A root cause analysis system to establish the most plausible explanation for post-endoscopy upper gastrointestinal cancer

Background Missing upper gastrointestinal cancer (UGIC) at endoscopy may prevent curative treatment. We have developed a root cause analysis system for potentially missed UGICs at endoscopy (post-endoscopy UGIC [PEUGIC]) to establish the most plausible explanations.

Methods The electronic records of patients with UGIC at two National Health Service providers were examined. PEUGICs were defined as UGICs diagnosed 6–36 months after an endoscopy that did not diagnose cancer. An algorithm based on the World Endoscopy Organization post-colonoscopy colorectal cancer algorithm was developed to categorize and identify potentially avoidable PEUGICs.

Results Of 1327 UGICs studied, 89 (6.7%) were PEUGICs (patient median [IQR] age at endoscopy 73.5 [63.5–81.0]; 60.7% men). Of the PEUGICs, 40% were diagnosed in patients with Barrett’s esophagus. PEUGICs were categorized as: A – lesion detected, adequate assessment and decision-making, but PEUGIC occurred (16.9%); B – lesion detected, inadequate assessment or decision-making (34.8%); C – possible missed lesion, endoscopy and decision-making

Authors
Umair Kamran1, Dominic King1, Abdullah Abbasi2, Ben Coupland3, Nosheen Umar1, Warren C. Chapman1, Srisha Hebbar2, Nigel J. Trudgill1

Institutions
1 Department of Gastroenterology, Sandwell and West Birmingham NHS Trust, West Bromwich, UK
2 Department of Gastroenterology, University Hospitals of North Midlands NHS Trust, Stoke-on-Trent, UK
3 Health Informatics, University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK

submitted 15.12.2021
accepted after revision 7.7.2022
published online 31.8.2022

Bibliography
Endoscopy 2023; 55: 109–118
DOI 10.1055/a-1917-0192
ISSN 0013-726X
© 2022, Thieme. All rights reserved.
Georg Thieme Verlag KG, Rüdigerstraße 14, 70469 Stuttgart, Germany

Table 1’s, Fig. 1’s
Supplementary material is available under https://doi.org/10.1055/a-1917-0192
adequate (8.9 %); D – possible missed lesion, endoscopy or decision-making inadequate (33.7 %); E – deviated from management pathway but appropriate (5.6 %); F – deviated appropriately from management pathway (3.4 %). The majority of PEUGICs (71 %) were potentially avoidable and in 45 % the cancer outcome could have been different if it had been diagnosed on the initial endoscopy. There was a negative correlation between endoscopists’ mean annual number of endoscopies and the technically attributable PEUGIC rate (correlation coefficient −0.57; P = 0.004).

Conclusion Missed opportunities to avoid PEUGIC were identified in 71 % of cases. Root cause analysis can standardize future investigation of PEUGIC and guide quality improvement efforts.

Introduction

Upper gastrointestinal cancers (UGICs) are usually diagnosed by endoscopy; however, UGIC can be diagnosed after an endoscopy that did not identify the cancer. These are termed post-endoscopy UGICs (PEUGICs). The British Society of Gastroenterology (BSG) recommends that PEUGIC should be a quality standard and regularly audited [1]. In a meta-analysis, 11.3 % of UGICs were not diagnosed at an endoscopy performed up to 3 years before the diagnosis [2], and more recent studies report PEUGIC rates of 6.7 %–9.4 % [3–6]. PEUGICs are less likely to present with alarm symptoms and are more commonly associated with less advanced clinical stage [7]. Other associated factors include younger age, female sex, increasing deprivation, and an inadequate number of biopsies [5, 6, 8], but not endoscopist experience [9]; however, the studies published to date have lacked a systematic analysis approach to the causes of PEUGIC.

