Comparison of short-term outcomes from the International Oesophago-Gastric Anastomosis Audit (OGAA), the Esophagectomy Complications Consensus Group (ECCG), and the Dutch Upper Gastrointestinal Cancer Audit (DUCA)

Oesophago-Gastric Anastomosis Study Group on behalf of the West Midlands Research Collaborative

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Abstract

Background: The Esophagectomy Complications Consensus Group (ECCG) and the Dutch Upper Gastrointestinal Cancer Audit (DUCA) have set standards in reporting outcomes after oesophagectomy. Reporting outcomes from selected high-volume centres or centralized national cancer programmes may not, however, be reflective of the true global prevalence of complications. This study aimed to compare complication rates after oesophagectomy from these existing sources with those of an unselected international cohort from the Oesophago-Gastric Anastomosis Audit (OGAA).

Methods: The OGAA was a prospective multicentre cohort study coordinated by the West Midlands Research Collaborative, and included patients undergoing oesophagectomy for oesophageal cancer between April and December 2018, with 90 days of follow-up.

Results: The OGAA study included 2247 oesophagectomies across 137 hospitals in 41 countries. Comparisons with the ECCG and DUCA found differences in baseline demographics between the three cohorts, including age, ASA grade, and rates of chronic pulmonary disease. The OGAA had the lowest rates of neoadjuvant treatment (OGAA 75.1 per cent, ECCG 78.9 per cent, DUCA 93.5 per cent; P < 0.001). DUCA exhibited the highest rates of minimally invasive surgery (OGAA 57.2 per cent, ECCG 47.9 per cent, DUCA 85.8 per cent; P < 0.001). Overall complication rates were similar in the three cohorts (OGAA 63.6 per cent, ECCG 59.0 per cent, DUCA 62.2 per cent), with no statistically significant difference in Clavien–Dindo grades (P = 0.752). However, a significant difference in 30-day mortality was observed, with DUCA reporting the lowest rate (OGAA 3.2 per cent, ECCG 2.4 per cent, DUCA 1.7 per cent; P = 0.013).

Conclusion: Despite differences in rates of co-morbidities, oncological treatment strategies, and access to minimal-access surgery, overall complication rates were similar in the three cohorts.

Introduction

Oesophageal cancer is a major cause of global mortality, accounting for more than 436 000 deaths annually¹. Late presentation frequently means that only 30–40 per cent of patients are suitable for curative treatment options^{2,3}. Oesophagectomy remains an integral part of the curative treatment in this latter group, but is associated with significant morbidity and mortality^{4,5}. Before the establishment of definitions for complications and quality measures by the Esophageal Complications Consensus Group (ECCG) in 2015⁶, it was challenging to evaluate the international variation in postoperative oesophagectomy outcomes. The ECCG has provided postoperative outcomes from 24 selected high-volume centres, setting a benchmark for high-quality oesophageal surgery reporting⁴. The Dutch Upper Gastrointestinal Cancer Audit (DUCA) has also provided detailed outcome data from a nationally centralized oesophageal cancer programme, further highlighting national and international variation in complications⁵.

Patient co-morbidity has a significant impact on postoperative outcomes and plays a critical role in achieving good outcomes^{7–9}, whether surgery involves a minimally invasive or open operation^{10,11}.

The Oesophago-Gastric Anastomosis Audit (OGAA)¹² was undertaken in 2018 to provide a comprehensive assessment of preoperative, intraoperative, and postoperative oesophagectomy outcomes, with a detailed appraisal of complications, in

Received: December 15, 2020. Accepted: January 27, 2021

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accordance with the ECCG framework. The study aimed to collect data from a large number of centres and countries, encompassing centres with varying levels of experience and patient volumes. Centre inclusion was by open invitation, as opposed to the invitation-only approach used by the international ECCG study and compulsory inclusion in DUCA, in an attempt to include this broader perspective.

The aims of the study were to report comprehensive shortterm postoperative outcomes from the OGAA cohort, and to provide detailed comparative analyses against the ECCG and DUCA benchmarking studies.

