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Several clinical conditions require adjunctive tube feeding in order to maintain a normal nutritional state for the patient. Commonly in clinical practice, a nasogastric tube is used for this purpose, but for longer periods of time, or for indefinite use, this method of delivering enteral feeds is less acceptable than gastrostomy tube feeding [13, 43]. For much of the twentieth century, the Stamm gastrostomy, which requires open surgical laparotomy, was the most commonly accepted insertion technique. This was until 1979 when Ponsky and Gauderer introduced the percutaneous endoscopic gastrostomy (PEG) technique [18]. PEG has the advantage that it is minimally invasive, it can be performed by a gastroenterologist, it is relatively inexpensive, and, if the patient's condition precludes use of a general

anesthetic, it can be performed under sedation. A PEG may also be placed with laparoscopic assistance when anatomical variants preclude the conventional PEG insertion technique [17].

Indications for Gastrostomy Tube Feeding

The range of indications for insertion of a PEG is extensive (see Table 99.1). The commonest indication for PEG insertion in pediatrics is to overcome oral-motor impairment and feeding difficulties in children with neurological impairment; the largest single group is children with cerebral palsy. Contraindications to gastrostomy tube insertion are listed in Table 99.2.

In children with neurological impairment gastrostomy, placement has been shown to significantly increase weight, reduce feeding time, and reduce both feed-related choking episodes and frequency of chest infections [27, 54, 57]. Family stress is significantly reduced [27], and quality of life of parents increases after PEG insertion to assist feeding [58]. Severe oral-motor dysfunction is a marker for the severity of degree of neurological dysfunction. Accordingly, children with severe neurological impairment who require gastrostomy feeding have a substantial long-term mortality. This is probably related more to the underlying neurological condition than it is to PEG placement [9].

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Table 99.1 Indications for insertion of a gastrostomy feeding tube

Failure of adequate nutritional intake
Oral-motor dysfunction (>50% the commonest indication)
Craniofacial abnormalities
Head and neck trauma
Supplemental alimentation in those with increased calorie requirements
Malignancy and chemotherapy
Chronic renal failure
Cystic fibrosis
Congenital heart disease
Crohn's disease
Short bowel syndrome
Human immunodeficiency viral infection
Prolonged dependence on nasogastric tube feeding (>6 weeks)
Unsafe airway
Recurrent aspiration
Gastric drainage/decompression
Motility disorders
Short-bowel syndrome

Table 99.2 Contraindications to insertion of a gastrostomy feeding tube

Sick, unstable patient, e.g., in heart failure
Coagulopathy/bleeding disorders
Peritonitis
Severe ascites
Gastric varices
Distorted anatomy
E.g., 2° to severe kyphoscoliosis
Colonic interposition
Oesophageal obstruction
Hepatosplenomegaly
Failed diaphanoscopy

Complications of Gastrostomy Tube Feeding

Insertion of a PEG feeding tube carries with it a relatively low risk of complications. The result from Larson's series, which includes both adults and children, is typical of the published literature and revealed a procedure-related mortality of

1%, a major complication rate of 3%, and a minor complication rate of 13% [35]. The commonest minor complication is infection of the gastrostomy insertion site and overgrowth of granulation tissue. Major complications are rare and include wound infection, cellulitis, oesophageal injury (probably sustained during extraction of the guide wire), abdominal wall abscesses, necrotizing fasciitis, gastrocolic fistula, colocolic fistula, duodenal hematoma, complicated pneumoperitoneum, gastric perforation, peritonitis, acute gastric dilatation, and gastroduodenal obstruction caused by the balloon of the gastrostomy catheter. Those patients with multisystem organ failure have an increased rate of complications and a poor response to nutritional support; for this population, the risk of PEG may outweigh its benefit [38].

Many of these complications can be avoided or reduced in likelihood by refinements to the technique of insertion [3]. A further complication and one which produces significant issues in relation to clinical management is symptomatic gastroesophageal reflux (GER) occurring after PEG insertion [27, 56].