Colorectal cancer (CRC) diagnosed following a colonoscopy that did not diagnose the CRC is termed post-colonoscopy CRC (PCCRC) [10]. The World Endoscopy Organization (WEO) has proposed that PCCRC should be categorized into interval and non-interval cancers and has provided a system to determine the most plausible etiologies [11]. This was subsequently validated, with suggestions made to improve the categorization system [12].

We have undertaken a detailed analysis of PEUGICs to establish how many were interval and non-interval cancers, and have developed a root cause analysis system based on the WEO-PCCRC system to identify the most plausible explanations.

Methods

Patient identification and data collection

Using International Classification of Diseases 10th revision codes, adults (> 18 years) diagnosed with esophageal (C15), gastric (C16), and duodenal (C17) cancers were identified at two UK endoscopy providers: Sandwell and West Birmingham NHS trust (January 2010 to December 2019) and University Hospitals of North Midlands NHS Trust (January 2017 to March 2020). Patients were excluded if they did not have an endoscopy prior to their diagnosis or were referred from other hospitals. Other exclusion criteria included: non-UGI cancers, neuroendocrine tumours, sarcomas, and gastrointestinal stromal tumours.

PEUGICs were defined as cancers in patients who had an endoscopy that did not diagnose their cancer 6–36 months prior to the UGIC diagnosis. Patients who had an endoscopy within 6 months of the UGIC diagnosis were deemed UGIC controls. If PEUGIC patients had more than one endoscopy 6–36 months prior to diagnosis, the endoscopy that did not diagnose cancer that was closest to the date of the cancer diagnosis was classified as the index endoscopy. For controls, the endoscopy closest to the date of the cancer diagnosis was the index endoscopy. All patients with UGIC or dysplasia at the two providers are reported by an expert GI pathologist and confirmed by a second pathologist.

Endoscopies were performed with the patient under conscious sedation (using midazolam) or with xylocaine throat spray alone, depending on the patient’s preference and the clinical judgment of the endoscopist. Data collected included: patient variables (age at endoscopy, sex); endoscopy variables (indication, photodocumentation of J maneuver [gastric retroflexion] and view quality in gastric body and second part of duodenum [D2], tolerance [well or poorly tolerated based on the endoscopy report], and endoscopy findings); endoscopist variables (total endoscopies performed over the study period); cancer information (diagnosis date, site, staging, differentiation, tumor size, treatment received [endoscopic resection, surgical resection, chemotherapy, or best supportive care only] and histological diagnosis); and other management information (surveillance or follow-up plan, reasons for deviation from plan [patient related or administrative]).

The total number of UGI endoscopies performed by each endoscopist was extracted from the endoscopy reporting systems. Endoscopies performed on training lists were considered to have been undertaken by the trainer in terms of endoscopy volume and PEUGIC analysis.

Interval and non-interval cancers and root cause analysis of the most plausible explanation for PEUGIC

Interval and non-interval cancers

Interval PEUGICs were identified before the next planned surveillance endoscopy [11]. Non-interval PEUGICs were identified at (type I) or after (type II) the next planned surveillance endoscopy, or when no further surveillance or follow-up was planned (type III). Examples of the PEUGIC subcategories are provided in Table 1s (see online-only Supplementary material).

Root cause analysis of the most plausible explanation for PEUGIC

PEUGICs were categorized into six types (A to F), involving a four-step process:
Step 1  Focal or cancer-associated lesion or premalignant condition detected in the same segment as the subsequent PEUGIC?

If Yes, proceed to Step 2; if No, proceed to Step 3.

Step 2  Lesion adequately described and photographed, adequate biopsy samples taken, and the surveillance/follow-up plan was appropriate?

If Yes, PEUGIC categorized as "A": lesion detected, adequate assessment and decision-making, but PEUGIC still occurred.

If No, PEUGIC categorized as "B": lesion detected, inadequate assessment or decision-making.

Step 3  Index endoscopy adequate or, if inadequate, recognized by the endoscopist as inadequate and planned follow-up was appropriate?