Methods

The OGAA study was run by the Oesophago-Gastric Anastomosis Study Group, on behalf of the West Midlands Research Collaborative. Centres performing oesophagectomy for cancer were invited to contribute. There was no minimum unit volume to register for the study and participation was voluntary. Committed surgeons from each country were invited to act as national lead as part of the organizing committee. This permitted language-specific dissemination of study material and advertising, to facilitate wider centre recruitment. Opportunities for voluntary participation were circulated through national surgical societies. A dedicated social media team facilitated global engagement of international oesophageal surgical centres through a number of platforms. All endeavours were made to ensure open and inclusive centre recruitment, to provide a comprehensive global cohort. The protocol for this study has been published^{12,13} along with the collaborative model that has successfully delivered a number of international and national cohort studies^{14–17}.

Centres were not required to standardize surgical or management pathways, and no changes were made to individual aspects of patient care as part of the study. Teams of surgeons, surgical trainees, research nurses or medical students prospectively identified eligible patients over a9-monthperiod from2 April2018 to31 December2018. Patients were followed for 90 daysfrom the date of surgical resection, with study follow-up closing on 31 March 2019. Data collection teams at each centre were supervised by a consultant surgeon, who took overall responsibility for local study conduct and data validation. No external data validation was performed on submitted data, in keeping with previously published data by the ECCG⁴. Data submitted to DUCA are subject to external validation to ensure completeness and accuracy.

Outcome measures

The primary aim was to assess the comparative frequency of postoperative complications (within 30 days) across the OGAA, ECCG, and DUCA cohorts. Complications were defined by the ECCG framework⁶, and classified based on the Clavien–Dindo grade¹⁸. Secondary outcomes were reoperation, readmission, and postoperative mortality rates. Outcome data for the ECCG and DUCA cohorts were sourced from the most recent publications at the time of conception of the present study. This therefore encompassed patients undergoing oesophagectomy from January 2015 to December 2016 and from January 2016 to December 2017 respectively^{4,5}. Data on oesophagectomy and gastrectomy were reported separately for the DUCA cohort, and only oesophagectomy data were included in the present analysis. As data were acquired from published materials, individual patient-level data were not available, so statistical adjustment for

Tumour staging was performed in accordance with the eighth edition of the TNM staging classification¹⁹. Positive tumour margins in the OGAA were defined as tumour identifiable at 1 mm or less, in accordance with the Royal College of Pathologists guidance²⁰. However, positive tumour margins in the ECCG and DUCA studies were defined as tumour identifiable at 0 mm, in accordance with College of American Pathologists guidance²¹. Comparison of margin positivity (R status) between the OGAA and ECCG/DUCA was therefore not possible.

Ethical approval and data sharing for OGAA

Ethical approval was dependent on local protocols and was country-specific. It was the responsibility of the local principal investigator of the enrolled unit to ensure that appropriate ethical or audit approval was gained before commencement of the study. Ongoing study approval was maintained locally throughout the duration of the study. In the UK, the study was registered at each site as either a clinical audit or service evaluation, on the basis that the information collected was routine and anonymized with no influence on the clinical care pathway.

Statistical analysis

For variables that were available for all three cohorts, comparisons were done using χ^2 tests for nominal variables, and Kruskal-Wallis tests for ordinal variables. Where significant differences were detected, post hoc pairwise comparisons were performed using χ^2 tests or Mann–Whitney *U* tests, as applicable, with Bonferroni correction for three comparisons applied to the resulting *P* values. Variables that were reported only for two cohorts were analysed using χ^2 tests or Mann–Whitney *U* tests, as applicable, All analyses were carried out using SPSS[®] version 22 (IBM, Armonk, New York, USA), with P < 0.050 deemed indicative of statistical significance throughout.

Results

Between April and December 2018, 2247 oesophagectomies for cancer were included in the OGGA. A summary of the characteristics of this study and those of the ECCG and DUCA cohorts is shown in Table 1. The OGAA included patients from 137 centres across 41 countries (106 centres in high-income countries, 31 centres in low-middle-income countries). Of centres contributing to the OGAA, 71 were located in 13 countries where oesophageal adenocarcinoma (OAC) was the predominant histological type²². The ECCG and DUCA studies included patients from fewer centres (24 and 22 respectively) in fewer countries (14 and 1 respectively). Fourteen centres contributing to the ECCG were located in six countries with a histological predominance of adenocarcinoma (OAC) over squamous cell carcinoma (SCC). The DUCA encompasses centres only in the Netherlands, where the age-standardized incidence per 100 000 population is 4.4 for OAC and 2.0 for SCC²².