Gastroesophageal Reflux

Gastroesophageal reflux (GER) is common in children with cerebral palsy, the largest group in whom a gastrostomy feeding tube is inserted, and occurs in 19–75% of such cases [21, 48, 53, 59]. Central nervous system dysfunction is the prime cause of this high incidence of GER in children with cerebral palsy. Additional contributory factors include hiatus hernia, adoption of a prolonged supine position, and increased intra-abdominal pressure secondary to spasticity, scoliosis, or seizures [22, 24]. As a result of neuromuscular incoordination in the foregut, the anti-reflux function of the lower esophageal sphincter mechanism and esophageal motility are significantly impaired. Gastric dysmotility and delayed gastric emptying may also predispose toward GER in children with neurological impairment [2, 4, 8, 42], although

this relationship has not been demonstrated in all studies [6, 30, 41, 52].

Insertion of a Stamm gastrostomy has been shown to reduce lower esophageal sphincter (LES) pressure and predispose to GER [7, 32]. Studies following PEG insertion have shown both an increase in LOS pressure [29], and no effect on basal LES pressure unless rapid bolus feeds are delivered via the tube [10].

Similarly, some authors have found no relationship between PEG insertion and GER [37, 47, 49, 56], whereas others have [20, 27]. The reported postoperative prevalence of GER as a complication of PEG insertion varies from 13% to 28% [26, 28, 33, 56]. It may be that the site of insertion of the gastrostomy tube has an influence on the development of postoperative GER, and some endoscopists have found that tube placement in the antrum or lesser curve is associated with less subsequent reflux [47, 50].

Given the uncertainty about whether PEG insertion will exacerbate GER in the individual patient, especially those with foregut dysmotility, it would seem prudent to establish whether or not GER exists preoperatively [19]. Unfortunately, no test has been shown reliably to predict which patients will develop clinically significant GER post-PEG insertion. Despite normal clinical history and preoperative radiological and lower esophageal pH studies, GER can become apparent in neurologically impaired children after gastrostomy tube placement [5]. Much of the evidence in the literature is conflicting as a result of relatively small studies in selected cases, but the larger studies have shown no significant difference in GER symptoms or median reflux index on 24-h lower esophageal pH monitoring before and after PEG insertion [33, 37, 47]. Even preoperative histological evidence of esophagitis is poorly predictive of subsequent significant GER [12, 26]. In practice a pragmatic attitude should be adopted which takes into account the extent of clinical symptoms of GER (vomiting, aspiration, etc.) prior to PEG insertion and then selects those patients with significant clinical symptoms for investigation by prolonged lower esophageal pH monitor-

ing and barium or water-soluble contrast studies to determine the need for a surgical anti-reflux procedure or jejunostomy [19].

Surgical Anti-reflux Procedures

The notion of a “prophylactic” anti-reflux procedure following gastrostomy insertion especially in children with neurological impairment was advocated by some [31]. The consensus’ view now, however, is that such an approach is not advisable [14, 23, 33, 34, 45, 51, 56, 61, 62]. A major reason for this view is that fundoplication is associated with a higher morbidity and mortality rates in neurologically impaired children, when compared with neurologically normal children [36, 40, 44, 46].

Postoperative morbidity rates of up to 50% and reoperation rates of up to 20% and mortality rates up to 50% are quoted following standard Nissen fundoplication [1, 40]. Major complications can occur both intra- and postoperatively including hepatic vein laceration, bowel perforation, tension pneumothorax, paraesophageal hernia, and small bowel obstruction [44]. Children with neurological impairment have more than twice the complication rate, three times the morbidity rate, and four times the anti-reflux reoperation rate than non-neurologically impaired children [15, 44]. In one study, for instance, more than 30% of children with neurological impairment had major complications or died within 30 days of surgery, and 25% had documented operative failure [39]. In another report, nearly one half of neurologically impaired children had documented recurrent GER after surgery [40]. Recurrent GER often leads to a second operation, but these repeats have a failure rate of around 30% [11, 55, 60].