If Yes, PEUGIC categorized as "C": possible missed lesion, endoscopy and decision-making adequate.

If No, PEUGIC categorized as "D": possible missed lesion, endoscopy or decision-making inadequate.

Step 4  If the management pathway deviated from the recommendations following the index endoscopy, the following categories were identified:

Where due to patient choice or the decision of the responsible clinician that the patient was not fit for further investigations, "E": deviated from management pathway but appropriate.

Where due to administrative delays (i.e., surveillance or follow-up procedures not booked within the recommended timeframe), "F": deviated inappropriately from management pathway.

More than one PEUGIC explanation was allowed for individual patients. Detected lesions in the PEUGIC segment included premalignant (Barrett's esophagus, gastric atrophy, or gastric intestinal metaplasia), and focal or cancer-associated lesions (esophageal ulcer or stricture, Los Angeles grade C or D reflux esophagitis, or gastric ulcer).

Endoscopies performed 6 weeks beyond the planned follow-up date (for focal lesions) and 12 weeks beyond the planned surveillance dates (for premalignant conditions) were categorized as inappropriate and related to administrative factors, in the absence of patient choice or an intercurrent illness that delayed the endoscopy.

An endoscopy was considered adequate if the following criteria were met:
1. high definition video-endoscopy with image capture and biopsies
2. J maneuver performed and photographed
3. D2 intubated and photographed
4. view quality in the stomach photographed and classified as excellent, good, or satisfactory, with no foam, mucus, blood, or food limiting the view
5. tolerance excellent, good, or satisfactory and not limiting the view.

Photodocumentation of the J maneuver, D2, and gastric body were the minimum criteria for adequate photodocumentation.

Avoidability

The previously described approach used to define avoidable PCRC [12] was used to determine whether a PEUGIC was potentially avoidable based on cancer size at diagnosis and the factors identified on root cause analysis. Small PEUGICs were categorized as unavoidable if they were growing by < 5 mm/year, as they would have been unlikely to be detectable during the index endoscopy. PEUGICs were also considered unavoidable if the recommended pathway was not followed because the patient declined investigations or was deemed by the responsible clinician to be too frail to proceed with further investigation. All other PEUGICs were considered potentially avoidable.

Potential impact of delay in diagnosis on PEUGIC clinical outcomes

The outcome for a PEUGIC was unlikely to be different if patients were diagnosed at an early stage despite a negative index endoscopy and later underwent successful endoscopic resection. The outcome was also unlikely to be different for patients who were frail at index endoscopy and were unlikely to be eligible for curative treatment at any stage. Patients diagnosed with their cancer at an advanced stage that precluded curative treatment or endoscopic resection were considered to have potentially had a different outcome had they been diagnosed at index endoscopy.

Attribution

PEUGICs were attributable to individual endoscopists if technical endoscopic or decision-making factors were identified on the root cause analysis [12]:
1. premalignant, focal, or cancer-associated lesion identified but not described according to the recommended criteria (e.g., Prague classification [13, 14]) or lesion site or morphology not recorded in the endoscopy report or photographed
2. premalignant, focal, or cancer-associated lesion identified but not biopsied appropriately or according to recommended guidelines where relevant (e.g., Seattle protocol for Barrett's esophagus [13, 15] and Sydney protocol for gastric atrophy and intestinal metaplasia [16])
3. endoscopist did not recommend an appropriate surveillance or follow-up plan
4. if the index endoscopy was inadequate, the endoscopist did not recognize it as inadequate or did not recommend a repeat procedure.

PEUGICs were not deemed attributable in the following situations:
1. small PEUGIC (growing at < 5 mm/year)
2. the patient declined further investigations or was deemed by the responsible clinician to be too unwell for further investigation.

