Introductions Persistent GI symptoms and malnutrition have been associated with poorer quality of life in upper-GI cancer patients. This study aims to assess GI symptoms and nutritional status in patients undergoing modern treatment.

Methods Patients with newly diagnosed upper-GI cancer were prospectively reviewed at the time of diagnosis and at 3- and 12-months following radical treatment. Nutritional assessment was performed using the patient-generated subjective global assessment (PG-SGA), which is considered the ‘gold-standard’ for nutritional assessment and has been validated in the oncology setting (score ≥ 4 intervention needed; score ≥ 9 critical intervention needed). The gastrointestinal symptom rating scale (GSRS) was used to evaluate the presence/absence and severity of 22 GI symptoms using a 4-point response scale. Total scores range from 0–66, where 0 = all symptoms absent and 66 = all symptoms severe.

Results 61 males and 19 females, median age 66 (range 46–89) years were recruited (61% oesophageal, 33% gastric, 6% gastro-oesophageal junction tumours). Of these, 68 were reviewed at 3-months and 25 at 12-months. Mean (SD) body weight and body mass index (BMI) were 76.7 kg (17.4) and 26.7 kg/m² (4.7) at baseline, 74.4 kg (14.8) and 25.9 kg/m² (4.4) at 3-months and 72.1 kg (16.3) and 24.7 kg/m² (4.4) at 12-months. There was a significant mean difference in weight (2.0 kg, p = 0.002) and BMI (0.56 kg/m², p = 0.006) at 3-months compared to baseline. These reduced further by 12-months. Mean (SD) PG-SGA score at baseline 9.0 (6.3), 3-months 7.8 (5.6), and 12-months 7.4 (5.0) indicated that intervention was required. At baseline, 3- and 12-months 61%, 52% and 68% of patients respectively were considered moderately or severely malnourished. Mean (SD) total GSRS scores were 14.2 (10.8), 12.0 (9.4) and 15.5 (11.5) at baseline, 3- and 12-months respectively. The symptoms with the greatest increase in prevalence (% more patients) from baseline to 3-months (n = 68) were nausea (+24%), loose stool (+16%), pain swallowing (+24%), regurgitation (+21%), belching (+15%) and acid reflux (+12%). Of the n = 25 followed up at 12-months, the most common symptoms reported were flatulence (76%), belching (72%), abdominal pain (68%), abdominal grumbling (56%) and early satiety (52%).

Conclusion After treatment commences there is progressive weight loss over time. Troublesome GI symptoms persist at 12-months and may be contributing to this weight loss. Optimising nutritional status and controlling GI symptoms is required throughout the treatment pathway.

Disclosure of Interest None Declared.

Introductions Eosinophilic oesophagitis (EO) is the underlying diagnosis in at least 10% of those with dysphagia. To make the diagnosis, oesophageal biopsies showing an eosinophil count >15 per hpf are required. It is most frequent in males under aged 50 years.

Methods All patients having a gastroscopy for dysphagia were identified retrospectively for 6 consecutive years from our endoscopy reporting system. Patient demographics, endoscopic findings and whether biopsies were taken were recorded together with histology results.

Results 3068 patients had a gastroscopy with an indication of dysphagia (1489 female, age 15–100 years, average 67.7 y). The number of patients varied little between years (486–550 patients/year). Common endoscopic diagnoses were normal (20.4%), benign stricture (12.6%), oesophagitis (18.1%), Barrett’s (4.8%), dysmotility (3.7%) and hiatus hernia (10%). 1620 (52.8%) had oesophageal biopsies.

44 patients (1.5% of all patients) were diagnosed with EO, 32 of who were males. This equates to 2.8% of those who were biopsied and 4.7% of those biopsied without cancer, stricture or Barrett’s. Although only 13.3% of those with dysphagia were aged 50 years or under, they equated to 45.4% of those diagnosed with EO. Of those with EO, 6 had food bolus, 6 “typical” EO changes e.g., feline oesophagus, ridges etc, 4 an irritable oesophagus and 3 Schatzki rings.

Conclusion EO is a relatively common cause of dysphagia but is almost certainly under-recognised due to lack of oesophageal biopsies at endoscopy. Reliance on endoscopic changes of EO at endoscopy will miss the majority of cases. Although biopsying only those under 50 years would be more cost effective than biopsying all, it would also miss the majority of cases. It may be appropriate for the BSG to use frequency of oesophageal biopsies in dysphagic patients as a quality assurance measure for upper GI endoscopy.

Disclosure of Interest None Declared.

Introductions Lymphocytic duodenosis (LD) is an early marker for coeliac disease (CD). However, the majority of cases are due to non-CD related conditions.

Disclosure of Interest None Declared.