The OGAA included eight centres that contributed to the ECCG, and four that contributed to the DUCA. Two centres contributed to both the ECCG and DUCA. Of the 137 centres in the OGAA cohort, the approximate case volume was less than 20, 20–60 and over 60 procedures per year in 78, 51, and eight centres, after extrapolating the 9-month numbers collected to provide an annual estimate. Centres in the DUCA and ECCG

| | OGAA | ECCG | DUCA |
|----------------------------------|--|---|--------------------------------|
| No. of centres | 137 | 24 | 22 |
| No. of countries | 41 | 14 | 1 |
| No. of patients | 2247 | 2704 | 1617 |
| Inclusion dates | Apr 2018 to Dec 2018 | Jan 2015 to Dec 2016 | Jan 2016 to Dec 2017 |
| Centre volume | Any | High volume only | > 20 resections/year |
| Centre inclusion | Open invitation (national leads dissemination, national societies, social media) | Invite only (2020— open to new applications) | Mandatory national audit |
| Type of surgery | Oesophagectomy only | Oesophagectomy only | Oesophagectomy and gastrectomy |
| Indication | Malignancy only | Any | Malignancy only |
| Definition of an involved margin | <1mm ²⁰ | 0 mm ²¹ | 0 mm ²¹ |
| Centres enrolled in OGAA | Ξ | 8 | 4 |

Table 1 Summary of study characteristics of cohort studies from the OGAA, ECCG and DUCA

OGAA, Oesophago-Gastric Anastomosis Audit; ECCG, Esophagectomy Complications Consensus Group; DUCA, Dutch Upper Gastrointestinal Cancer Audit.

studies performed an average mean of 37 and 56 oesophagectomies per year respectively.

Comparisons of patient characteristics, tumour staging, treatment, and outcomes between the three cohorts are summarized in *Tables* 2–4, with further detail reported in *Tables* S1–S3.

Patient characteristics

Baseline patient characteristics of patients in the three studies are shown in *Table* 2. The three studies had a similar sex distribution, all having a preponderance of men. A significant difference in age was observed, which increased progressively across the ECCG, OGAA, and DUCA cohorts (22.2, 29.1 and 29.7 per cent of the patients respectively were aged over 70 years; P < 0.001). The ECCG cohort tended to have lower BMI (BMI at least 25 kg/m²: OGAA 55.9 per cent, ECCG 53.1 per cent, DUCA 56.3 per cent; P = 0.037), but also had the greatest number of co-morbidities on the basis of ASA grade (ASA grade III or higher: OGAA 30.7 per cent, ECCG 38.5 per cent, DUCA 21.5 per cent; P < 0.001).

Specific co-morbidity indices were not reported, although the OGAA and DUCA reported total numbers of co-morbidities, which were significantly higher in the latter (2 or more co-morbidities: OGAA 11.2 per cent, DUCA 29.6 per cent; P < 0.001). Assessment of individual co-morbidities showed the DUCA to have the highest rate of chronic pulmonary disease, which was almost twice that of the other cohorts (OGAA 13.7 per cent, ECCG 10.5 per cent, DUCA 20.2 per cent; P<0.001). Of the other co-morbidities reported, the ECCG had the highest rates of both congestive heart failure (OGAA 3.0 per cent, ECCG 4.6 per cent, DUCA 0.7 per cent; P < 0.001) and peripheral vascular disease (OGAA 5.2 per cent, ECCG 6.8 per cent, DUCA 4.5 per cent; P = 0.03), and the OGAA had the highest rate of moderate-to-severe renal disease (OGAA 2.6 per cent, ECCG 1.3 per cent, DUCA 1.3 per cent; P = 0.001). Rates of diabetes mellitus were similar in the three cohorts (OGAA 12.0 per cent, ECCG 13.5 per cent, DUCA 14.5 per cent; P = 0.089). Comparison of pathological tumour staging between the three cohorts showed the OGAA to have more advanced tumours, with the highest T and N categories, followed by the ECCG and DUCA (pT3 or higher: OGAA 49.8 per cent, ECCG 44.6 per cent, DUCA 38.8 per cent, P < 0.001; pN+: OGAA 46.2 per cent, ECCG 42.6 per cent, DUCA 39.3 per cent; P < 0.001) (Table 3).