For each endoscopist, the total and mean annual number of UGI endoscopies performed over the study period were extracted. The "technically attributable" rate per 1000 endoscopies was
calculated for each endoscopist by dividing the technically attributable PEUGIC number by the total number of UGI endoscopies.

**Statistical analysis**
The Mann–Whitney U test and chi-squared test were used for continuous and categorical variables respectively, and two-sided P values <0.05 were considered significant. Spearman’s rank correlation was used to assess correlations between the technically attributable PEUGIC rate per 1000 endoscopies and the mean annual number of endoscopies by endoscopists.

A funnel plot examined variation in technically attributable PEUGIC rates between endoscopists. It was constructed as a scatter plot with superimposed control limits, representing one and two SDs from the mean. Endoscopists outside the control limits had a significantly higher technically attributable PEUGIC rate than the mean. Scatter plots were used to correlate the delay in diagnosis (interval from index endoscopy to PEUGIC diagnosis) and cancer stage (I, II, III, or IV) and histological differentiation (categorized as well, moderately, and poorly differentiated) for all PEUGICs and also separately for PEUGICs with and without Barrett’s oesophagus.

Stata statistical software, release 16, was used for the statistical analysis.

**Ethics**
This work was undertaken as a service improvement project and ethics approval was not sought. It was registered with Trust Audit and Quality Improvement Departments of Sandwell and West Birmingham NHS Trust and University Hospitals of North Midlands NHS Trust.

**Patient and public involvement**
There was no patient or public involvement in this study.

**Results**

**Study subjects**
A total of 1327 UGICs met the inclusion criteria (Fig. 1); 89 (6.7 %) were PEUGICs. Of these, 48 % were diagnosed 6–18 months after the index endoscopy and 52 % were diagnosed 18–36 months after the index endoscopy. The patient demographic details and characteristics of the index endoscopy for patients with PEUGIC and the UGIC controls are shown in Table 1.

**Cancer details**
The majority of PEUGICs were esophageal (83 %), with 17 % being gastric. Data on the clinical staging, treatment received, and stratification based on an index endoscopy finding of Barrett’s oesophagus are presented in Table 2. Among the PEUGICs, 57 % were early stage (i.e. stage I or II), compared with 22 % of the UGIC controls. No correlation was found between the interval from the index endoscopy to PEUGIC diagnosis and tumor size, stage at diagnosis, or histological differentiation (Fig. 1). More than half of PEUGIC patients received treatment with curative intent (53 %), compared with 29 % of the UGIC controls (P = 0.002). Patients with PEUGICs were more likely to undergo endoscopic resection than the UGIC controls (31.3 % vs. 5.1 %; P = 0.002).

**Index endoscopy details for PEUGIC patients**
In 27 % of the PEUGIC patients, the index endoscopy was for Barrett’s surveillance compared with 1.1 % in the UGIC control group. In PEUGIC patients, views were excellent or good in 47.2 % and satisfactory in 25.8 %; four patients (4 %) had poor views due to gastric food residue and in 22.5 % the view quality was not recorded. Procedure tolerance was not recorded in 30 % of patients, but the procedure was reported as well tolerated in 64 % of the PEUGIC endoscopies.

Photodocumentation of gastric retroflexion was found in 38 % of the index endoscopies. Duodenal intubation was reported in 89 % of the PEUGIC patients, but D2 photodocumentation was found in only 32.6 %. No images were recorded in 34.8 % of endoscopies, with only one recording that the endoscopy reporting system failed to capture images. The indications for the index endoscopy and endoscopic diagnoses differed between the PEUGIC and UGIC control groups (Table 1).