No single symptom is reliably predictive of recurrent GER so it is necessary to have a high index of suspicion for the development of recurrent GER after anti-reflux procedure in neurologically impaired children and to have a low threshold for proceeding to upper GI contrast study and lower esophageal pH study or endoscopy to investigate this possibility [39].

Medical Management of Gastroesophageal Reflux

The advent of proton-pump inhibitors (PPI) for use in children has had a very significant impact on the treatment of GER. Just as increasing experience of the complications following fundoplication has been shown to raise the threshold for performing this operation in children with neurological impairment [51] so has the efficacy of PPI as medical treatment been associated with a dramatic decrease in the number of surgical anti-reflux procedures performed in children [25].

In conjunction with PPI therapy, strategies to control reflux include a change from bolus to continuous pump feeding [10] and use of whey-predominant enteral milk formulae which have been shown to be associated with faster gastric emptying and less reflux [16].

References

- Albanese CT, Towbin RB, Ulman I, Lewis J, Smith SD. Percutaneous gastrojejunostomy versus Nissen fundoplication for enteral feeding of the neurologically impaired child with gastroesophageal reflux. *J Pediatr*. 1993;123:371–5.
- Alexander F, Wyllie R, Jirousek K, Secic M, Porvasnik S. Delayed gastric emptying affects outcome of Nissen fundoplication in neurologically impaired children. *Surgery*. 1997;122:690–8.
- Beasley SW, Catto-Smith AG, Davidson PM. How to avoid complications during percutaneous endoscopic gastrostomy. *J Pediatr Surg*. 1995;30:671–3.
- Brown RA, Wynchank S, Rode H, Millar AJ, Mann MD. Is a gastric drainage procedure necessary at the time of antireflux surgery? *J Pediatr Gastroenterol Nutr*. 1997;25:377–80.
- Burd RS, Price MR, Whalen TV. The role of protective antireflux procedures in neurologically impaired children: a decision analysis. *J Pediatr Surg*. 2002;37:500–6.
- Campbell JR, Gilchrist BF, Harrison MW. Pyloroplasty in association with Nissen fundoplication in children with neurologic disorders. *J Pediatr Surg*. 1989;24:375–7.
- Canal DF, Vane DW, Goto S, Gardner GP, Grosfeld JL. Reduction of lower esophageal sphincter pressure with Stamm gastrostomy. *J Pediatr Surg*. 1987;22:54–7.
- Carroccio A, Iacono G, Li Voti G, Montalto G, Cavataio F, Tulone V, Lorello D, Kazmierska I, Acerno C, Notarbartolo A. Gastric emptying in infants with gastroesophageal reflux. Ultrasound evaluation before and after cisapride administration. *Scand J Gastroenterol*. 1992;27:799–804.
- Catto-Smith AG, Jimenez S. Morbidity and mortality after percutaneous endoscopic gastrostomy in children with neurological disability. *J Gastroenterol Hepatol*. 2006;21:734–8.
- Coben RM, Weintraub A, Di Marino Jr AJ, Cohen S. Gastroesophageal reflux during gastrostomy feeding. *Gastroenterology*. 1994;106:13–8.
- Dalla Vecchia LK, Grosfeld JL, West KW, Rescorla FJ, Scherer LR, Engum SA. Reoperation after Nissen fundoplication in children with gastroesophageal reflux: experience with 130 patients. *Ann Surg*. 1997;226:315–3.
- Douzinas EE, Andrianakis I, Livaditi O, Bakos D, Flevari K, Goutas N, Vlachodimitropoulos D, Tasoulis MK, Betrosian AP. Reasons of PEG failure to eliminate gastroesophageal reflux in mechanically ventilated patients. *World J Gastroenterol*. 2009;15:5455–60.
- Fay DE, Poplasky M, Gruber M, Lance P. Long-term enteral feeding: a retrospective comparison of delivery via percutaneous endoscopic gastrostomy and nasoenteric tubes. *Am J Gastroenterol*. 1991;86:1604–9.
- Flake AW, Shopene C, Ziegler MM. Anti-reflux gastrointestinal surgery in the neurologically handicapped child. *Pediatr Surg Int*. 1991;6:92–4.
- Fonkalsrud EW, Ashcraft KW, Coran AG, Ellis DG, Grosfeld JL, Tunell WP, Weber TR. Surgical treatment of gastroesophageal reflux in children: a combined hospital study of 7467 patients. *Pediatrics*. 1998;101:419–22.
- Fried MD, Khoshoo V, Secker DJ, Gilday DL, Ash JM, Pencharz PB. Decrease in gastric emptying time and episodes of regurgitation in children with spastic quadriplegia fed a whey-based formula. *J Pediatr*. 1992;120:569–72.
- Frohlich T, Richter M, Carbon R, Barth B, Kohler H. Review article: percutaneous endoscopic gastrostomy in infants and children. *Aliment Pharmacol Ther*. 2010;31(8):788–801.
- Gauderer MW, Ponsky JL, Izant Jr RJ. Gastrostomy without laparotomy: a percutaneous endoscopic technique. *J Pediatr Surg*. 1980;15:872–5.
- Gottrand F, Michaud L. Percutaneous endoscopic gastrostomy and gastro-esophageal reflux: are we correctly addressing the question? *J Pediatr Gastroenterol Nutr*. 2002;35:22–4.
- Grunow JE, Al-Hafidh A, Tunell WP. Gastroesophageal reflux following percutaneous endoscopic gastrostomy in children. *J Pediatr Surg*. 1989;24:42–4; Discuss.
- Gustafsson PM, Tibbling L. Gastro-oesophageal reflux and oesophageal dysfunction in children and adolescents with brain damage. *Acta Paediatr*. 1994;83:1081–5.

