital stays. Complications of PEG are rare and the majority prevented with appropriated nursing cares.

Conclusions: PEG feeding provides the most valuable access for nutrition in patients with a functional gastrointestinal system. Its high effectiveness, safety and reduced cost underlie increasing worldwide popularity.

Key words: PEG. Gastrostomy. Tube feeding. Enteral nutrition. Gastric feeding tubes. Intubation. Gastrointestinal. Nursing care. Complication.

INTRODUCTION

Percutaneous endoscopic gastrostomies (PEG), first described in 1980, have become widely used to provide enteral nutritional support to patients who are unable to ingest solid or liquid foods due to many disorders, despite having preserved absorption and motility functions of the gastrointestinal tract. In these cases, PEG tubes have arisen as an alternative to artificial parenteral nutrition and especially to nasogastric tubes, for the administration of food directly into the stomach (which is recognized as the most suitable and physiological feeding option).

PEG placement is an endoscopic technique that allows the placement of a flexible tube to create a temporary or permanent communication between the abdominal wall and the gastric cavity, ensuring the direct passing of food into the patient's digestive tract.

Even when the use of PEG tube feeding has not been universally demonstrated to decrease risks of aspiration pneumonia (1) or long term mortality, nor outcomes regarding to weight maintenance when compared with nasogastric tube feeding in several groups of patients (2), PEG feeding has been consistently demonstrated to be the feeding method with a lower probability of intervention failure, suggesting the endoscopic procedure is more effective and safe than nasogastric tube feeding, according with a Cochrane systematic review (3).

Since Ponsky and Gauderer described this technique (4), PEG tubes have replaced other surgical (5) and radiological (6) gastrostomy techniques as the method of choice for long term feeding of patients who are unable to maintain adequate nutrition in the presence of a normal gastro-

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intestinal functioning. As a result, PEG use is recognized as a minimally invasive procedure that eliminates the need for general anesthesia and requires less instrumentation, it is therefore a valuable source of nutrition by enteral feeding in nursing homes and domiciliary environments (7) when the administration period is expected to exceed 4 weeks and life expectancy of patients exceeds two months (8). It is favored by its simplicity, usefulness, safety, ease of operation and low cost (4).

This article aims to review current evidence of the indications for and advantages of PEG tube placement in variety of settings and pathological conditions. Placement techniques and procedural management of PEG tubes will also be explained and risks and potential complications discussed. Finally, specific nursing care will be provided.

A PubMed library-based search was carried out for the period between 1990 and July 2014, using the following individual and combined key words: PEG tube, PEG tube feeding, complications, diet, dietary intervention, dietary treatment, enteral or parenteral nutrition, and risk factors. References cited in the articles obtained were also searched in order to identify other potential sources of information. The results were limited to human studies available in English.

INDICATIONS FOR PERCUTANEOUS ENDOSCOPIC GASTROSTOMY

The option to feed a patient through a PEG tube should be considered in different situations, both in hospital and at home (9). In fact, several acute and chronic conditions may be alleviated by feeding sufferers with an intact digestive tract through a PEG tube. A reduction in oral intake, generally due to neurodegenerative processes (10) represents the main reason for PEG placement in up to 90 % of cases. However, PEG tube feeding in dementia patients has been largely controversial: The extensive use of these devices in situations of oral nutrition failure contrast with of the lack of proven benefits in patients with advanced dementia, that were not demonstrated in a systematic review that included seven observational studies (10): There were no evidences of increased survival, improvement of nutritional status or reduction of pressure ulcers prevalence rates in patients receiving enteral tube feeding. Therefore, the final decision for PEG tube placement in patients with dementia and other neurodegenerative diseases should be assessed between the physician, family and caregivers, bearing in mind the patient's advance directives (11).

Additionally, a repeated bronchial aspiration of food, or obstruction derived from oropharyngeal, neck or esophageal tumors (12) are other common indications. Table I includes the most frequent indications for PEG placement, classifying patients according to the chronicity of underlying diseases and its ability to recovery.