**Correlation between the attributable PEUGIC rate and endoscopist data**
Technical endoscopic factors were identified in 52 PEUGIC patients (58.4 %). It was not possible to calculate the mean annual endoscopy volume for one endoscopist, who was consequently excluded from further analyses. The technically attributable PEUGIC rate was calculated for 23 endoscopists. A negative cor-

![Table 1](Image)

| UGI cancers at SWBNHST and UHNM NHST n = 1678 |
| Tertiary care referrals n = 174 |
| Critical clinical or endoscopy data not available n = 27 |
| No endoscopy prior to cancer diagnosis n = 57 |
| Non-UGI cancers or unknown primary n = 46 |
| Nonepithelial cancers: neuroendocrine tumours/gastrointestinal stromal tumours/sarcomas n = 47 |
| Included patients n = 1327 |

![Fig. 1 Flowchart describing reasons for exclusion and selection of study patients.](Image)

UGI, upper gastrointestinal; SWBNHST, Sandwell and West Birmingham NHS trust; UHNM NHST, University Hospitals of North Midlands NHS Trust.
A root cause analysis of the most plausible explanation for PEUGIC

PEUGICs were classified as non-interval in 98% of patients: 48% (n = 42) type I; 5% (n = 4) type II; and 47% (n = 41) type III. Only two PEUGICs were interval cancers, both of which occurred in patients with Barrett’s esophagus.

Preamalignant lesions were identified in 39 patients (43.8%): 36 Barrett’s esophagus, two gastric adenoma, and one gastric atrophy. Focal or cancer-associated lesions were identified in 12 patients (13.5%): four esophageal stricture, three abnormal

Root cause analysis of the most plausible explanation for PEUGIC

PEUGICs were classified as non-interval in 98% of patients: 48% (n = 42) type I; 5% (n = 4) type II; and 47% (n = 41) type III. Only two PEUGICs were interval cancers, both of which occurred in patients with Barrett’s esophagus.

Preamalignant lesions were identified in 39 patients (43.8%): 36 Barrett’s esophagus, two gastric adenoma, and one gastric atrophy. Focal or cancer-associated lesions were identified in 12 patients (13.5%): four esophageal stricture, three abnormal
esophageal area, two esophageal ulcer, one severe esophagitis, one cardia inflammation, and one gastroesophageal junction nodule. No lesion was found in 38 patients (42.7 %) (▶Fig.3 shows PEUGIC examples).

The results of root cause analysis of the PEUGIC patients are shown in ▶Fig. 4 and ▶Table 3. More than one plausible explanation was found in seven patients (7.9 %): six had inadequate biopsies and an inadequate surveillance plan, and one had an inadequate surveillance plan and an administrative delay.

### Table 2  Comparison of clinical staging and treatment received by all patients, with further stratification of the post-endoscopy upper gastrointestinal cancer (PEUGIC) patients by index endoscopy findings of Barrett’s esophagus.

<table>
<thead>
<tr>
<th>Total, n (%)</th>
<th>UGIC control, n (%)</th>
<th>PEUGIC, n (%)</th>
<th>P value</th>
<th>Barrett’s esophagus, n (%)</th>
<th>No Barrett’s esophagus, n (%)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Clinical stage at diagnosis</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Early</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I</td>
<td>148 (11.2)</td>
<td>112 (9.0)</td>
<td>36 (40.5)</td>
<td>&lt;0.001</td>
<td>29 (80.6)</td>
<td>7 (13.2)</td>
</tr>
<tr>
<td>II</td>
<td>179 (13.5)</td>
<td>164 (13.2)</td>
<td>15 (16.9)</td>
<td></td>
<td>3 (8.3)</td>
<td>12 (22.6)</td>
</tr>
<tr>
<td><strong>Advanced</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>401 (30.2)</td>
<td>390 (31.5)</td>
<td>11 (12.4)</td>
<td></td>
<td>1 (2.8)</td>
<td>10 (18.9)</td>
</tr>
<tr>
<td>IV</td>
<td>450 (33.9)</td>
<td>429 (34.7)</td>
<td>21 (23.6)</td>
<td></td>
<td>3 (8.3)</td>
<td>18 (34.0)</td>
</tr>
<tr>
<td>Not available</td>
<td>149 (11.2)</td>
<td>143 (11.6)</td>
<td>6 (6.7)</td>
<td>0.260</td>
<td>0 (0.0)</td>
<td>6 (11.3)</td>
</tr>
<tr>
<td><strong>Treatment</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Curative</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Endoscopic resection</td>
<td>86 (6.5)</td>
<td>60 (4.8)</td>
<td>26 (29.2)</td>
<td>&lt;0.001</td>
<td>20 (55.6)</td>
<td>6 (11.3)</td>
</tr>
<tr>
<td>Surgical resection</td>
<td>225 (17.0)</td>
<td>209 (16.9)</td>
<td>16 (18.0)</td>
<td></td>
<td>9 (25.0)</td>
<td>7 (13.2)</td>
</tr>
<tr>
<td>Chemo-radiotherapy</td>
<td>73 (5.5)</td>
<td>71 (5.7)</td>
<td>2 (2.2)</td>
<td></td>
<td>0 (0.0)</td>
<td>2 (3.8)</td>
</tr>
<tr>
<td>Palliative therapy*</td>
<td>943 (71.1)</td>
<td>898 (72.5)</td>
<td>45 (50.6)</td>
<td></td>
<td>7 (19.4)</td>
<td>38 (71.7)</td>
</tr>
</tbody>
</table>

