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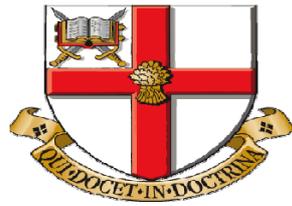
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Malnutrition, enteral nutrition and the use of the percutaneous endoscopic
gastrostomy.

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Abbreviations

ANS	Artificial nutrition support
BMI	Body Mass Index
BANS	British artificial nutrition survey
BAPEN	British association of parenteral and enteral nutrition
EN	Enteral nutrition
ETF	Enteral tube feeding
HETF	Home enteral tube feeding
GP	General Practitioner
LOS	Length of stay
'MUST'	Malnutrition Universal Screening Tool
MAC	Mid arm circumference
MND	Motor neuron Disease
MDT	Multidisciplinary team
MS	Multiple sclerosis
NHS	National Health Service

NICE	National Institute for Health and Clinical Excellence
NBM	Nil by mouth
NG	Nasogastric
PN	Parenteral nutrition
PEG	Percutaneous endoscopic gastrostomy
PIG/PIGG/PRG	Per-oral image-guided gastrostomy
RCTs	Randomized control trials
RIG	Radiologically inserted gastrostomy
RTT	Referral to treatment
TST	Tricep skinfold thickness
UK	United Kingdom

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Introduction

Context of the Exploratory Study

This review will critically appraise the literature on issues surrounding percutaneous endoscopic gastrostomy (PEG) placement for patients who need enteral nutrition (EN) support. The following sections outline the nature of related studies which have been conducted in the past, helping in the establishment of a framework in which the exploratory study can be located and thus providing rationale for the study. There are a number of distinct subject areas which are as follows:

- Malnutrition; definition, consequences and identification;
- Treatment of malnutrition; EN support; orogastric and nasogastric (NG) tube feeding;
- NG versus PEG;
- Care pathways and the multidisciplinary (MDT) team in relation to PEG;
- Trends in home enteral tube feeding (HETF).

Malnutrition; definition, consequences and identification.

There is no universally accepted term to define malnutrition and as a result the term is used interchangeably in literature (Azam, 2007). The term malnutrition for the purpose of this review will be referred to as under-nutrition; “a deficiency of energy, protein and other nutrients that causes adverse effects on the body (shape, size and composition), on the way it functions and on clinical outcomes” (Elia, 2003).

Malnutrition has a number of clinical consequences which include:

- Impaired immune response;
- Reduced muscle strength;
- Impaired wound healing;
- Impaired recovery from illness and surgery;
- Impaired psycho-social function;
- Poorer clinical outcomes.

(Brotherton et al. 2012).

Malnutrition is cited as a major clinical and public health problem and is a burden to patients, health and social care services and society in general (Schenker, 2003). The healthcare cost of managing individuals with malnutrition is cited to be twice that of managing non-malnourished individuals due to higher use of healthcare resources. There is evidence to suggest that malnourished patients need to visit their GP more often, need more prescriptions, have more admissions to hospital and have a longer hospital stay (National Institute for Health and Clinical Excellence, 2006a). The most recent published figures (2007 data) for public expenditure on disease related malnutrition in the United Kingdom (UK) have estimated a cost in excess of £13 billion per annum. This figure is based on malnutrition prevalence figures and the associated costs of both health and social care using the British Association of Parenteral and Enteral Nutrition's (BAPEN's) Malnutrition Universal Screening Tool ('MUST') as the main basis of the calculations. This cost has increased from an estimate of £7.3 billion in 2003, of which most of the healthcare cost was due to the treatment of malnourished patients in hospital (~£3.8 billion)(Elia, Stratton, Russell, Green & Pan, 2006). The National Institute for Health and Clinical Excellence [NICE] (2006a) has identified a substantial cost saving of £28,472 (estimated saving per 100,000 population) which can result from identifying and treating malnutrition. In the UK malnutrition is therefore ranked third in the top clinical guidelines shown to produce cost savings after hypertension and long-acting reversible contraception.

Malnutrition can be identified by screening patients using a screening tool and this is therefore the first step in identifying those at risk of malnutrition (Nutricia, 2009). In 2003 'MUST' was launched - a five step, validated screening tool that can be used across healthcare settings to identify adults who are malnourished or at risk of malnutrition (Brotherton et al. 2012). The 'MUST' screening tool measures height, weight, body mass index (BMI) and any unplanned/unintentional weight loss. An unintentional weight loss of 10% of body weight is usually associated with poorer clinical outcomes (McKinlay, 2008). The presence of an acute disease effect resulting in no dietary intake for more than five days is also considered and an overall risk score for malnutrition obtained. Tricep skinfold thickness (TST) and mid upper arm circumference (MAC) are not included. The 'MUST' can be viewed as tracing the clinical journey of a patient from the past history (history of unintentional weight change) to the present (current weight status and BMI) and into the future (likely effect of underlying condition) (Elia, 2004). Subtle nutritional deficiency states are therefore not identified with 'MUST' but it will potentially identify patients at risk of malnutrition (McKinlay, 2008). A copy of the 'MUST' screening tool is shown in Figure 1 (BAPEN, 2013a, p.3).

Figure 1: A copy of the 'MUST' screening tool.

BAPEN (2013a, p.3).

The 'MUST' screening tool has been identified to be the most commonly used nutritional screening tool in all care settings (85% hospitals and 92% care homes) within the UK (BAPEN, 2012). The reliability of 'MUST' has been established by the inter-rater agreement by different healthcare workers on the same group of patients (kappa values between 0.8 and 1.0), where a kappa value of 1 is where the raters are in complete agreement and a kappa value of ≤ 0 is where there is no agreement among the raters, other than what would be expected by chance (Elia, 2003; Elia, 2004).

The charitable association – BAPEN, raises awareness of malnutrition. For over twenty years the organisation has been integral for a number of changes in the way in which nutrition is regarded within the healthcare system (BAPEN, 2013b). Six key pieces of work in the field of malnutrition have been identified; Kings Fund Report (1992), research by McWhirter and Pennington (1994) and four nutrition screening surveys by BAPEN in 2007, 2008, 2010 and 2011. The work by The Kings Fund (1992) was revolutionary. It set out in straightforward terms why nutrition was an issue within the National Health Service (NHS) by ensuring that patients were well nourished. The Kings Fund Report (1992) was based on a number of references originating as far back as the 1950's. Malnutrition was highlighted as a risk for people with severe illness, resulting in delayed recovery from medical and surgical disorders and increased length of stay (LOS) – duration of a single episode of hospital admission. The importance of

recognising and treating malnutrition before hospital admission was noted as the ideal by nutritional assessment; monitoring of height and weight in general practice and further monitoring within the hospital environment. Therefore the recognition of malnutrition originating from the community was established; with the emphasis for detecting malnutrition being predominantly focused within the hospital setting by improving the education and awareness of malnutrition by doctors and nurses and a hospital nutrition team offering services to the community health team in the area of artificial nutrition support (ANS). The establishment of a care plan was highlighted as an important step after the detection of malnutrition where a plan of treatment and ongoing monitoring was acknowledged. The Kings Fund Report (1992) has been a basis for ongoing research in the field of malnutrition in the UK.