An increasing body of literature is documenting the potential value of prophylactic PEG tube placement at

treatment initiation in patients with head and neck cancer, who are at increased risk of malnutrition and dysphagia (13). In these patients, enteral tube feeding is often required in response to dysphagia, odynophagia or other side effects of treatment that lead to dehydration and/or weight-loss during or after cancer treatment. The majority of studies published in the literature generally commence nutrition support by a PEG tube when clinically indicated in response to deterioration in swallowing or nutritional status (14-16). In contrast, some studies have reported on the commencement of enteral feeds prior to treatment (17-20), showing that prophylactic PEG placement and early tube enteral feeding was associated with a limited loss of weight, allowing an effective and safe nutrition and hydration of the patient during chemoradiation, according to retrospective chart reviews (16,18); Additionally, patients who require therapeutic PEG tube placement in response to significant weight loss during treatment suffered greater morbidity than patients who received PEG tubes prophylactically (21).

The evidence to clearly support the early placement and use of a PEG tube in patients undergoing treatment for head and neck cancer is weak however and the benefits versus risks have not been definitely established (22). Increasing concern that gastrostomy placement leads to prolonged tube dependency and long term dysphagia exist (23,24).

An ongoing randomized controlled trial (RCT) aimed at assessing the nutritional and clinical outcomes of patients with head and neck cancer undergoing prophylactic gastrostomy prior to treatment compared with standard practice of commencement of tube feeding (25) will shed light on this particular topic.

In the pediatric population, PEG insertion for enteral nutrition has become widely accepted, after having been demonstrated as an efficient and safe technique even in small infants, and associated with an acceptable rate of complications (26). A range of experience from clinical showing an improvement in or maintenance of adequate nutritional status in patients with a variety of underlying disorders (as well as a high level of acceptance by caregivers), has been reflected in the rising number of medical conditions for which PEG feeding is indicated in children. These include not only neurological disorders, or congenital malformations leading to oropharyngeal dysphagia, but also medical and surgical conditions impairing an adequate caloric intake, special feeding requirements (i.e. unpalatable formula in multiple food allergies or metabolic diseases) or the need for continuous enteral feeding in short bowel syndrome and malabsorption.

The use of PEG feeding in pediatric oncology has increased in last few years. In these particular situations of early PEG feeding, PEG placement is able to reverse weight loss (27) and represents a relatively safe way to prevent malnutrition in children with cancer, and subsequently might play a role in the oncological outcome (28).

Table I. Indications for the placement of a percutaneous endoscopic gastrostomy

<i>Indications</i>
<p><i>I. Patients with potentially reversible diseases in which it is expected that the PEG can be removed once the process is solved:</i></p> <p>Neurological diseases: Guillain-Barre syndrome, stroke, cranial trauma Anorexia nervosa Hyperemesis gravidarum Severe burns Multiple injuries and facial trauma Transplants prior malnutrition Head and neck tumors treated with chemotherapy and radiotherapy Diseases of the esophagus</p> <p><i>II. Patients with irreversible diseases with prolonged survival in which the PEG is placed permanently and helps improve their quality of life:</i></p> <p>Neurological diseases: ALS, multiple sclerosis, dementia, Parkinson's disease, Alzheimer's disease, stroke, post-anoxic encephalopathy, brain metastases, brain tumours, poliomyelitis, brain injury (traumatic or surgical) Progressive muscular dystrophy Head and neck tumors Facial malformations and oropharyngeal Neoplasms of the esophagus and cardias Oropharynx tumors Dermatomyositis and polymyositis Amyloidosis Cystic fibrosis Short bowel syndrome Inflammatory bowel disease Scleroderma</p> <p><i>III. Patients with terminal and debilitating diseases with a relatively long life expectancy (this indication should be individualized and consensual):</i></p> <p>Encephalitis Repeated stroke Advanced malignancies AIDS terminal stages Intestinal obstruction by peritoneal carcinomatosis Radiation enteritis Severe acute pancreatitis</p> <p><i>IV. Preventing malnutrition in pediatric illnesses:</i></p> <p>Chemotherapy in oncologic disease. Unpalatable formula in multiple food allergies Inadequate caloric intake Multiple congenital malformations Short bowel syndrome Oropharyngeal dysmotility Epidermolysis bullosa Unpalatable medications in renal failure</p> <p><i>V. Improving morbidity in patients undergoing radiotherapy for head and neck carcinomas</i></p>

ALS: Amyotrophic lateral sclerosis; AIDS: Acquired immune deficiency syndrome.