UGIC, upper gastrointestinal cancer.  
* 657 patients had best supportive care only.
Barrett’s esophagus

Around 40% of PEUGICs were diagnosed in patients with Barrett’s esophagus. The Prague criteria were used in 94% of patients and the Seattle biopsy protocol was followed in 42%. Planned surveillance intervals were incorrect in 25%. Among the PEUGICs in patients with Barrett’s esophagus, 89% were diagnosed at an early stage (stage I or II) and 81% received treatment with curative intent (56% endoscopic resection and 25% surgery).

Of 38 non-Barrett’s esophageal PEUGICs, 18 were squamous cell cancers and 20 were adenocarcinomas, including 17 that involved the gastroesophageal junction. Categories D (22/38) and B (12/38) were identified as the commonest most plausible explanations.

Avoidability

Of PEUGICs, 71% were categorized as potentially avoidable. The unavoidable PEUGICs included 21 small PEUGICs; there were also three related to patient’s choice not to undergo further investigations and two where decisions were taken not to investigate further owing to multiple co-morbidities and patient frailty.

Impact on clinical outcome

The clinical outcome could potentially have been different for 45% of PEUGICs. This included 23 patients with advanced stage (stage III or IV) at diagnosis, 14 patients who underwent esophagectomy, and three patients who were too frail at the time of cancer diagnosis but could potentially have been offered endoscopic resection if their cancer had been detected at index endoscopy.

Discussion

This is the first study to report a detailed root cause analysis of unselected PEUGICs, develop a system of analysis to categorize the causes of PEUGIC, and identify contributing factors and missed opportunities to potentially avoid PEUGIC in 71% of patients. Inadequate assessment of premalignant or focal lesions, inadequate endoscopy quality, and poor decision-making around surveillance or follow-up plans were identified as the commonest explanations for PEUGIC. A negative correlation between the annual number of endoscopies performed by individual endoscopists and the technically attributable PEUGIC rate was noted.
The unadjusted PEUGIC rate was 6.7%, which was within the target of <10% proposed in a position statement on UK endoscopy quality standards [1]. However, both endoscopy providers are part of large conurbations and some patients may have been diagnosed with PEUGIC at different providers and would not have been captured in an analysis limited to local hospital records, meaning this is therefore likely to be an underestimate. Studying national datasets can circumvent this problem, as seen in the national UK PCCRC analysis [17], when 13% of PCCRCs were diagnosed in a different provider from the one that performed the index colonoscopy (personal communication from Drs. Roland Valori and Nicholas Burr).