The paper by Mc Whiter and Pennington (1994) is one of the first seminal pieces of published work in the UK to determine the incidence of malnutrition of patients on admission to hospital from the community. One hundred elderly patients were recruited into the study from consecutive admissions into hospital. Each patient's nutritional status was determined from anthropometric data and weight loss before illness; height and weight were recorded and used to determine BMI, MAC and TST. Values were compared to those in tables of normal values for MAC and TST measurements (standardised for sex and age, 16-64 years drawn from data published in the United States and data drawn directly from the elderly population in the UK). It

was noted that 43% of elderly patients admitted into hospital were malnourished of which 4% suffer from mild malnutrition (BMI <20 kg/m², TST or MAC below 15th centile), 20% moderate malnutrition (BMI < 18kg/m², TST or MAC below 5th centile) and 19% severe malnutrition (BMI <16 kg/m², TST or MAC below 5th centile). No statistical details were included in the study by McWhirter and Pennington (1994) to determine the significance of the results. It was acknowledged that BMI by itself is not a sensitive indicator of protein energy malnutrition as it does not distinguish between the depletion of fat or muscle which can introduce error; a low BMI will include individuals who normally weigh less than usual for their height. As a result of the potential misinterpretation when using BMI, other measurements of fat and muscle mass were used; TST to estimate fat reserves and MAC as a measure of muscle protein stores and unintentional weight loss before illness as a measure of nutritional status to identify those at risk of complications as a result of getting thinner. The results provide a “snap shot” picture of the percentage of elderly admitted to hospital with malnutrition. Implications highlighted for clinical practice as a result of the study included the need to improve the recognition of malnutrition in hospital by doctors, nurses and other health care staff (to identify malnourished elderly patients from the community) and to heighten the awareness of nutrition within the community, as there are potentially a number of individuals within the community who are in a suboptimal nutritional state – high risk category of malnutrition.

Between 2007 and 2011, four nutrition screening week surveys by BAPEN aimed to establish the prevalence of malnutrition on admission or recent admission to different care settings, to document screening practice and provide feedback to improve practice. The results based on April 2011 data and the 'MUST' criteria, established that malnutrition affected one in four adults on admission to hospital, more than one in three adults admitted to care homes in the previous six months and up to one in five adults on admission to mental health units in the UK. Most of those affected by malnutrition were found to be in the high risk category ('MUST' Score 2 or more). These results were found to be similar to those obtained in the other three screening surveys with the exception of a lower prevalence of malnutrition on admission to hospital, and the prevalence of malnutrition found on recent admission to care homes similar to the 2008 survey which was higher than in both 2010 and 2007. It was highlighted that nutritional screening policies and practices vary between, and within, healthcare settings and malnutrition continues to be under-recognised and under-treated with much of the malnutrition present on admission to institutions cited to originate in the community (Russell & Elia, 2012). Participation within the screening surveys was voluntary and therefore potential problems with selection bias could result. Higher values of malnutrition could potentially be expected for settings with a screening policy in place; as a result of using a screening tool malnutrition is being detected. In settings where no screening tool is in place malnutrition is not being detected and therefore not treated. This assumption supports the statement that malnutrition is often under-recognised and under-treated, an implication for practice.

No detail has been included in the nutrition screening surveys about the formal determination of a sample size calculation, however with four surveys in total being completed to date this helps to ensure that the surveys are large enough to help determine clinically important results. The four surveys provide the most up to date data on malnutrition within the UK to date.

Treatment of malnutrition.

In most cases malnutrition is a treatable condition that can be managed using first line dietary advice to optimise food intake. Oral nutritional supplements can be used in addition to first line dietary advice where dietary advice alone is unable to prevent and treat malnutrition. ANS in the form of enteral tube feeding (ETF) or parenteral feeding is indicated when oral intake is insufficient or unsafe in the case of swallowing disturbances, unintentional weight loss (>10% of body weight), partial failure of intestinal function requiring supplementary nutrition support and oncology disorders for example (NICE 2006b; Murphy, 2010; Macmillan Cancer Support, 2014). The route and level of feeding is decided on an individual basis according to the clinical indications, treatment plan and nutritional status of the individual. Feeding via a tube into the stomach is considered unless there is upper gastrointestinal dysfunction. (NICE, 2006b). Previous work by Stratton, Green and Elia (2003) concluded that ETF used within the hospital setting, can increase nutritional intake (98% of trials reviewed, all random control trials (RCTs)), significantly reduce mortality (11% Vs. 22%, with

meta-analysis, suggested odds ratio 0.48, 95% CI 0.30-0.78) and significantly lower complication rates including sepsis, wound and urinary infections and pneumonia (33% vs. 48% with meta-analysis, suggested odds ratio 0.5, 95% CI 0.35-0.70). This study has been noted by the European Society for Clinical Nutrition and Metabolism (2006) as a comprehensive and extensive systematic review and meta-analysis concerning the benefit of ETF in the hospital setting and has been the basis of supporting the use of nutritional support as a therapeutic intervention in clinical practice. The study is cited within the American Society for Parenteral and Enteral Nutrition, Disease-Related Malnutrition and Enteral Nutrition Therapy (2010). It is noted however that out of the seventy four trials reviewed as part of the study (n = 2769) only 46% of the trials were randomized (n = 33) and that poor study designs due to low Jadad scores and small sample sizes were limitations of the data set (80% of the trials had a Jadad score of 2 or less, the higher the score the better the study design, highest score 5). Therefore future larger RCTs to assess the clinical effectiveness of ETF is noted as a requirement for future practice.

The NICE Nutrition Support Guidelines (2006) provide recommendations for clinical practice and were the result of the examination of systematic reviews, meta-analyses and RCTs in ETF. The Guidelines concluded that ETF does increase nutritional intake however the evidence to support outcomes such as a reduced LOS or mortality is not clear. The 2006 Guidelines were reviewed in 2012 and the Nutrition Support in Adults,

Quality Standards, QS24 issued (NICE, 2012). The quality standards are based upon the 2006 Guidelines and define clinical best practice within the area of nutrition support including ETF. An integrated MDT approach to the provision of services is cited by NICE (2012) as fundamental to the delivery of high-quality care to adults who need nutrition support. Five quality statements are included within the table below.

Table 1: NICE quality standard statements for nutrition support

Statement number	Quality statement
Statement 1	A validated screening tool ('MUST' for example) is used to screen people for the risk of malnutrition in care settings.
Statement 2	People who are malnourished or at risk of malnutrition have a management care plan.
Statement 3	People who are screened for the risk of malnutrition have their screening results and nutrition support goals documented and communicated between care settings.
Statement 4	People managing their own ANS and/or their carers are trained to manage their nutrition delivery system and monitor their wellbeing.
Statement 5	People receiving nutrition support are "offered a review of the indicators, route, risks, benefits and goals of nutrition support at planned intervals"

(NICE, 2012, p. 7).

The quality standard for nutrition support in adults (2012) provides the most up to date recommendations for current clinical practice.

Parenteral nutrition (PN) is administered intravenously and is usually used when a patient is unable to have enteral nutrition. This may result from small bowel surgery

or obstruction for example where insertion of a tube for EN would be difficult or contra-indicated (Macmillan, 2014). For the purpose of the review PN has not been considered as participants included within the author's exploratory study had no clinical indication for PN.

Enteral nutrition support.

There are a number of ways of administering EN. EN may be delivered either;

- Directly into the stomach via an orogastric or NG tube, gastrostomy; PEG, Per-oral image-guided gastrostomy (PIG/PIGG/ PRG), or radiologically inserted gastrostomy (RIG);
- Beyond the stomach (postpyloric) via a nasoduodenal/nasojejunal tube, gastrojejunostomy or jejunostomy.

(Thomas & Bishop, 2007).

For the purpose of the review EN beyond the stomach has not been considered.

Participants included within the author's exploratory study had no clinical indication for postpyloric feeding. The postpyloric route is considered where there is upper gastrointestinal dysfunction for example delayed gastric emptying.

Orogastric and NG feeding.

Orogastric and NG feeding are considered for short term feeding; four weeks or less

(Parenteral and Enteral Nutrition Group of the British Dietetic Association, 2011).

Orogastric feeding may be used in adults with basal skull fractures (NG feeding is associated with patient death due to potential intracranial placement) (Spurrier, 2008)

and tends to be used extensively in the neonatal unit setting (infants are reported to be obligatory nose breathers and the presence of a NG tube can increase airway

resistance and should therefore be used in ventilated infants only) (National Patient

Safety Agency, 2005; Rogahn, 1998). The orogastric procedure involves the insertion

of a thin flexible NG tube through the mouth directly into the stomach. NG feeding

involves the insertion of a NG tube through the nostril, down the back of the throat

and oesophagus and directly into the stomach as shown in Figure 2 (Macmillan Cancer

Support 2014).