22. Halpern LM, Jolley SG, Johnson DG. Gastroesophageal reflux: a significant association with central nervous system disease in children. *J Pediatr Surg.* 1991;26:171-3.
23. Hament JM, Bax NM, van der Zee DC, De Schryver JE, Nesselaaar C. Complications of percutaneous endoscopic gastrostomy with or without concomitant antireflux surgery in 96 children. *J Pediatr Surg.* 2001;36:1412-5.
24. Harrington JW, Brand DA, Edwards KS. Seizure disorder as a risk factor for gastroesophageal reflux in children with neurodevelopmental disabilities. *Clin Pediatr (Phila).* 2004;43:557-62.
25. Hassall E. Decisions in diagnosing and managing chronic gastroesophageal reflux disease in children. *J Pediatr.* 2005;146:S3-12.
26. Heikenen JB, Werlin SL. Esophageal biopsy does not predict clinical outcome after percutaneous endoscopic gastrostomy in children. *Dysphagia.* 2000;15:167-9.
27. Heine RG, Reddihough DS, Catto-Smith AG. Gastroesophageal reflux and feeding problems after gastrostomy in children with severe neurological impairment. *Dev Med Child Neurol.* 1995;37:320-9.
28. Isch JA, Rescorla FJ, Scherer LR, West KW, Grosfeld JL. The development of gastroesophageal reflux after percutaneous endoscopic gastrostomy. *J Pediatr Surg.* 1997;32:321-3.
29. Johnson DA, Hacker III JF, Benjamin SB, Ciarleglio CA, Chobanian SJ, Van Ness MM, Cattau Jr EL. Percutaneous endoscopic gastrostomy effects on gastroesophageal reflux and the lower esophageal sphincter. *Am J Gastroenterol.* 1987;82:622-4.
30. Jolley SG, Leonard JC, Tunell WP. Gastric emptying in children with gastroesophageal reflux. I. An estimate of effective gastric emptying. *J Pediatr Surg.* 1987;22:923-6.
31. Jolley SG, Smith EI, Tunell WP. Protective antireflux operation with feeding gastrostomy. Experience with children. *Ann Surg.* 1985;201:736-40.
32. Jolley SG, Tunell WP, Hoelzer DJ, Thomas S, Smith EI. Lower esophageal pressure changes with tube gastrostomy: a causative factor of gastroesophageal reflux in children? *J Pediatr Surg.* 1986;21:624-7.
33. Khattak IU, Kimber C, Kiely EM, Spitz L. Percutaneous endoscopic gastrostomy in paediatric practice: complications and outcome. *J Pediatr Surg.* 1998;33:67-72.
34. Langer JC, Wesson DE, Ein SH, Filler RM, Shandling B, Superina RA, Papa M. Feeding gastrostomy in neurologically impaired children: is an antireflux procedure necessary? *J Pediatr Gastroenterol Nutr.* 1988;7:837-41.
35. Larson DE, Burton DD, Schroeder KW, DiMagno EP. Percutaneous endoscopic gastrostomy. Indications, success, complications, and mortality in 314 consecutive patients. *Gastroenterology.* 1987;93:48-52.