CONTRAINDICATIONS FOR PEG PLACEMENT

There are few absolute contraindications to PEG placement, and these mainly include technical limitations as a result of anatomical particularities such as lack of transillumination with an inability to access the

anterior gastric wall, including colonic interposition and severe ascitis, uncorrectable advanced coagulopathy, portal hypertension with significant gastric varices leading to unassumable risk of bleeding; finally, pharyngeal or esophageal obstruction blocking the passage of the gastroscope to the stomach will prevent a PEG

tube placement. The remaining are considered relative contraindications (Table II).

Prior abdominal surgery is currently not considered as a contraindication to PEG placement, with clinical studies showing that it can be safely placed in these patients (29), with a high success rate (30). Gastric surgery may represent a unique challenge to the endoscopist, with a 28 % of placement failures recorded in a retrospective report (29).

PREPARING THE PATIENT FOR A PEG TUBE PLACEMENT

Informed consent

Informed consent should be obtained from patients or their legal surrogate decision makers in a consensuated way with by health professionals. Patients with advanced dementia and dysphagia usually undergo to PEG placement, so consent for a treatment in a patient without legal capacity should be guaranteed from nominated legal substitutes. The intention of informed consent is to enhance the patient's care by providing them or their caregiver with complete information on the benefits and risks of tube feeding and medications before PEG insertion (31).

Antiplatelet and anticoagulant medication

PEG is classified as an invasive interventional endoscopic procedure (32) that can result in bleeding, a complication that has been reported in approximately 2.5 % of procedures in the early literature (33,34). Patients undergoing PEG are commonly treated with aspirin and/or other antithrombotic agents, which are commonly used for treating or preventing several cardio- and cerebrovascular diseases. A major dilemma concerning patients taking these medications includes the potential risk of bleeding as a result of endoscopic intervention and the risk of thromboembolic events when such medications are withheld. Recent guidelines from the American Society of Gastrointestinal Endoscopy (ASGE) published in 2009 for the use of anticoagulant and antiplatelet therapy for endoscopic procedures recommends that patients who are taking clopidogrel or ticlopidine should have these medications discontinued 7-10 days before PEG placement; with regard to aspirin and others non-steroidal anti-inflammatory drugs (NSAIDs) endoscopic procedures may be performed while the patient is receiving this medication in the absence of a pre-existing bleeding diathesis (35). However, the ASGE guidelines are based on expert opinion and best clinical practice, since no prospective RCT trials to support them are available.

Several recent large retrospective cohort studies have been carried out to determine whether there is an association between periprocedural aspirin, clopidogrel, or

Table II. Contraindications for the placement of a percutaneous endoscopic gastrostomy

Contraindications
<i>I. Due to local problems:</i>
Nonswelling esophageal obstruction
Active gastric pathology
Total gastrectomy
Extreme obesity
Previous midline laparotomy (can hinder the location of the puncture site)
<i>II. Absolute contraindications:</i>
Colonic interposition
Partial or subtotal gastrectomy
Massive ascites
Portal hypertension (gastric varices)
Peritoneal dialysis
Active gastric pathology
Coagulation disorders
Sepsis
Cardiorespiratory disease that prevents the endoscopy
Pyloric stenosis
Expected survival < 2 months (NGT is preferred)

NGT: Nasogastric tube.

ticlopidine use and bleeding in patients who underwent PEG tube placement. According to these studies, post-PEG bleeding were rare events (0 % to 2.8 %), and the use of aspirin or clopidogrel before or after PEG was not associated with procedure-related bleeding in any study (36-40). The use of dual antiplatelet therapy was not a risk factor for postprocedure bleeding, according to a retrospective multicenter study (39).