Figure 2: Diagram to show the position of a NG feeding tube.

(Macmillan Cancer Support, 2014, p. 3).

Harm caused by misplaced NG tubes and NG tubes for orogastric feeding, tend to be documented simultaneously as the type of tube used and insertion route is almost identical. With both NG and orogastric feeding tubes there is a reported risk that the tube can become misplaced into the lungs during insertion, or move out of the stomach at a later stage. Patients may experience problems tolerating their enteral feeding regime due to gastro-oesophageal reflux or delayed gastric emptying which could result in aspiration pneumonia (National Collaborating Centre for Acute Care, 2006). The tube is visible and some patients may find initial insertion of a NG tube uncomfortable (Macmillan Cancer Support, 2014).

Gastrostomy feeding.

Gastrostomy feeding is indicated for patients with a functioning gastrointestinal tract and should be considered for long-term feeding (greater than four weeks) (NICE, 2006b). A gastrostomy feeding tube is a narrow tube that is placed directly through the abdominal wall into the stomach shown in Figure 3. A gastrostomy feeding tube can be inserted a number of ways shown in the Table 2 overleaf. The gastrostomy tube is used to deliver nutrition, fluid and medications directly into the stomach (Abbott Nutrition, 2006; White, 1998) and therefore results in an improved nutritional, hydration or clinical status (Draper, 2011). Feeding via a gastrostomy tube permits the maintenance of tissue metabolism and structure even though a patient cannot eat anything, or enough to regain health (Clinical Resource Efficiency Support Team, 2004).

Figure 3: Diagram to show the position of a PEG tube in the stomach.

(Macmillan Cancer Support, 2014, p.5).

Table 2: Types of gastrostomies, methods of insertion and tube characteristics.

Type of gastrostomy	Method of gastrostomy insertion	Gastrostomy tube characteristics
PEG	Endoscopic placement (use of a thin flexible optic camera)	Held in place with internal retention disc/bumper/dome
RIG	X-rays and other imaging techniques used	Held in place with internal retention balloon (water filled)
PIG/PIGG/PRG)	X-ray and other imaging techniques used	Held in place with internal retention disc/bumper/dome

(Abbott Nutrition, 2006; Great Ormond Street, 2014).

There have been a number of studies conducted and evaluated around the effectiveness of NG, PEG, RIG and PIG/PIGG/PRG feeding. NG feeding is cited as a classic, time-proven technique (Gomes et al. 2011). The NG tube can be inserted at the patient's bedside and is inexpensive (Murphy, 2010). The first published report of a PEG was in 1980 by Gauderer, Ponsky and Izant, and was cited as an innovative technique. The technique has become widely accepted and was widely utilised and favoured over the alternative laparotomy gastrostomy of the time. As a result, the majority of research conducted and evaluated around the effectiveness of NG feeding compared to gastrostomy feeding is based upon the PEG technique. There have now been advancements in radiological imaging techniques permitting the use of the fluoroscopically guided version of the PEG techniques; the RIG and PIG/PIGG/PRG (Dickinson, 2013). For the purpose of the review the author will concentrate on the PEG technique. Placement of RIG and PIG/PIGG/PRG are not technically facilitated within the hospital setting where the study was conducted and therefore not included in the review and subsequent exploratory study.

NG versus PEG.

PEG feeding tubes are increasingly used for EN in adults where patients cannot maintain adequate nutrition with oral intake. Indications for PEG feeding include obstruction of the gastrointestinal tract where the passing of a NG tube would be difficult. In 2006 the National Collaborating Centre for Acute Care reviewed NG versus PEG feeding. Four studies were included and reviewed; Baeten and Hoefnagels (1992), Norton, Homer-Ward, Donnelly, Long and Holmes (1996), Park et al. (1992) and The FOOD Trial Collaboration (2005). A number of methodological problems were noted; there was a greater proportion of sick patients in the PEG group when compared to the NG group in the Baeten and Hoefnagels (1992) study and most of the patients in the NG group in the study by Park et al. (1992) transferred to PEG feeding during the study. The study by Norton et al. (1996) cited a significantly lower mortality in the gastrostomy group (2 deaths, 12%) compared to the NG group (8 deaths, 57%), ($P < 0.05$). NG patients received significantly ($P < 0.001$) less of their feed (78%, 95% CI 63% to 94%) compared with the gastrostomy group (100%). It is noted that the study focused on patients with a diagnosis of acute dysphagic stroke; therefore the results may not necessarily be significant when compared to other diagnoses. Results from three large ($n = 859$), randomized multicentre FOOD collaboration trials conducted between November 1996 and July 2003 with stroke patients highlighted that early ETF was associated with a reduction in the risk of death of 5.8% (95% CI 0.8-12.5, $p=0.09$) and a reduction in death or poor outcome of 1.2% (95% CI -4.2 to 6.6, $p=0.7$). In the NG versus PEG trial ($n = 321$), PEG feeding was associated with an increase in risk of

death by 1.0% (95% CI -10 to 11.9, p=0.9) and an increased risk of death or poor outcome of 7.8% (95% CI 0.0 to 15.5, p=0.05). As a result the data was reported not to support the use of PEG feeding (within 30 days) in the dysphagic stroke patient.

Research by Lockett, Templeton, Byrne and Norcross (2002) and a study by Longcroft-Wheaton et al. (2009) (summarised in Table 3 overleaf) suggests that PEG insertion does no harm, supports the view that deaths observed were due to underlying comorbidities and supports the opinion that nutrition support is associated with a better outcome.

Table 3: Summary of Studies conducted by Lockett et al. (2002) and Longcroft-Wheaton et al. (2009).

	Lockett et al. (2002)	Longcroft-Wheaton et al. (2009)
Study design	<p>n = 166</p> <p>Conducted over 2 years</p> <p>Indications for surgery, complications and antibiotic use determined</p>	<p>n = 44 (women n = 22), mean age 75</p> <p>Retrospective</p> <p>Conducted over 1 year</p> <p>Reviewed mortality</p> <p>Based on previous study by Levine, Sachs, Jin and Meltzer (2007); study included a survival curve which the Longcroft-Wheaton et al. (2009) study was compared</p> <p>Retrospective study by Levine et. al (2007) n = 6382, conducted over 4 years, model initially developed on n = 2739 patients, mean age 78 years, 68% female, internally validated n = 3643 from same sample, mean age 78 years, 68% female</p>
Results	<p>1 death directly related to PEG Insertion (peritonitis), 27 complications (16.3%). Mortality 30 days post PEG placement = 13 (7.8%)</p> <p>30 patients died within 30 days post PEG insertion</p> <p>Wound infections occurred in 9 patients</p> <p>4/153 patients with prophylactic antibiotics developed infections</p>	<p>Noted comorbidities (84% neurological, 23% cardiovascular)</p> <p>Mortality at 1, 3, 6 and 12 months = 16%, 20%, 25% and 28% respectively</p> <p>Risk factors from Levine et al. (2007) obtained from administrative data, including comorbidities. Risk factor assigned a weight based on odds ratios. Study not PEG specific, neurological comorbidity not included, 81% sample African American, therefore query comparable.</p>
Conclusions	<p>PEG safe in establishing enteral access in most patients</p> <p>Antibiotics prevent wound infection</p>	<p>PEG does no harm; study supports opinion that nutrition support is associated with a better outcome</p> <p>Higher deaths initially related to poor pre morbid state</p>
Observations		<p>Further comparison study to include PEG and comorbidities would be beneficial</p> <p>Should alternative feeding tube be used initially such as an NG until prognosis more predictable.</p>