Treatment variation

Data on treatment and tumour staging across the three cohort studies are presented in *Table 3*. The DUCA contained higher rates of neoadjuvant chemoradiotherapy (CRT) and minimally invasive surgery (CRT rate: OGAA 35.6 per cent, ECCG 46.1 per cent, DUCA 87.8 per cent, P < 0.001; minimally invasive approach: OGAA 57.2

per cent, ECCG 47.9 per cent, DUCA 85.8 per cent; P < 0.001), with a greater proportion of three-stage operations (abdomen, chest, neck), resulting in higher rates of cervical anastomoses (OGAA 22.8 per cent, ECCG 37.9 per cent, DUCA 43.0 per cent; P < 0.001). Comparisons between the OGAA and ECCG showed smaller differences, although the former had higher rates of minimally invasive surgery, whereas the ECCG had higher rates of neoadjuvant CRT and neck anastomoses.

Postoperative outcomes

Overall complication rates were similar in the three studies (OGAA 63.6 per cent, ECCG 59.0 per cent, DUCA 62.2 per cent; P = 0.752), with no significant difference in complication severity, as classified by the highest Clavien–Dindo grade (Fig. 1). Despite this, rates of individual complication types differed between the studies (Table 4). For example, the OGAA had the highest rates of respiratory and infective complications, but significantly lower rates of gastrointestinal and cardiac complications. Rates of anastomotic leak and conduit necrosis in the OGAA were 14.2 and 2.7 per cent respectively. Combining these outcomes, it was found that the composite anastomotic leak/conduit necrosis rate differed significantly between the cohorts, being highest in the DUCA, and lowest in the ECCG cohort (OGAA 14.6 per cent, ECCG 11.4 per cent, DUCA 19.0 per cent; P < 0.001). There were no significant differences in chyle leak rates between pairs of studies.

Mortality rates at 30 and 90 days were similar in the OGAA and ECCG cohorts (90-day mortality: OGAA 4.5 per cent, ECCG 4.5 per cent; P = 0.967). The DUCA did not report 90-day deaths, but had a significantly lower 30-day mortality rate than the other studies (30-day mortality: OGAA 3.2 per cent, ECCG 2.4 per cent, DUCA 1.7 per cent; P = 0.013). The Dutch audit had a significantly higher 30-day readmission rate than the other two cohorts (OGAA 11.5 per cent, ECCG 10.2 per cent, DUCA 14.4 per cent; P < 0.001).

Discussion

Postoperative oesophagectomy outcomes across an unselected global cohort have been evaluated and compared with those in previous benchmarking studies from the ECCG and DUCA. Overall rates of complications were comparable in all three studies. The OGAA and ECCG reported similar 30-day mortality rates, but the DUCA had significantly lower mortality rates than the OGAA. This lower mortality rate may reflect a well resourced, centralized, national oesophageal cancer programme facilitating an ability to recognize and rescue the deteriorating patient. Evidence from the Agency for Healthcare Research and Quality