Regarding to anticoagulation, ASGE guidelines recommend that warfarin should be discontinued 3-5 days before the procedure and bridged with low molecular weight heparins (LMWH) or unfractionated heparin (UFH) in the case of a high risk of thromboembolic complications. LMWH should be discontinued at least 8 hours before the PEG procedure; UFH infusion is recommended to be discontinued 4-6 hours before PEG and restarted 2-6 hours after the procedure is completed (35). The safety of these recommendations has been demonstrated, since the use of LMWH did not increase the risk of bleeding in the aforementioned observational studies (37). In addition, one study have suggested that patients undergoing therapeutic anticoagulation or those with increased INR values have no elevated risk of bleeding during PEG placement (40).

The safety of maintaining antiplatelet therapy in a PEG placement tube should be evaluated with further RCT, but available data supporting the individual decision to maintaining these drugs in those patients for whom the thrombotic risk is high if withdrawn, cannot be afforded.

Preventing peristomal infection

Although PEG is considered a relatively minor surgical procedure, it is associated with general complications, among which wound infection is the most common problem. The placement of a PEG tube is not considered a sterile technique and patients undergoing to it are often vulnerable to infection for a variety of reasons including old age, compromised nutritional intake, immunosuppression and underlying disease such as malignancy and diabetes (41). Bacteria colonizing the nasopharyngeal and upper digestive tract may cause peristomal infection in PEG placement using the pull technique (42), a complication that is described with a frequency of up to 32 % without antibiotic prophylaxis (33,43,44).

A systematic review with meta-analysis of RCT demonstrated a significant reduction in the incidence of peristomal infection when intravenous prophylactic antibiotics were administered (pooled OR 0.31, 95 % CI, 0.22-0.44) (45). The most commonly used antibiotics to prevent peristomal infection are intravenously administered betalactams, including co-amoxiclav, cefotaxime, cefoxitin or cefazolin, prior to PEG. A recent RCT however, comparing the administration of 20 mL of co-trimoxazole solution deposited in a newly inserted PEG catheter to cefuroxime prophylaxis given intravenously before PEG was at least as effective at preventing wound infections (46).

PEG PLACEMENT TECHNIQUE

Patient preparation

After fasting for at least 6 hours and having a recent normal blood coagulation analysis, the medication the patient receives should be checked, especially regarding the suspension of anticoagulants or antiplatelets, if needed. A venous access should be channeled, and to prevent septic complications, broad spectrum antibiotics should be administered intravenously 30 minutes before, unless a 20 mL liquid solution of co-trimoxazole is going to be deposited through the PEG tube immediately after being inserted (47).

The abdominal skin should be shaved if needed and disinfected with a colorless disinfectant. Dentures must be removed and oral secretions vacuumed if necessary. After this, cleaning and disinfection of the oropharyngeal cavity is required, by using a swab with a suitable antiseptic solution.

Materials

The PEG device is usually marketed as a kit, including: Syringe and needle, scalpel, trocar, thread-guide, tube and snare. In addition to this material, medication for sedoan-

algnesia and local anesthesia should be provided, together with the tools to administer them and to aspirate oropharyngeal secretions if required.

Placement technique

To insert a PEG tube usually requires a team of 3 people (generally 2 endoscopists/gastroenterologists and a nurse). The patient is placed supine, monitored, and oxygen by nasal cannula administered. After disinfecting the abdominal wall to create a sterile field, the patients should undergo to a complete esophago-gastro-duodenoscopy (EGD), with maximal air/carbon dioxide insufflation for the extension of the wall of the stomach. The exact site of PEG insertion is determined by gastroscopic transillumination and manual palpation from outside for visualized confirmation of the appropriate placement into the lower part of the stomach.

The insertion site of the PEG tube is ideally in the median line (linea alba) to prevent hematoma and infections in the rectus muscle compartments. Next, a needle will be entering through the skin into the stomach at the location where the PEG tube is to be placed.