A review conducted by Gomes et al. on behalf of the Cochrane Collaboration (2011) evaluated the effectiveness and safety of PEG as compared to NG for adults with swallowing disturbances. Nine randomised control studies were included. It was highlighted that research has been conducted in this field previously; Langmore, Kasarkis, Manca and Olney (2007), Bath, Kerr and Collins (2009), Dennis, Lewis, Cranswick and Forbes (2005), McClave et al. (2005), Douzinas et al. (2006) and Hamidon et al. (2006). The studies by Langmore et al. (2007) and McClave et al. (2005) were excluded from the review as the study by Langmore et al. (2007) was a meta-analysis of which the author was unable to find any controlled or randomized studies and the study by McClave et al. (2005) contained no interventions of interest for the purpose of the Cochrane Review; the study investigated the use of gastric residual values as a marker for aspiration in critically ill patients, not the benefits of NG versus PEG. There are a number of variations noted between the nine studies; follow-up times, age range and range of diagnoses including neurological disease, stroke and neoplasia of the ear, nose and throat. It is noted that some of the authors in these studies failed to report the methods used in sequence generation during randomisation of participants into the NG versus PEG groups and failed to note allocation concealment; a technique used to prevent bias by the researchers (unconsciously or otherwise) influencing which participants are assigned to a given intervention group. The main results were: intervention failure occurred in 19/156 patients in then PEG group and 63/158 patients in the NGT group (95% CI 0.08 to 0.76, $p = 0.01$) in favour of PEG. No statistically significant difference between comparison

groups in complications (risk ratio 1.00) or mortality (relative risk 0.96) were observed (95% CI 0.91 to 1.11, $p = 0.93$) and (95% CI 0.64 to 1.44, $p = 0.64$) respectively. No specific details on the types of intervention failure and complications were noted in the review. Intervention failure could be interpreted as unsuccessful tube placement or even tube misplacement but this cannot be determined.

Complications in the region of 8-30% are cited following the endoscopic placement of enteral feeding tubes (Löser et al. 2005). Two important patient outcomes in regard to PEG placement are improving mortality and quality of life (Plonk, 2005). Thirty day mortality rates have been cited to vary in previous studies. In the National Confidential Enquiry into Patient Outcome and Death [NCEPOD] (2004), mortality rates of 6% (986 deaths) were noted out of a total of 16,648 PEG procedures obtained from UK hospital episode statistics during 2002/2003. A degree of under reporting was believed due to PEG procedures being undertaken in the outpatient setting which would not have been covered by the dataset. Deaths post hospital discharge was not included, potentially underestimating total mortality figures. NCEPOD (2004) conducted a review of mortality in three data sets collected retrospectively by questionnaire during 2002-2003, including deaths within thirty days of PEG insertion. In relation to PEG, 719 cases were identified for review; 76% had respiratory complications and 26% had cardiovascular disease implicated in the causes of death,

indicating that present co-morbidities may be the cause of death, as opposed to mortality directly associated with PEG insertion.

Although some evidence suggests that PEG placement is preferable, patient selection should be timely and appropriate (Ponsky & Gauderer, 1989) and should only be performed if its benefits clearly outweigh its risks and burdens (Plonk, 2005). The General Medical Council reinforce this viewpoint in the document “Treatment and care towards the end of life: good practice in decision making” (2010). It highlights that the most challenging decision is generally about withdrawing or not starting artificial nutrition and hydration when it has the potential to prolong the patient’s life. Overall there is a lack of information on the assessment of benefits with invasive procedures such as PEG and ethical issues associated with randomised controlled trials. The studies raise the question of what is acceptable mortality associated with PEG and does low mortality merely indicate that patients who are sick have been excluded.

Table 4 overleaf details some of the major and minor complications associated with PEG placement. Major complications can be defined as requiring the need for surgery, non-prophylactic antibiotics, or blood transfusion or procedure related death (Vervloessem et al. 2009).

Table 4: Major and minor complications associated with PEG.

Major Complications	Minor Complications
Intra-peritoneal bleeding	Tube blockages
Peritonitis	Tube misplacement
Bowel perforation	Leakage of gastric contents
Haemorrhage	Stoma site infection
Buried bumper syndrome	
Aspiration pneumonia	

(Joanna Briggs Institute, 2010), (National Patient Safety Agency, 2010).

Care pathways and the MDT in relation to PEG.

Care pathways have been linked to an improvement in outcomes including reduced complications, hospital admissions and cost savings in relation to gastrostomy insertion (Talwar & Hewitt, 2009). It is noted by Campbell et al. (2010) that despite the sound principles which underline integrated care pathways, there have been few evaluations undertaken to ascertain the pathways benefit in changing practice and improving outcomes. A total of three PEG care pathways were identified for consultation; Leicestershire and Rutland MND Supportive and Palliative Care Group (2011), Warrington and Halton Hospitals NHS Foundation Trust (2011) (basis of exploratory study) and Wirral Universal Teaching Hospital NHS Foundation Trust (2012). Table 5 overleaf compares and contrasts the content of the three highlighted pathways:

Table 5: Comparison of three PEG care pathways

Author	Leicestershire and Rutland MND Supportive Palliative Care Group (2011)	Warrington and Halton Hospitals NHS Foundation Trust (2011)	Wirral Universal Teaching Hospital NHS Foundation Trust (2012)
Pathway diagnosis specific	Yes, MND	No	No
Pathway care setting specific	Not stated	Community setting	Not stated
Pathway design	Flowchart	Booklet	Booklet
Content of pathway	<p>Referral process stating MDT and contact details</p> <p>Biochemistry review</p> <p>Respiratory assessment</p> <p>If patient suitable for PEG and willing consent; procedure for booking endoscopy appointment, copying of clinic letters to relevant MDT, organisation of EN training detailed.</p> <p>Discharge to HEN dietetic care, biochemistry review by district nurse post PEG insertion</p> <p>If patient not suitable for PEG; RIG or NG considered.</p>	<p>Pre PEG assessment section considering living arrangements, carer support, whether PEG has been discussed with patient/family/ carer, capacity, lasting power of attorney, requirement of a best interest meeting</p> <p>Detailed pre PEG GP, speech and language therapist, dietitian (uses 'MUST'), other professional section and gastroenterology section, which include assessment around current medical condition, contraindications, medication, biochemistry, whether PEG placement is in the patient's best interest, capacity to make decision re PEG insertion</p>	<p>Pre PEG assessment referrals checklist to MDT</p> <p>Explanation of procedure offered to patient/relatives/carers. Written literature provided on procedure, risks/benefits</p> <p>Consent</p> <p>Biochemistry review</p> <p>Medication review</p> <p>Day of procedure checks; prophylactic antibiotics, analgesia, transport,</p> <p>Post procedure checks; clinical observations, analgesia</p> <p>Day one post PEG checks; clinical, biochemistry, medication, feeding, nausea and vomiting, diarrhoea, wound site,</p> <p>Day two/three/four post PEG checks; as per day one checks</p> <p>Ongoing care; daily, biochemistry, feeding, stoma,</p> <p>Discharge checklist included</p>
Overall comments	<p>A4 format</p> <p>No scope to document directly on to pathway</p> <p>Day case admission for PEG insertion</p>	<p>Large booklet</p> <p>Documentation straight on to pathway by various health care professionals</p> <p>Pathway is intended to be used in conjunction with acute PEG pathway which details pre and post PEG placement procedure observations, prophylactic antibiotics, wound management</p>	<p>Large booklet</p> <p>Nursing observations can be documented on to the pathway directly, variance section included</p> <p>No MDT sections to enable documentation of MDT assessments, for example dietitian and speech and language therapist</p> <p>No community involvement prior to PEG insertion. Potentially more appropriate as an acute pathway</p> <p>Query design of pathway anticipates in patient admission following procedure with day one, two, three and four checks detailed.</p>

It is noted by Westaby, Young and O'Toole (2010) that a defined referral pathway should be completed for all cases considered for PEG. Care pathways are noted in research undertaken by Norris (2005) to enable the seamless delivery of care to patients across the primary and secondary care interface of the NHS. The content of the three care pathways reviewed varied considerably, however content which is reflected across all three of the pathways include the MDT, consent, biochemistry and some mention of discharge. Physicians are noted to inform patients and carers poorly regarding PEG benefits, burdens and alternatives (Plonk, 2005), with patients and carers not always aware of the long term implications of PEG (Rickman, 1998). In a study by Todd, Van Rosendaal, Duregon, and Verhoef (2005) it was observed that individuals were insufficiently informed regarding the PEG and the ramifications of the PEG placement. HETF is noted to place a considerable burden on family and carers (Best & Hitchings, 2010) with PEG feeding noted as time consuming and interfering with daily life (Martin, Blomberg & Lagergren, 2012). Research by McKee (1999) highlights that patients must understand what a procedure involves and how the procedure may or may not affect them in the future in order to provide informed consent. In respect to consent and decision making and PEG placement, a team-orientated approach and more active dialogue with regard to care planning among health care professionals and carers has been highlighted to promote effective communication (Todd, Van Rosendaal, Duregon & Verhoef, 2005). In both the Warrington and Halton Hospitals NHS Foundation Trust (2011) and the Wirral

University Teaching Hospitals NHS Trust (2012) pathways, specific reference to the explanation of the PEG procedure with the patient/family and carers are noted.