36. Lasser MS, Liao JG, Burd RS. National trends in the use of antireflux procedures for children. *Pediatrics.* 2006;118:1828-35.
37. Launay V, Gottrand F, Turck D, Michaud L, Atego S, Farriaux JP. Percutaneous endoscopic gastrostomy in children: influence on gastroesophageal reflux. *Pediatrics.* 1996;97:726-8.
38. Marin OE, Glassman MS, Schoen BT, Caplan DB. Safety and efficacy of percutaneous endoscopic gastrostomy in children. *Am J Gastroenterol.* 1994;89:357-61.
39. Martinez DA, Ginn-Pease ME, Caniano DA. Recognition of recurrent gastroesophageal reflux following antireflux surgery in the neurologically disabled child: high index of suspicion and definitive evaluation. *J Pediatr Surg.* 1992;27:983-8; discuss.
40. Martinez DA, Ginn-Pease ME, Caniano DA. Sequelae of antireflux surgery in profoundly disabled children. *J Pediatr Surg.* 1992;27:267-71; discuss.
41. Mollitt DL, Golladay ES, Seibert JJ. Symptomatic gastroesophageal reflux following gastrostomy in neurologically impaired patients. *Pediatrics.* 1985;75:1124-6.
42. Okada T, Sasaki F, Asaka M, Kato M, Nakagawa M, Todo S. Delay of gastric emptying measured by ¹³C-acetate breath test in neurologically impaired children with gastroesophageal reflux. *Eur J Pediatr Surg.* 2005;15:77-81.
43. Park RH, Allison MC, Lang J, Spence E, Morris AJ, Danesh BJ, Russell RI, Mills PR. Randomised comparison of percutaneous endoscopic gastrostomy and nasogastric tube feeding in patients with persisting neurological dysphagia. *Br Med J.* 1992;304:1406-9.
44. Pearl RH, Robie DK, Ein SH, Shandling B, Wesson DE, Superina R, Mctaggart K, Garcia VF, O'Connor JA, Filler RM. Complications of gastroesophageal antireflux surgery in neurologically impaired versus neurologically normal children. *J Pediatr Surg.* 1990;25:1169-73.
45. Puntis JW, Thwaites R, Abel G, Stringer MD. Children with neurological disorders do not always need fundoplication concomitant with percutaneous endoscopic gastrostomy. *Dev Med Child Neurol.* 2000;42:97-9.
46. Ravelli A, Richards CA, Spitz L, Milla PJ. Is Nissen fundoplication the optimal treatment for gastroesophageal reflux in children with neurological impairment? *J Pediatr Gastroenterol Nutr.* 1996;22:411.
47. Razeghi S, Lang T, Behrens R. Influence of percutaneous endoscopic gastrostomy on gastroesophageal reflux: a prospective study in 68 children. *J Pediatr Gastroenterol Nutr.* 2002;35:27-30.
48. Reyes AL, Cash AJ, Green SH, Booth IW. Gastroesophageal reflux in children with cerebral palsy. *Child Care Health Dev.* 1993;19:109-18.
49. Samuel M, Holmes K. Quantitative and qualitative analysis of gastroesophageal reflux after percutaneous endoscopic gastrostomy. *J Pediatr Surg.* 2002;37:256-61.