Three different methods have been described for PEG tube insertion: The two most widely established techniques are the pull-through method –initially described by Sacks & Vine (48), and the push method– originally described by Gauderer and Ponsky (4), due to their safety and effectiveness. In both cases, the endoscope enters through the mouth of the patient to the stomach to localize the best point to place the tube. Next, in the pull-through method, a needle is entered through the skin into the stomach at the location where the PEG tube is to be placed. A pull wire is introduced into the stomach and detected with an endoscopic snare or forceps. Then, the endoscope is slowly withdrawn until the wire appears at the mouth of the patient, and fixed to the PEG device. The PEG tube is introduced through the mouth into the stomach; indicated by meeting a resistance in the inner part of the tube to reach its final position, appearing from inside of the stomach.

The push method requires the puncture of the stomach with a double gastropexy scalpel performed under general anesthesia, with a distance of 2 cm between the two points. Between these two fixations, a puncture cannula is advanced into the stomach and a feeding tube is inserted through it. Thereafter, the puncture cannula is removed. The intragastral fixation balloon is filled with a syringe with saline solution to prevent a dislocation. Gastropexy sutures will be removed after some days. This technique avoids the passage of the PEG tube along the patient's upper aero-digestive tract.

The third method for PEG insertion, described by Russell (49), consists of inserting the tube through the abdominal wall after using stents and should be considered when the passage of the tube through the mouth needs to be prevented.

Several retrospective series have compared the pull-through and push methods (50-53). In general, pull-through PEG carried out by endoscopic teams were technically easier; push PEG showed an overall significantly higher rate of complications, dislocations and occlusions, but not in patients with advanced head, neck and esophageal cancer, among whom push-PEGs are preferred. As such, the final decision as to which PEG tube should be used depends on individual conditions.

A repeat endoscopic monitoring to determine optimal placement and to ensure the absence of immediate complication is always recommended; in particular, to set the external bumper under direct vision, which is paramount to prevent a buried bumper syndrome (BBS) (54).

Patient care after PEG placement

It is recommended to take bed rest for at least 6 hours after placement and to monitor closely all vital signs as well as any occurrence of abdominal pain, fever or gastrointestinal bleeding. It is advisable to keep a peripheral venous line inserted for at least 6 hours in case complications arise. Additionally, some analgesia may be required during the first two days, especially in the case of children (55).

THE MOMENT FOR INITIATING PEG FEEDING

Feeding through PEG tubes have traditionally been delayed until the following day after its placement due to the fear of immediate post procedural complications, including peritoneal leakage and bleeding. Several observational studies however (56,57), RCTs (58,59) and a systematic review with a meta-analysis (60) have evaluated the differences between early feeding (i.e. starting liquid and/or nutritional formula administrations in the first 3 to 6 hours after placement) compared with delayed feeding (i.e. from 12 hours after insertion up to the following day). In the case of early feeding, no significant differences in local infections, diarrhea, bleeding, GERD, fever, vomiting, stomatitis, leakage, and death were noted among patients. Furthermore, in addition to early feeding being safe and well tolerated, it also results in a reduction of costs and a decrease in hospitalization.

Parallel results have been also reproduced among pediatric patients (61). Therefore, early feeding through PEG tube is recommended as it provides the patient and health-care systems with the safest and most cost-effective results.

COMPLICATIONS OF PEG

The insertion of a PEG tube is a safe method with few complications (that are clinically minor and easily

resolved). The incidence rates for serious and minor complication have been estimated to be 3 % and 6 %, respectively. Immediate mortality after the procedure appears to be less than 1% (62,63). Table III describes the most common complications, their causes and measures for resolution.

Identifying risk factors for complications

There are several retrospective reports raising awareness of the risk factors for PEG-related complications, with the aim of decreasing patient discomfort and healthcare costs (64-70). Among the non-modifiable risk factors, advanced age is recognized as increasing the risk of death after PEG insertion in 1 %/year (65); specifically an age of more than 75 years has been identified as a predictive factor for early death 1 month after PEG placement (OR = 2.49; 95 % CI = 1.47-4.21) (64). Malnutrition, expressed both as a decreased body mass index and low serum albumin levels, is repeatedly associated with a high mortality and high complication rate after PEG, as well as the presence of comorbidities. In fact, the subrogates high C-reactive protein (CRP) levels and abnormal leukocyte counts were related with an increased early mortality rate (66). The coexistence of congestive heart failure, renal failure, urinary tract infection, previous aspiration, chronic pulmonary disease, coagulopathy, circulation disorders, metastatic cancer, and liver disease were all of them strongly associated with an increased mortality. The sum of several risk factors in the same patient also greatly increases the likelihood of early death after insertion of a PEG tube; thus, the presence of 3 risk factors multiplied by 6 increases the probability of death at 1 month compared to patients who had no risk factors (64).