Integrated care pathways are defined as care plans that detail the essential steps with a specific clinical problem (Campbell, Hotchkiss, Bradshaw & Porteous, 1998). The care pathways by Warrington and Halton Hospitals NHS Foundation Trust (2011) and the Wirral University Teaching Hospital NHS Foundation Trust (2012) could be considered to meet the integrated care pathway definition; they are presented in a structured care plan format, describe the expected progress and variance of the patient's clinical condition and contribute towards positive outcomes. The pathways act as a resource and allow users to access recommendations on PEG insertion and monitoring. This is comparable to a NICE pathway for example (NICE, 2014), however links to other relevant topics to create a network of information are not included.

The pathways reviewed were noted to be suited to either the acute or community setting. The primary placement of a PEG is undertaken in the acute hospital setting (Best & Hitchings, 2010). Pressure on hospital beds often leads to early discharge of EF patients. In some instances, patients will attend the hospital for the insertion of their tube as a day case and be discharged home on the same day (Thomas & Bishop, 2007) as per the Leicestershire and Rutland MND Supportive Palliative Care Group (2011) pathway. It is stated by Kurien et al. (2012) that complications are reported to "often

occur" following discharge in to the community post-PEG insertion and it is acknowledged that there is limited research in the evaluation of community teams in the management of PEG complications. It has been documented by Madigan (2003) and Lowry and Johnston (2007) that community follow up is often inadequate for patients on HETF with only a small number of patients and carers feeling that they receive sufficient information to enable them to manage their feeding tube prior to leaving hospital. This situation poses potential problems if the patient is in hospital for a shortened LOS and then discharged back into the care of community services. The literature published by Madigan (2003), was recorded to reflect a conducted symposium on EF and therefore reflects the views of participants in the symposium and not necessarily documented research. In the study by Lowry and Johnston (2007) a postal questionnaire was used to assess if patients/carers were trained in the care of PEG tube pre-discharge and to ascertain whether appropriate follow up was in place. Sixty six patients were identified for inclusion in the study and a response rate of 44% (29 patients) was noted. Out of the twenty nine respondents, twenty one (72%) reported that they had been taught how to manage the tube, feeds and feeding pumps prior to discharge and 24 had been reviewed by the dietitian post discharge. Fifteen patients had encountered PEG problems, fourteen of whom knew who to contact in the event of a problem. The main problems encountered were the PEG tube falling out (n =7), PEG site infection (n = 2), migration/loosening of the PEG (n = 2) and tube blockage (n = 1). In order to resolve the majority problem of the PEG falling out and migration/loosening of the PEG, the situation would need to be potentially

addressed in the acute setting to facilitate tube re-insertion/review, not the community setting.

Work by Ditchburn (2005) highlights a number of patient benefits following the implementation of a care pathway in the secondary care setting:

- Improved consistency of advice from pre-PEG assessment to post-PEG community follow up;
- Improved continuity of care;
- Provision of information booklets on PEG;
- Involvement of patients/carers with education reducing tube related problems and complications;
- Quicker referrals.

The study was observational; it documented the development of a care pathway to support patients and maintain standards of PEG feeding.

There would appear to be a lack of PEG pathways dedicated for specific diagnoses. For example in the diagnosis of head and neck cancer it is noted that there is no specific published care pathway to coordinate prophylactic gastrostomy insertion before treatment (Talwar & Hewitt, 2009). The use of pathways for a specific diagnosis can be useful. For example in the MND pathway by the Leicestershire and Rutland MND Supportive Palliative Care Group (2011), a respiratory assessment is highlighted as an

example of a diagnosis specific assessment. An alternatively placed gastrostomy tube would need to be considered if the patient failed the respiratory assessment as a result of sedation used during PEG insertion.

Trends in home enteral tube feeding.

PEG is reported to be one of the most commonly performed gastrointestinal procedures (Potack, 2008) with a low incidence of complications (Draper, 2011). For this reason, home enteral tube feeding (HETF) is an expanding area of nutritional support with an increasing number of patients receiving HETF via a gastrostomy (Kurien et al. 2012). The British Artificial Nutrition Survey (BANS) was established in 1996 and has been reporting the trends in ANS for more than 10 years. In 1998 it was estimated that there were more than 12,000 patients on HETF (BANS, 1998). Data from the 2011 British Artificial Nutrition Survey (BANS) suggests that the number of HETF episodes increased by 44% from 12,190 episodes in 2010 to 17,550 episodes in 2011 (BAPEN, 2012). A reduction in reporting rates was noted due to the Health and Social Care Act (2006) (Health Service, 2007), where a requirement for reporters to obtain consent from patients prior to submitting data to BANS was imposed. In 2010 it was confirmed that BANS reporters were no longer required to obtain consent from patients and therefore a subsequent increase in reporting rates was observed. Data suggests that the gastrostomy is the primary route of feeding (75%), as it has been over the last 10 years (BAPEN, 2011). A number of factors have been quoted to have

contributed to the growth of HETF including the reduction in the number of hospital beds and increase in the elderly population (Best & Hitchings, 2010), with 63% of new registrations in 2010 being aged over 60 years and 41% over 70 years of age (BANS, 2011). Neurological conditions including progressive conditions such as MND, MS, bulbar palsy and Parkinson's disease are reported to be the most common indication for PEG (Draper, 2011) (Department of Health, 2005). BANS data from 2008 highlights that 48% of new patient HETF registrations suffered from neurological conditions. Recent BANS data from 2011 notes a decrease in this overall figure from 58% in 2000 compared to 46% in 2010 of which 12% and 8% accounting for MND and MS primary diagnoses respectively. Head and neck cancer accounts for 77% of new HETF registrations.

Summary

In conclusion this review has been useful in considering how to approach PEG placement and the services that should be provided throughout primary and secondary care and across the multidisciplinary teams.

It has established a vision of what the priorities are in relation to PEG insertion in light of the current research evidence available in this field:

- Early and appropriate patient assessment;
- Care of PEG tubes;
- Awareness of complications.

Staff education and the provision of standardised guidelines, care planning and care pathways should be standard practice in the management of PEG placement. The importance of patients and relatives understanding PEG placement, together with the implications for the long term have been highlighted. Assessment of patient suitability for PEG placement should be made in collaboration with the patient and relatives whenever possible.

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Implementation of a pilot community percutaneous endoscopic gastrostomy
placement care pathway: An exploratory study

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The journal appropriate for study submission is the *Journal of Human Nutrition and Dietetics*, an international peer reviewed journal which publishes papers in applied nutrition and dietetics. The following study would potentially be included in the category of clinical nutrition and the practice of therapeutic dietetics.