Table 2 Baseline demographics of OGAA, ECCG, and DUCA cohort studies

| | % of patients | | | Overall P^* | P for pairwise comparisons [§] | | |
|-------------------------------|---------------|-------------|------------|----------------------|---|----------------------|------------------|
| | OGAA | ECCG | DUCA | | OGAA versus ECCG | OGAA versus DUCA | ECCG versus DUCA |
| Male sex | 78.6 | 77.5 | 76.0 | 0.151 | _ | _ | _ |
| Age (vears) | | | | < 0.001 [†] | 0.001 [¶] | < 0.001 [¶] | < 0.001¶ |
| < 40 | 2.5 | 2.4 | 0.4 | | | | |
| 41-50 | 8.1 | 8.0 | 47 | | | | |
| 51-60 | 24.5 | 26.7 | 19.5 | | | | |
| 61–70 | 35.8 | 40.7 | 45.7 | | | | |
| 71-80 | 26.2 | 19.7 | 27.9 | | | | |
| × 80 | 20.2 | 25 | 1.8 | | | | |
| $PMI (kg/m^2)$ | 2.5 | 2.5 | 1.0 | 0.037 | 0 0391 | 1 0001 | 0 3721 |
| 2 10 E | 1 2 | 60 | 2.0 | 0.057 | 0.059 | 1.000 | 0.372 |
| < 10.J | 4.2 | 10.0 | 2.9 | | | | |
| 16.5-25.0 | 39.9 | 40.1 | 40.8 | | | | |
| 25.0-30.0 | 35.3 | 33.0 | 39.9 | | | | |
| > 30 | 20.6 | 19.5 | 16.4 | | | | |
| Smoking status | 00 C | | | - | - | - | - |
| Never smoked | 38.6 | - | - | | | | |
| Ex-smoker (> 6 weeks) | 40.3 | - | - | | | | |
| Ex-smoker (< 6 weeks) | 5.5 | - | - | | | | |
| Current smoker | 15.6 | - | - | | | | |
| ASA fitness grade | | | | < 0.001 ⁺ | < 0.001 | < 0.001 | < 0.001" |
| Ι | 13.3 | 15.2 | 15.8 | | | | |
| II | 56.1 | 46.2 | 62.7 | | | | |
| III | 29.7 | 36.7 | 21.1 | | | | |
| IV | 1.0 | 1.8 | 0.4 | | | | |
| V | 0.0 | 0.0 | 0.0 | | | | |
| No. of co-morbidities | | | | $< 0.001^{\ddagger}$ | -# | -# | —# |
| 0 | 59.3 | _ | 46.6 | | | | |
| 1 | 29.5 | _ | 23.8 | | | | |
| > 2 | 11.2 | _ | 29.6 | | | | |
| ECOG status | | | | _ | _ | _ | _ |
| 0 | 60.9 | _ | _ | | | | |
| 1 | 32.8 | _ | _ | | | | |
| > 2 | 63 | _ | _ | | | | |
| Dishetes mellitus | 0.5 | | | 0 089+ | _ | _ | _ |
| No | 87.0 | 865 | 85 5 | 0.005 | | | |
| Incomplicated | 11.2 | 12.0 | 127 | | | | |
| End organ damage | 0.8 | 12.9 | 13.7 | | | | |
| Museerdial information | 0.8 | U.0 | 0.8 | 0.006 | | | |
| Congostivo hoort foiluro | 0.4 | 5.4 4.6 | D.D D 7 | 0.220 | - | - | - 0.001 |
| Chronic nulmonary diagon | 3.U | 4.0 10 F | 0.7 | < 0.001 | 0.000 | < 0.001 | < 0.001 |
| Chronic pulmonary disease | 13./ | 10.5 | 20.2 | < 0.001 | 0.002 | < 0.001 | < 0.001 |
| Peripheral Vascular disease | 5.2 | 6.ð | 4.5 | 0.003 | 0.051 | 1.000 | 0.005 |
| Moderate–severe renai disease | 2.6 | 1.3 | 1.3 | 0.001 | 0.003 | 0.016 | 1.000 |

Data are reported only as percentages, in order to simplify the table; the associated numerators and denominators are reported in *Table S1*. OGAA, Oesophago-Gastric Anastomosis Audit; ECCG, Esophagectomy Complications Consensus Group; DUCA, Dutch Upper Gastrointestinal Cancer Audit; ECOG, Eastern Cooperative Oncology Group.^{*}χ² test, except [†]Kruskal–Wallis test and [†]Mann–Whitney U test for ordinal variables. [§]Bonferroni-corrected χ² test, except [§]Bonferroni-corrected Mann–Whitney U test for ordinal variables. [§]Conferroni-corrected χ² test, except [§]Bonferroni-corrected X² test, except [§]B

Nationwide Readmission Database²³ showed that failure-to-rescue rates were 21.2 per cent in low-volume centres, compared with 13.4 per cent in high-volume centres.

Patient selection has often been perceived as a key determinant of successful postoperative outcomes. Although definitive CRT has good outcomes for SCC, its efficacy is limited for adenocarcinoma^{24–26}. The extent to which patient selection contributed to differences in patient outcome between the three studies remains elusive. The Dutch audit reported the lowest 30-day mortality rate, but contained the highest rate of chronic pulmonary disease and had the greatest proportion of patients aged over 70 years. The OGAA cohort, on the other hand, contained more patients with locally advanced disease (higher T categories and rates of nodal positivity). This might reflect access to, and use of, diagnostic and staging modalities, as well as availability and cultural attitudes to non-surgical treatments (indicated by lower neoadjuvant treatment rates), especially in middle- and low-income societies¹³. The OGAA also had the highest rates of conduit necrosis, which may highlight the challenges of maintaining high surgical standards in lower-volume units. Despite these differences between the three studies, overall complication rates were broadly similar.