The risk of complications, including death, should always be assessed individually in each patient undergoing PEG tube insertions; however, we must always bear in mind that enteral feeding is superior to parenteral feeding in the nutritionally depleted patient, and PEG feeding remains the safer, easier and less expensive method for tube feeding for a wide range of severely compromised patients. Indeed, the indication for PEG is strongly associated itself with mortality (65).

PEG tube placement by an inexperienced endoscopist has been identified as a modifiable risk factor related to early complications. Furthermore, the insertion of the internal bumper of a PEG tube in the upper body of the stomach also was a significant risk for early and late complications (66).

Interestingly, some recent multicenter retrospective research has shown that proton pump inhibitors (PPIs) users (defined as patients who were taking standard doses of PPIs at least 48 hours before PEG placement) were associated with adverse PEG-related complications (including mortality, bowel perforation, post-procedural gastrointestinal bleeding, peritonitis, fever, pneumonia, peristomal

Table III. Complications of PEG: Causes and attitudes of resolution

<i>Problem</i>	<i>Possible cause</i>	<i>Attitude</i>
Necrotizing fasciitis	Necrosis of the superficial fascia	Broad-spectrum antibiotics. Surgical debridement
Bleeding from the puncture site or the gastric mucosa	A surrounding vessel injury	Producing compressive hemostasis by increasing the traction from the tube. If it does not stop, remove the tube and undergo to endoscopic coagulation
Aspiration	Aspiration of refluxed content from the stomach	Prevent it with postural treatment. Applying feeding technique correctly. If this happens, feeding should be stopped, respiratory therapy started and antibiotics should be prescribed
Irritation or infection in the skin around the stoma	Excessive pressure on the stoma. Lack of periostomal hygiene. Output gastric fluid	Adjust the distance between the external retention ring and the stoma. Clean the stoma following the rules indicated. Put gauze below the retention ring and change it daily. Consult an expert physician
Obstruction of the PEG tube	Dried food or drug product clogging inside the probe. Lack of flushing water after and between administering food or medication	Always flush water after administration of food or drugs. Flushing with warm water and carbonated beverages with a syringe. Not to place objects through the lumen in an attempt to dislodge a clog, preventing the tube rupture or perforation of the stomach. Using pancreatic enzymes mixed with bicarbonate solution If not enough, proceed to change the tube
Tube extraction	The PEG tube comes out of accidental or voluntary	Immediately replace the tube. If not immediately available, place a Foley catheter temporarily
The tube cannot be rotated	Burial of the tube in the abdominal wall	Rotate and push the probe gently inward. If not turn, remove and substitute the tube
Nausea and/or vomiting	High osmolarity of the formula. Infusion excessively fast. Lactose-intolerance. Excessive fat content in the diet	Appropriate dilute the formula. Return to previous infusion rate. Manage lactose-free diets. Use low-fat diets
Diarrhea	Hyperosmolar solution. Deficit lactose. Poor absorption of fats. Diet-cold	Use isotonic diets and/or dilute the hypertonic ones. Suppress lactose. Use low-fat formulas
Constipation	Low fluid administration. Insufficient fiber intake	Administering fluids in adequate amounts. Increasing the amount of fiber in the nutritional formula
Peristomal granuloma	Proliferation of granulation tissue through the stoma	Resection and/or cauterization of tissue

leaks, or infection) when compared with patients non PPIs users (71).

Head and neck cancer patients have a higher risk for procedure related mortality following gastrostomy than mixed patient populations, according to a systematic review specifically conducted to defined the optimum technique for gastrostomy placement in this particular patients (72). This research also showed that major complication rates following radiologically inserted gastrostomy were greater than those following PEG in patients with head and neck cancer.