Abstract

Background: The number of adult patients in the community receiving enteral feeding via a percutaneous endoscopic gastrostomy (PEG) is increasing. Identified problems in relation to PEG were highlighted by a community multidisciplinary team including delayed referrals and discharges. The study aimed to explore retrospectively outcomes in relation to PEG insertion following the implementation of a pilot community PEG placement care pathway.

Methods: Data were analysed for a sample of participants over 18 years of age in three communities, served by a district general hospital in the North West of England. Group 1; ten participants managed on the community PEG placement care pathway and Group 2; ten participants who were not managed on community PEG placement care pathway with a similar primary diagnosis to Group 1. PEG insertion required to maintain nutritional status, hydration and/or medication administration for greater than fourteen days. Group 1 data for referral to treatment (RTT) waiting time was compared with the National Health Service (NHS) RTT waiting times for gastroenterology. Group 1 data for length of stay (LOS) following PEG insertion was compared to Group 2 data by conducting an Independent t-test to analyse LOS between the two groups. A measure of central tendency obtained for LOS for Group 1 and Group 2 data was used in the calculation to estimate treatment cost. Group 1 data to estimate treatment cost was compared to Group 2 data by conducting an Independent t-test to analyse treatment cost between the two groups. Data

collection was obtained to establish if the hospital's PEG information booklet was provided prior to PEG insertion.

Results: 6/10 participants in Group 1 had a RTT waiting time of 1 to 58 days. Median LOS for Group 1 was 4 days; Median LOS for Group 2 was 10 days. Group 1 had an estimated treatment cost of £1114.15 per patient; Group 2 had an estimated treatment cost of £2314.15 per patient. 7/10 Group 1 participants were provided with the hospital's PEG information booklet at least one week prior to PEG insertion.

Conclusions: A reduction in LOS, a RTT waiting time within 18 weeks and a lower estimated mean treatment cost were noted for Group 1 participants. Expansion of the exploratory study is required so the objectives generated can be challenged further.

Introduction

A gastrostomy feeding tube is a narrow tube that is placed directly through the abdominal wall in to the stomach. The gastrostomy tube is generally kept in place with a soft spongy balloon or plastic disc internally and a fixation device externally.

Percutaneous endoscopic gastrostomy (PEG) is a technique used to put a gastrostomy tube in place using an endoscope (Abbott Nutrition, 2006). The first published report of a PEG was in 1980 by Gauderer, Ponsky and Izant (1980). PEG is reported to be one of the most commonly performed gastrointestinal procedures (Potack, 2008). The gastrostomy tube is used to deliver nutrition, fluid and medications directly into the stomach (Abbott Nutrition, 2006). Gastrostomy tube feeding should be considered for long-term (4 weeks or more) enteral tube feeding (National Institute for Health and Clinical Excellence, 2006). Feeding via a gastrostomy tube mitigates malnutrition and permits the maintenance of tissue metabolism and structure even if the patient cannot eat anything or sufficient to regain health (Clinical Resource Efficiency Support Team, 2004).

The primary placement of a gastrostomy is undertaken in the acute hospital setting (Best & Hitchings, 2010). Pressure on hospital beds often leads to early discharge of enterally fed patients. Home enteral tube feeding (HETF) is an expanding area of nutritional support. In some instances, patients will attend the hospital for the insertion of their tube as a day case, so reducing costs compared with other placement methods and discharged home on the same day (Thomas & Bishop, 2007). To ensure

the safe transfer of care and the appropriate monitoring of patients clear communication needs to be established between health professionals in both primary and secondary care (Best & Hitchings 2010). It has been documented that community follow up is often inadequate for patients on HETF with only a small number of patients and carers feeling that they receive sufficient information to enable them to manage their feeding tube prior to leaving hospital (Best & Hitchings 2010). This situation potentially poses problems if the patient is in hospital for a shortened length of stay (LOS) and then discharged back into the care of community services. HETF is noted to place a considerable burden on family and carers (Best & Hitchings, 2010). Adequate training and support to reduce problems and complications are required, making coordination of patient care by a multidisciplinary team (MDT) paramount.

A community HETF service in the North West (NW) of England problems highlighted the lack of formal community assessment procedure prior to PEG insertion, increased referral to treatment (RTT) waiting time from community referral to PEG insertion, increased LOS post insertion and standardised information for PEG insertion was available but not routinely used. A care pathway was developed (Appendix 1) to address the identified issues. The concept of the care pathway is noted to have originated in the United States. It is a “structured multidisciplinary care plan developed to take in to account of current knowledge and best practice” (Ditchburn, 2005, p.34). It details the essential steps in the care of a patient with a specific clinical problem” encouraging evidence based practice (Ditchburn, 2005, p.34). Therefore the

community PEG placement pathway is anticipated to promote best practice and incorporates a MDT checklist which identifies specific interventions at specified times. The presence of contraindications which could affect aftercare post PEG insertion is noted and issues in relation to the provision of information concerning PEG placement. The benchmark that has been used to initiate the community PEG placement pathway is based upon an acute PEG placement pathway used within the acute trust.

Aim

To explore outcomes in relation to PEG insertion subsequent to the implementation of a pilot community PEG placement care pathway.

Objectives

- 1) To explore whether patients who were managed on the pilot community PEG placement care pathway had a RTT waiting time from community referral to PEG insertion of eighteen weeks or less;
- 2) To explore whether patients who were managed on the pilot community PEG placement care pathway had a reduced LOS following PEG insertion versus patients with a similar primary diagnosis who were not managed on the pilot community PEG placement care pathway;
- 3) To explore whether there is a reduction in estimated treatment cost gained from the integrated pilot community PEG placement care pathway.

- 4) To explore if the hospital's PEG information booklet was provided to the patient at least one week prior to PEG insertion.

Materials and methods

The retrospective exploratory study was completed using data on community patients who had a PEG inserted during October 2010 to November 2011. Purposive sampling of twenty patients was conducted (Marshall, 2006). Data has been collected on a sample of the population which the author is interested in (NHS Institute for Innovation and Improvement, 2005). Ten patients over 18 years of age, requiring artificial nutrition support (ANS) for greater than 14 days who were managed the pilot community PEG placement care pathway were identified (Group 1). Ten patients with a similar primary diagnosis who were not managed on the pilot community PEG care pathway (Group 2) (control) were also identified. PEG insertion for both groups was required to maintain nutritional status, hydration and/or medication administration. The timings of the evaluation coincide with the author's dietetic involvement in a working group which developed and piloted the community PEG placement pathway during this period. From a clinical perspective no further participants were identified for PEG insertion to date after the period of November 2011 and no further work has been completed by the working group since the beginning of 2012.

Inclusion Criteria

- ANS required for greater than 14 days;
- Greater than 18 years of age;
- PEG insertion required to facilitate the maintenance of nutritional status, hydration and/or medication administration.

Exclusion Criteria

- ANS required for less than 14 days;
- Less than 18 years of age;
- Diagnosed eating disorder;
- Alternative feeding tube to PEG.

Exploratory Study Setting

The study has been conducted using patients from three communities served by one district general hospital in the North West of England. The three communities were selected as the author is employed within the dietetic department at the local district general hospital which provides the service. Three towns are located in the area which is split into two boroughs. Borough one consists of two towns and has a population of 119,000. The health of the population in borough one is reported to be worse than the England average. Deprivation is a major issue where 21 out of the 79 'Super Output Areas' fall in 10% of the most deprived areas in England. Unemployment is noted as a challenge where one in three adults in one ward is claiming an out-of-work

benefit (Department of Health, 2012a). Borough two consists of one town and has a population of 199,000. At a glance, the health of the population in borough two is reported to be mixed compared with the England average. (Department of Health, 2012b).