Around 40% of PEUGICs occurred in patients with Barrett’s esophagus. A systematic review has also described an esophageal cancer miss rate of 24% in Barrett’s esophagus [18]; however, 89% of the PEUGICs in patients with Barrett’s esophagus were diagnosed at an early stage and 81% were amenable to curative endoscopic or surgical resection. These results are supported by previous studies that have shown a positive impact of Barrett’s surveillance on tumor staging and the survival of patients [19, 20]. Of the Barrett’s PEUGICs, 56% were treated by endoscopic resection and can therefore be regarded as surveillance successes; nine underwent surgical resection when earlier detection and endoscopic intervention might have avoided this outcome. The main reasons for PEUGIC included inadequate numbers of biopsies and inadequate surveillance plans. We would recommend that surveillance of Barrett’s esophagus, and gastric intestinal metaplasia and atrophy should only be performed by endoscopists with adequate training, on dedicated lists with adequate time, and using optimal mucosal enhancement techniques [21–25].

Photographs of D2 were recorded in 33% of index endoscopies among the PEUGIC patients and of retroflexion in 38%. National and international guidelines recommend photodocumentation of anatomical landmarks [1, 28–30] and the widespread availability of electronic image capture means there should be no excuse for not obtaining adequate endoscopic views.

Table 3  Summary of the results of the root cause analysis of the most plausible explanation for the 89 post-endoscopy upper gastrointestinal cancers (PEUGICs).

<table>
<thead>
<tr>
<th>Type 1</th>
<th>Premalignant lesion noted (e. g. Barrett’s esophagus, gastric intestinal metaplasia or atrophy) in the same segment as the PEUGIC</th>
<th>39 (43.8%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>a</td>
<td>Biopsies adequate and, if Barrett’s esophagus found, segment was adequately measured, and surveillance plan adequate and within correct timeframe, but PEUGIC still occurred</td>
<td>15 (16.9%)</td>
</tr>
<tr>
<td>b</td>
<td>Biopsies inadequate and/or Barrett’s segment not measured</td>
<td>17 (19.1%)</td>
</tr>
<tr>
<td>c</td>
<td>Surveillance plan inadequate</td>
<td>11 (12.4%)</td>
</tr>
<tr>
<td>d</td>
<td>Surveillance not undertaken or not undertaken within the correct timeframe but appropriate owing to patient choice or co-morbidity</td>
<td>2 (2.2%)</td>
</tr>
<tr>
<td>e</td>
<td>Surveillance not undertaken or not within the correct timeframe and inappropriate</td>
<td>3 (3.4%)</td>
</tr>
</tbody>
</table>

Type 2  Focal or cancer-associated lesion noted in the same segment as the PEUGIC (e. g. esophageal ulcer or stricture, grade C or D reflux esophagitis, gastric ulcer)

| a      | Site and morphology described and photographed, adequate biopsy sampling and follow-up undertaken in the correct timeframe but PEUGIC still occurred | 0        |
| b      | Site or morphology not described or not photographed or biopsy sampling inadequate                                              | 7 (7.9%)  |
| c      | Follow-up plan inadequate                                                                                                       | 7 (7.9%)  |
| d      | Follow-up not undertaken or not undertaken within the correct timeframe but appropriate owing to patient choice or co-morbidity | 3 (3.4%)  |
| e      | Follow-up not undertaken or not within correct timeframe and inappropriate                                                    | 0        |

Type 3  No premalignant lesion/focal or cancer-associated lesion noted in the same segment as the PEUGIC

| a      | Possible missed lesion but prior endoscopy adequate                                                                               | 8 (8.9%)  |
| b      | Possible missed lesion, with prior endoscopy not recognized by endoscopist as inadequate                                         | 28 (31.5%) |
| c      | Possible missed lesion, with prior endoscopy recognized as inadequate but follow-up plan inadequate                               | 3 (3.4%)  |
| d      | Possible missed lesion, with prior endoscopy recognized as inadequate and follow-up plan adequate, including no follow-up owing to patient choice or co-morbidity | 0        |

More than one possible explanation was found in seven cases.
images. Photodocumentation of cecal intubation and rectal retroflexion are established for colonoscopy to ensure examination completeness and the examination of high risk areas [31]. Similar efforts are needed for UGI endoscopy to ensure that high risk areas are adequately examined. Finally, an accurate description according to the established classification systems [32] is of critical importance to the ongoing management and follow-up of lesions, including correlation with histology.