Removal and replacement of the PEG

After 2-3 weeks of being placed, a fistulous gastrocutaneous tract is formed, allowing the easy removal of the gastrostomy tube. A PEG tube can be removed when the reason for its placement has been resolved: in these cases, the gastrocutaneous fistula will spontaneously close after 24-72 hours. Most of PEG tubes, however, are placed due to chronic or progressive disorders, so the tube should be periodically replaced, after a half-life of

3-6 months that can be extended up to 12-18 months if properly cared for.

A PEG tube can be removed by strong and sustained traction until the internal bumper goes through the stoma (percutaneous method); alternatively, the tube can be removed with aid of endoscopy, by linking the gastric bumper of the tube with a polypectomy snare (endoscopic method). A recent observational retrospective study has analyzed the advantages of both methods of PEG removal in terms of associated complications (73): The immediate complication rate was lower with the percutaneous removal method, with no significant differences in the late complications rate between the two methods. Peristomal bleeding was not associated with antiplatelet or warfarin use, age, gender, or short interval tube replacement. In contrast, old age was a significant risk factor of mechanical complication during PEG tube replacement (OR, 3.83; 95 % CI, 1.04-14.07, $p = 0.043$). The authors concluded that the percutaneous method may be safer and more feasible for replacing PEG tubes in older patients in order to prevent such mechanical complications as esophageal injury. These results should be further validated with prospective RCTs.

Subsequently, a replacement gastrostomy tube is inserted through the stoma into the stomach and the balloon in its tip is filled in with saline or methylene blue (between 6 and 20 mL, depending on the manufacturer and model); the tube is fixed externally with a retention ring.

The substitution of a PEG tube is an easy technique that should be learned by primary care professionals, to reduce economic costs, patient anxiety and of their caregivers (thus providing greater comfort) (74).

In case of tube removal –accidental or intentional–, its early re-implantation is a priority in order to avoid the closure of the gastrocutaneous fistula. Where immediate accessing to an Endoscopy Unit is not possible, or the necessary equipment is not available, a Foley-type catheter with an inflated balloon in the gastric lumen can be used to preserve the tract and to ensure the nutrition and hydration of the patient.

“Buried bumper syndrome”: A potentially fatal complication

Buried bumper syndrome (BBS) is an uncommon and late complication of PEG (with most cases occurring from months to years after placement) that occurs when the internal bumper of the PEG tube erodes into the gastric wall and lodges itself between the gastric wall and the skin. If not adverted, it can lead to a variety of additional severe complications, including wound infection, peritonitis, and necrotizing fasciitis (75,76). The most common management of BBS consists of removing the PEG tube smoothly, by external traction and replacing it with a new PEG tube using the pull-through method or balloon replacement tube after dilation of the old tract (77). An alternative and successful endoscopic method has also been described, that

consists of introducing a conventional papillotome over a wire into the stomach, drawing it back as far as possible, and making incisions in all four directions to advance the tube with the internal bumper into the stomach (78,79).

Concern over BBS in the endoscopic literature, however, has led increasingly to recommendations for loose placement of the external bolster. It should be noted that leaving the external bolster too loose at the time of PEG placement increases the risk of leakage and peritonitis (54), due to internal leakage of gastrointestinal secretions and enteral formula into the peritoneal cavity. In almost all cases, the technique of PEG placement itself brings the gastric and anterior abdominal walls into apposition, forming a seal, which is also ensured by the contraction of the thick gastric musculature around the PEG tube (80).

CARE OF THE PATIENT WITH A PEG TUBE

Proper long lasting care is essential in avoiding PEG-related complications, in guaranteeing the correct nutritional status of the patient and in ensuring an extended half-life for the tube. Nursing care should include three distinct aspects.

PEG tube care

As highlighted above, the PEG tube may be used immediately after insertion, but it is recommended to wait approximately 3-6 hours before administering solutions to the patient however, in order to observe any early complication, in particular bleeding. Small amounts of water and nutritional formula should be administered and progressively increased up to the fully prescribed volume within a 2-3 days period (81).