Sample Size

Data from the records relating to a total of twenty patients were purposively sampled (Marshall, 1996). Group 1 consisted of ten patients who were managed on the pilot community PEG placement care pathway and Group 2 consisted of ten patients who were not managed on the pilot community PEG placement care pathway with a similar primary diagnosis to Group 1. An activity analysis conducted by the district general hospital's finance department for the period of October 2010 to November 2011 for endoscopic/open procedure codes enabled the identification of suitable Group 2 patients. The endoscopic/open procedure codes were clarified with the coding department to capture total PEG insertion activity. All data obtained for inclusion in the exploratory study was originally collected as part of routine care by professionals involved in the patients overall care. It was necessary to conduct the exploratory study including such patients as they were considered for PEG insertion during the chosen retrospective study period. Patients were not specifically recruited from vulnerable groups, however it is acknowledged by the author that some of the patients included in the study were classed as vulnerable; elderly, adults with learning disabilities and other vulnerable groups.

A community PEG placement pathway evaluation proforma (Appendix 2) was developed to capture qualitative data from the patient's medical notes.

Statistical Analysis

The exploratory study uses quantitative data collection. The quantitative approaches used in the evaluation involve the collection of nominal data and the collection of numerical data through statistics.

Objective One

For objective one, the data collected is nominal. The NHS RTT waiting time for gastroenterology is "within a maximum of 18 weeks of referral". Therefore the author explored the proportion of patients managed on the pilot community PEG placement care pathway who had a RTT within or beyond 18 weeks. The responses will be summarised and commented upon in this report.

Objective Two

The dependent variable is LOS and the independent variable is the two treatment regimes. An Independent t-test was conducted. All data was analysed using the Statistical Package for Social Sciences (SPSS) Version 21 (IBM software Limited, 2012).

The two data sets were analysed to test for any significant difference in LOS in days following PEG insertion amongst patients in both groups.

Objective Three

The median LOS for Group 1 and Group 2 patients was used in the calculation of estimated cost of treatment. An endoscopy examination profile (Appendix 3) and subsequent PEG insertion cost analysis (Appendix 4) have been generated by the hospital's finance and endoscopy departments to provide a figure of £314.15 per PEG insertion. The analysis includes staffing, consumables, equipment, coding, medical records, cleaning and overhead costs. A figure of £200.00 for cost per day of in-patient stay has been provided by the hospital. This figure was multiplied with the respective LOS data and added to the PEG insertion cost analysis to obtain an estimate of total cost. The total cost data were analysed to test for any significant difference between the two groups.

Objective Four

For objective four the data collected is nominal. Therefore the author explored whether patients who were managed on the pilot community PEG placement care pathway were provided with the hospital's PEG insertion information booklet at least one week prior to PEG insertion.

Results

Table 1 below shows the primary diagnoses of the Group 1 and Group 2 patients; Appendix 5 details the primary diagnoses, abbreviations and medical description of the diagnoses.

Table 1: Primary diagnoses for Group 1 and Group 2 patients.

Primary Diagnosis	Group 1	Group 2
CVA	1	2
CVA with DM	1	1
CVA with Dementia/Parkinson's disease	1	1
CVA with Neurological disease	1	-
Friederich's ataxia with DM	-	1
Huntington's chorea	-	1
MND	3	2
Mitochondrial Cytopathy	1	-
PSP	2	1
Dementia	-	1

Where possible, patients for inclusion in Group 2 were selected with the same diagnoses to patients in Group 1. When this was not possible the author selected patients with a comparable diagnosis, however it is noted that this was not achievable in all circumstances. Indications for PEG insertion for Group 1 and Group 2 patients were mainly neurological (7/10) and (5/10) respectively. Difficulty swallowing resulting in the inadequate maintenance of nutrition was indicated in all patients. PEG was not required solely to maintain hydration and/or medication status.

Data collected from the Group 1 pilot PEG placement care pathways, highlighted that six patients were seen by the gastroenterologist within the required time frame; “within a maximum of 18 weeks (126 days) of referral”. A RTT waiting time from 1 to 58 days is noted in Table 2.

Table 2: RTT for Group 1 participants.

Participant	RTT (days)	Outcome
1a	10	
1b	17	
1c	-	Referred to other trust for radiologically inserted gastrostomy (RIG)
1d	-	Deceased
1e	8	
1f	58	
1g	-	Not suitable for PEG
1h	-	Deceased
1i	41	
1j	1	

Objective Two

A review of the statistics for Group 1 reveals a mean LOS of 4.5 days, a median LOS of 4 days (Range 1-9 days). The Group 2 statistics reveal a mean LOS of 17 days, a median LOS of 10 days (Range 0-81 days). $p = 0.127$.

Objective Three

Estimated calculated treatment costs for Group 1 and Group 2 patients are outlined in

Table 3;

Table 3: Estimated treatment cost of PEG insertion for Group 1 and Group 2 patients.

	Cost of PEG + (cost per day of in-patient stay x median LOS)	Total cost	Range
Group 1	314.15 + (200.00 x 4)	£1114.15	£514.15 to £2114.15
Group 2	314.15 + (200.0 x 10)	£2314.15	£314.15 to £16,514.15

$p = 0.127$

Objective Four

Refer to Appendix 6 for a copy of the PEG information booklet. Data collected from the Group 1 pilot PEG placement care pathways highlighted that seven patients/carers were provided with the PEG information booklet at least one week prior to PEG insertion. No pilot community PEG care pathways were recorded in three patients medical notes; therefore the author was not able to establish whether the patients/carers were provided with the PEG information booklet prior to PEG insertion.

Discussion

Clinical conditions and diagnoses may be relevant in particular to “control” participants as it is noted that some of these participants were admitted to hospital and had a PEG inserted as the result of another condition. For example one participant had a diagnosis of MND and was scheduled to have a routine PEG insertion as an out-patient. Unfortunately the participant sustained a fracture in the interim and was therefore admitted in response to the fracture which could have increased the overall LOS, rather than if the patient had been admitted solely for planned PEG insertion. Therefore potentially the inclusion/exclusion criteria required modification.

The study was conducted over a thirteen month period, October 2010 – November 2011. It is noted by Clegg (1990) that using generalised data from a sample can depend upon when the sampling took place. The exploratory study captured PEG insertion activity over the four seasons of the year; a winter bed crisis can be a feature in NHS hospitals, which can result in variability in bed occupancy (Fullerton & Crawford, 1999). Public health concerns such as a flu outbreak can affect staffing, bed occupancy and priority of use. For example there may be a need for the cancelling of routine procedures during an outbreak to ensure the maintenance of other essential NHS services. The study period was limited from a clinical perspective as no further participants were identified for PEG insertion to date (at that time) after the period of November 2011 and no further work has been completed by the working group since

the beginning of 2012. This could potentially influence RTT waiting time, LOS and the provision of information pre PEG insertion.

In objective one, data collected from the Group 1 participants pilot PEG placement care pathway highlighted that six patients were seen “within a maximum of 18 weeks (126 days) of referral”, therefore patients managed on the pilot community PEG placement care pathway did not have a RTT waiting time from community referral to PEG insertion of 18 weeks or more. At first glance this appears acceptable that patients experienced a RTT Waiting Time from 1 to 58 days. As part of the NHS Constitution (March, 2012) patients have the right to access services within maximum waiting times, or for the NHS to take all reasonable steps to offer a range of alternative providers if this is not possible. Patients do have the option to choose to wait longer or, if it is clinically appropriate, for the patient to wait longer. In order to interpret the RTT waiting time values and use in practice, the measurements should not be interpreted on the basis of numerical values alone but in the context of overall risk. From a clinical perspective an increased RTT waiting time can have a detrimental effect if a participant is unable to maintain their nutritional and hydration status resulting in weight loss and a risk of malnutrition and dehydration. The retrospective extraction of file data does not enable the author to capture the experiences of participants/carers prior to PEG insertion. Data was not obtainable for four of the selected patients. Unfortunately there were no other patient records that could be included in the study. This may have affected analysis for objective two where it was noted there is no

significant difference in LOS between the two groups, despite the range of LOS in Group 1 being between 1-9 days, median LOS 4 days and Group 2; 0-81 days, median 10 days; median LOS reduced for the pilot community PEG care pathway group. Despite there being no significant difference in LOS a demonstrated reduction in LOS would be a desirable outcome if confirmed. LOS is a measure to show the mean time that a patient has stayed in hospital. Reducing LOS can free beds and staff time, enabling more patients to be treated and therefore helping to reduce waiting times resulting in an improved patient experience (NHS Institute for Innovation and Improvement, 2008). In order to interpret the LOS values and use in practice, the measurements cannot be interpreted on the basis of numerical values alone.