We found a negative correlation between endoscopists’ annual endoscopy number and the technically attributable PEUGIC rate. The BSG recommends that endoscopists should perform a minimum of 100 procedures each year to maintain proficiency [1]; however, in the current study, all of the endoscopists who had performed a PEUGIC endoscopy where an endoscopy-related contributing factor was identified had performed more than 100 annual endoscopies. This suggests that the annual endoscopy volume currently recommended may not be adequate, but it is important to emphasize that this assessment was based on the analysis of only a small number of endoscopists. These findings also highlight that further quality indicators are needed for endoscopy.

Although Barrett’s esophagus was identified as the predominant premalignant condition in the current study, other premalignant conditions (e.g., gastric atrophy and intestinal metaplasia) may be more common in other regions. The root cause analysis system developed is however generalizable and will provide a framework to investigate PEUGIC in other settings.

The present study has a number of limitations. It was a retrospective study and although the most plausible explanations were identified, causality cannot be established. Clinical staging was not available for 11% of patients owing to the patient’s choice not to have further investigations or because they had moved out of the catchment area.

Advanced imaging techniques and longer inspection times improve the diagnostic yield of endoscopy [22, 25, 33–35]; however, the recording of these parameters was not mandatory in the endoscopy reporting systems at the study providers. Owing to uncertainty around whether these techniques were used, they were not included in the proposed criteria for an adequate endoscopy examination. We would suggest that these important measures should be included in future PEUGIC studies. The impact of patient tolerance and sedation could not be assessed, but this clearly merits further study as a contributing factor to PEUGIC.

It is possible that some of the endoscopists had performed endoscopies outside of their national health service (NHS) provider and it was not possible to capture data on these endoscopies. This could potentially bias the results of the correlation between annual endoscopy number and technically attributable PEUGIC rate.

Evaluation of only the index endoscopy, as recommended by the WEO for PCCRC, has the potential limitation of missing important information on a small number of patients in whom a premalignant, focal, or cancer-associated lesion in the same segment as the PEUGIC was detected at a prior endoscopy (before the index endoscopy), with the lesion not being seen or recognized at the index endoscopy. Future studies should consider examining all endoscopies prior to a cancer diagnosis, to identify if there is any additional benefit to reviewing all endoscopies within the 3 years prior to diagnosis. Finally, this study included two NHS providers in the UK, and the study findings and root cause analysis system should be validated in future studies in other parts of the world.

In conclusion, in a retrospective analysis of PEUGIC, the most common plausible explanations were inadequate assessment or decision-making concerning premalignant, focal, or cancer-associated lesions, and possible missed lesions in the context of an inadequate endoscopy or decision-making following endoscopy. A systematic approach using the root cause analysis framework developed can differentiate the technical endoscopic, decision-making, and administrative factors that can lead to missing UGICs at both endoscopist and institutional level, and guide quality improvement efforts to reduce the PEUGIC rate.

**Acknowledgments**

We gratefully acknowledge the advice of Dr. Roland Valori and Professor Pradeep Bhandari in the development of the PEUGIC root cause analysis system. We are also grateful for the help in data collection provided by Drs. Malik Magrabi, Sarah Faloon, Mohammed Dhanji, Rahman Hameed, Mohammed Abdul, and Mohit Inani.

**Competing interests**

The authors declare that they have no conflict of interest.

**References**