The tube and its components (plugs and retention rings), should be cleaned daily with a swab, mild soap and warm water, rinsing and drying well after being used. The caps will remain closed when the tube is not in use. Checking periodically for proper inflation of the balloon in replacement tubes is also necessary.

To avoid injury from decubitus over the abdominal and gastric walls, the tube should be daily rotated, clockwise and counterclockwise. Daily monitoring to ensure that the external support does not press onto the patient's skin is required, as is changing the mounting location of the tube. A dressing between the skin and external fixation should not be placed, as this would cause undue pressure. Only in cases where drainage is present, a dressing may be used, but should be changed frequently when soiled.

Stoma care

During the first two weeks after PEG insertion, the peristomal area should be clean daily with soft soap and water,

from the inside out, drying well, and disinfecting with anti-septic and sterile gauze around the stoma –checking that there is no irritation, inflammation or gastric secretions. A small liquid drainage from granulation tissue of the stoma may be normal during these first weeks however.

It is recommended that the patient uses loose clothing so as not to press the stoma. If the stoma is not red, the patient can shower within a week.

Care during feeding

An adapted nutrition formula should be used, rather than grinding regular foods, as this will contain high amounts of water or oil reach a proper consistency for it to be administered through the tube; it will not have an adequate and balanced supply of nutrients and will be generally deficient in protein and excessive in fat. The formula may be administered by gravity, in a syringe or with a low-pressure feeding pump, either continuously or intermittently. The patient must be positioned at a 30-45° angle to facilitate gastric emptying and prevent reflux. This position must be maintained for an hour after completion of the feeding. The feeding formula should be administered at room temperature, starting at low volumes, increasing progressively as tolerance rises.

After food or drugs administration, it is necessary to instill 50 mL of water to flush any residue from the tube. In absence of a fluid restriction, it is recommended to use a large flushing volume, when possible. In case of continuous nutrition, it should be done every 4-6 hours. A syringe of 30 mL or greater is recommended, in order to avoid too much pressure and consequently the rupture of any component of the PEG tube (82).

In case of PEG tube obstruction, the use of pancreatitis enzymes mixed with a bicarbonate solution has been shown to be an effective method for unclogging the tube; after that, the PEG should be flushed with warm water and carbonated beverages. Finally, an effective method for unclogging the tube in some studies consists of using pancreatic enzymes mixed with bicarbonate solution, prior to flushing with warm water and carbonated beverages (63).

The patency of the tube can be checked by slowly aspirating gastric contents. It has been recommended that if greater than 100 mL, the content should be reintroduced and waiting for an hour before increasing the volume.

Administering medication through the PEG tube

The evidence regarding the effectiveness of nursing interventions in minimizing the complications associated with administering medication via enteral tubes is limited, with a lack of high-quality research on many important issues (83). However, a systematic review allows us to provide some recommendations to be considered when administering medications to a patient carrying a PEG tube:

All kinds of medications will be given diluted in water and unmixed, providing 5-30 mL of water after each and should never be mixed with formula. Enteric coated and sustained released pills should never be crushed; chewable, cytotoxic preparations, or sublingual tablets are not recommended. Bulk-forming tablets, such as Metamucil are prohibited. Hypertonic and concentrated drugs should be diluted in water before administration. Warfarin, phenitoin, morphine sulphate and aluminum-containing antacids should not be given in conjunction with feeding because of delayed drug response (84).

If available, liquid form medications are preferable since they may prevent occlusion of silicone PEG tube and nasoenteral tubes compared to solid forms. Diarrhea has been attributed to the sorbitol content in many liquid medications, rather than the drug itself, so are not advised. Effervescent drug preparations should also be avoided in order to prevent tube occlusion (85).

CONCLUSIONS

Feeding through a PEG tube is the desirable method to feed patients with dysphagia or in those patients who are unable to feed orally but have a functioning digestive system. The technique has become more widespread due to its simplicity, safety and low cost. For proper conduct, specific training for professionals responsible for these procedures is required, and in turn, from them to provide training and information to other professionals and caregivers involved in patient care. Administering proper care tailored and customized to each case, adopting preventive strategies, identifying and treating early complications will maximize safety and effectiveness outcomes for patients.

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