The estimated median treatment costs in objective three for Group 1 and Group 2 patients are £1114.15 and £2314.15 respectively. Based on the figures there would appear to be a reduced cost of treatment episode based on the median LOS for Group 1 patients. Despite there being no significant difference in treatment cost, a demonstrated cost reduction would be a desirable outcome to support the implementation of the pilot community PEG placement care pathway.

In objective four, from the data which was available, the hospital's PEG insertion information booklet was provided to seven patients/carers at least one week prior to PEG insertion. At first glance this would appear to be acceptable. Some patients, relatives and carers had commented to the author at review post PEG insertion (prior

to implementation of the pilot community PEG placement care pathway), that information about PEG insertion, PEG/stoma aftercare and feeding options were not discussed prior to PEG insertion. Standardised information on PEG was available but wasn't routinely being used. The literature available is suitable for use in both primary and secondary care. It outlines what patients can expect before and after PEG placement as well as identifying risks associated with PEG placement. In order to interpret this information and use in practice a qualitative analysis to answer subjective questions are required; was the information useful? Was it in the correct format? Do patients/carers feel informed? Are the patients/carers able to understand the information? Such information on the relevance of patient information could be linked to further study outcomes and effectiveness of integrated care pathways.

Conclusions and recommendations

The investigation into the four objectives provided insight for the author. A reduction in LOS, a RTT waiting time within 18 weeks and a lower estimated mean treatment cost were associated with implementation of the care pathway. These are positive outcomes. A number of recommendations have been noted by the author for further development and evaluation of the service.

- **A larger sample size;** possible inclusion of more hospitals. This potentially would enable the author to include more participants with comparable diagnoses and more data for RTT, LOS, cost of treatment episode and whether

the hospital's PEG insertion information booklet was provided prior to PEG insertion.

- **Prospective audit and evaluation of the service;** would enable the identification of a larger number of patients to include within the study. Seasonal and specific public health concerns can also be monitored and accounted for during the study.
- **Incorporation of qualitative data, evaluation from the patient/carer and staff perspective;** for example case studies and interviews would provide an in depth description of the situation, noting patients/carers experiences and expectations. Review of the MDT approach, completion of the pathway, caseload management and the collaboration between primary and secondary care could be reviewed. Case studies and interviews to provide feedback on the PEG information booklet, noting patients/carers opinions on whether the information provided is useful.
- **Study to assess morbidity and mortality post PEG insertion;** such a study would provide additional information to help ascertain if there was any effect from an increased RTT Waiting Time and overall morbidity and mortality.
- **PEG insertion and improvement of patient outcomes;** delivery of consistent nutrition, hydration and medication as the result of PEG insertion.

- **Assessment of complications post PEG insertion;** does frequency of community follow result in a significant reduction of complications post PEG insertion.
- **Review of quality of life post PEG placement;** a study incorporating appropriate markers to assess quality of life post PEG placement at specific time intervals. Benefits and burdens of PEG tube and stoma maintenance, as well as patients/carers opinions on feeding – method of administration and frequency.

Non-probability sampling was required for the exploratory study. The patients were selected on the basis of the accessibility of their data/records and purposive personal judgement of the author. The downside of this method is that an unknown proportion of the entire population was not sampled and therefore may not represent the entire population accurately; however the study conducted was merely exploratory (Non-Probability Sampling, 2013).

The pilot community PEG placement care pathway requires review and auditing to assess and document its effectiveness in clinical practice. Problems identified as a result of review and potential implementation would need to be addressed as they arise.

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Appendix 1: Community PEG placement care pathway

Print copy only.

Appendix 2

Community PEG Placement Pathway Evaluation Proforma

Group Number:

Primary Diagnosis:

Date of referral for PEG Insertion:

Date of outpatient appointment with Gastroenterologist:

Date of admission for PEG insertion:

Date of PEG insertion:

Date of discharge:

Was the Trust's PEG information booklet made available for use by patient/carer prior to PEG insertion?

Yes/No

Was it noted on what date the information was provided?

Yes/No

If yes what was the date.

Appendix 3: Endoscopy Examination Profile.

Question	Answer	Comments
Examination type Therapeutic Gastroscopy – Insertion of PEG	PEG placement	For therapeutic treatment and management of specific conditions
Performed by <ul style="list-style-type: none"> • Consultant/ Registrar • Nurse specialist 	Consultant/ Registrar/Nurse Specialist	For PEG placement assistance needed; Nurse specialist for example
Staffing allocated in support of list x1 band 6 nurse and x1 band 2 carer	2 qualified nurses band 5 and above 2 Band 3 Carers (decontamination of Scopes)	Airway management Vital signs observation, Meditech documentation, specimen collection
Consumables routinely used for these cases drugs, guide wires, etc.	Lignocaine spray 6p Midazolam 60p Disposable mouth guard 49p PEG Tube kit £80 Drawing up needles, syringes, Chloroprep, cannula and venflon dressings, gloves, aprons £2.50	Appropriate consumables used to treat these various conditions
Average examination time +/- number of cases per standard (unmixed list)	This procedures can take 30 minutes and are best performed on an unmixed list due to the urgency and need of individual patients	Examination times are dependent on individual needs

Appendix 4: PEG Insertion Cost Analysis

	Cost
Consultant	£ 66.00
Registrar	£ 18.00
Nursing (Procedure/Recovery)	£ 32.00
Decontamination	£ 5.00
Reception	£ 4.00
Administration	£ 11.00
Lignocaine	£ 0.06
Midazolam	£ 1.00
Disposable Mouth Guard	£ 0.49
PEG Tube Kit	£ 80.00
Drawing up needles, syringes, Chloroprep, cannula, gloves	£ 3.00
Managed Service – Endoscopy equipment	£ 35.00
Domestics	£ 3.00
Overheads	£ 57.00
TOTAL COST	£ 314.55

Appendix 5

Primary Diagnosis Abbreviations and Definitions

- Cerebrovascular Accident (CVA): Also known as stroke, resulting from infarction of part of the brain, or from intracerebral haemorrhage (Hope, Longmore, Hodgetts & Ramrakha, 1993).
- Diabetes Mellitus (DM): A syndrome caused by the lack or diminished effectiveness of endogenous insulin (Hope et al. 1993).
- Dementia: A chronic or persistent disorder of brain process due to organic brain disease. Symptoms include memory disorders, impaired reasoning ability and disorientation (Martin, 1994).
- Friederich's ataxia: An inherited disorder resulting in the clumsiness of willed movements, staggering when walking, unable to pronounce words appropriately, rapid involuntary movements of the eyes and spasticity of the limbs (Martin, 1994).
- Huntington's chorea: A hereditary disease caused by the defect of a single gene inherited as a dominant characteristic. Symptoms include involuntary movements accompanied by behavioural changes and progressive dementia (Martin, 1994).
- Motor Neurone Disease (MND): Progressive degenerative disease of unknown cause, affects upper and lower neurones, causes muscle weakness and wasting, no sensory abnormality present (Martin, 1994).
- Mitochondrial cytoplasty: Genetic disease.
- Neurological disease: Any disorder of the body's nervous system. A range of symptoms can be experienced which depends upon which nerves of the various systems within the body are affected.
- Parkinson's disease: A progressive disease that may lead to dementia. The disease is caused by the degeneration of dopaminergic neurones (Hope et al. 1993)
- Progressive Supranuclear Palsy (PSP): Degenerative condition that affects movement, personality, balance and later symptoms signs of dementia. Slurring of speech, moving of eyes and difficulty swallowing.

Appendix 6

Hospital's PEG information booklet

Print copy only.