50. Seekri IK, Rescorla FJ, Canal DF, Zollinger TW, Saywell Jr R, Grosfeld JL. Lesser curvature gastrostomy reduces the incidence of postoperative gastroesophageal reflux. *J Pediatr Surg.* 1991;26:982-4; discuss.
51. Smith CD, Othersen Jr HB, Gogan NJ, Walker JD. Nissen fundoplication in children with profound neurologic disability. High risks and unmet goals. *Ann Surg.* 1992;215:654-8; discuss.
52. Spiroglou K, Xinias I, Karatzas N, Karatza E, Arsos G, Panteliadis C. Gastric emptying in children with cerebral palsy and gastroesophageal reflux. *Pediatr Neurol.* 2004;31:177-82.
53. Srivastava R, Berry JG, Hall M, Downey EC, O'Gorman M, Dean JM, Barnhart DC. Reflux related hospital admissions after fundoplication in children with neurological impairment: retrospective cohort study. *BMJ.* 2009;339:b4411.
54. Stringel G, Delgado M, Guertin L, Cook JD, Maravilla A, Worthen H. Gastrostomy and Nissen fundoplication in neurologically impaired children. *J Pediatr Surg.* 1989;24:1044-8.
55. Subramaniam R, Dickson AP. Long-term outcome of Boix-Ochoa and Nissen fundoplication in normal and neurologically impaired children. *J Pediatr Surg.* 2000;35:1214-6.
56. Sulaeman E, Udall Jr JN, Brown RF, Mannick EE, Loe WA, Hill CB, Schmidt-Sommerfeld E. Gastroesophageal reflux and Nissen fundoplication following percutaneous endoscopic gastrostomy in children. *J Pediatr Gastroenterol Nutr.* 1998;26:269-73.
57. Sullivan PB, Juszczak E, Bachlet AM, Lambert B, Vernon-Roberts A, Grant HW, Eltumi M, McLean L, Alder N, Thomas AG. Gastrostomy tube feeding in children with cerebral palsy: a prospective, longitudinal study. *Dev Med Child Neurol.* 2005;47:77-85.
58. Sullivan PB, Juszczak E, Bachlet AM, Thomas AG, Lambert B, Vernon-Roberts A, Grant HW, Eltumi M, Alder N, Jenkinson C. Impact of gastrostomy tube feeding on the quality of life of carers of children with cerebral palsy. *Dev Med Child Neurol.* 2004;46:796-800.
59. Wesley JR, Coran AG, Sarahan TM, Klein MD, White SJ. The need for evaluation of gastroesophageal reflux in brain-damaged children referred for feeding gastrostomy. *J Pediatr Surg.* 1981;16:866-71.
60. Wheatley MJ, Coran AG, Wesley JR, Oldham KT, Turnage RH. Redo fundoplication in infants and children with recurrent gastroesophageal reflux. *J Pediatr Surg.* 1991;26:758-61.
61. Wheatley MJ, Wesley JR, Tkach DM, Coran AG. Long-term follow-up of brain-damaged children requiring feeding gastrostomy: should an antireflux procedure always be performed? *J Pediatr Surg.* 1991;26:301-4; discuss.
62. Wilson GJ, Van Der Zee, Bax NM. Endoscopic gastrostomy placement in the child with gastroesophageal reflux: is concomitant antireflux surgery indicated? *J Pediatr Surg.* 2006;41:1441-5.