There is conflicting evidence regarding the impact of minimally invasive oesophagectomy compared with open surgery on complication rates and other outcomes^{27–30}. The TIME and MIRO trials, which demonstrated the superiority of minimally invasive techniques, have further driven rapid adoption^{27,30,31}, although it should be recognized that transition to a new operative technique can be associated with increased complications that are likely to influence outcomes outside a trial setting^{32,33}. The DUCA involved significantly more minimally invasive surgery than the OGAA and ECCG studies, yet had similar levels of complications. The explanation may be complex. Rates of respiratory complications did not differ significantly between the OGAA and DUCA, despite the significantly higher rates of chronic pulmonary disease in the DUCA cohort, but there were higher rates of open

Table 3 Treatment and tumour staging across OGAA, ECCG, and DUCA cohort studies

| | % of patients | | | Overall | P for pairwise comparisons ‡ | | | |
|----------------------------------|---------------|------|------|---------------|--|------------------|--------------------|--|
| | OGAA | ECCG | DUCA | P^{\dagger} | OGAA versus ECCG | OGAA versus DUCA | ECCG versus DUCA | |
| Neoadiuvant therapy [*] | | | | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| None | 24.9 | 21.1 | 65 | 0.001 | 0.001 | 0.001 | (0.001 | |
| Chemotherapy only | 39.1 | 29.5 | 53 | | | | | |
| Radiotherapy only | 03 | 0.2 | 0.4 | | | | | |
| CRT | 35.6 | 46.1 | 87.8 | | | | | |
| Definitive CRT | 0.0 | 3 1 | 0.0 | | | | | |
| Surgical approach | 0.0 | 5.1 | 0.0 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Open | 42.8 | 52.1 | 14.2 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| MI | 57.2 | 47.9 | 85.8 | | | | | |
| Open surgery type | 57.2 | ±7.5 | 05.0 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Thoracophdominal | 8.0 | 0.0 | 0.0 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Tranchiatal | 9.0 | 20.1 | 47.6 | | | | | |
| Transthoracic | 2.7 | 79.0 | 52.4 | | | | | |
| | 02.5 | 19.9 | J2.4 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Abdomon and chost | E1 G | 107 | 70.7 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Abdomon only | 12.0 | 40.7 | 16.0 | | | | | |
| Chast only | 42.0 | 40.2 | 10.0 | | | | | |
| A masternasia site | 0.4 | 11.1 | 4.5 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Chast | 77.0 | CO 7 | E4 0 | < 0.001 | < 0.001 | < 0.001 | < 0.001 | |
| Criest | 77.0 | 60.7 | 54.2 | | | | | |
| Neck | 22.8 | 37.9 | 43.0 | | | | | |
| Abdomen | 0.0 | 0.0 | 0.4 | | | | | |
| No anastomosis | 0.2 | 1.4 | 2.4 | 0.004 | 0.001 | 0.005 | 0.001 | |
| Gastric tube | 100.0 | 06.0 | 00.4 | < 0.001 | < 0.001 | 0.005 | < 0.001 | |
| Stomach | 100.0 | 96.0 | 99.4 | | | | | |
| Colon | 0.0 | 1.3 | 0.3 | | | | | |
| Small bowel | 0.0 | 2.7 | 0.0 | | | | | |
| Roux-en-Y | 0.0 | 0.0 | 0.3 | | | 0.004 | | |
| Pathological T category | | | | < 0.001 | 0.009 | < 0.001 | < 0.001 | |
| p'l'x /'l'is | 2.0 | 2.1 | 0.0 | | | | | |
| p10-12 | 48.2 | 53.3 | 61.2 | | | | | |
| pT3 | 45.8 | 41.6 | 37.5 | | | | | |
| pT4 | 4.0 | 3.0 | 1.3 | | | | | |
| Pathological N status | | | | < 0.001 | 0.008 | < 0.001 | 0.324 | |
| pNx | 0.0 | 0.3 | 0.3 | | | | | |
| pN– | 53.8 | 57.1 | 60.5 | | | | | |
| pN+ | 46.2 | 42.6 | 39.3 | | | | | |
| Pathological M status | | | | < 0.001 | < 0.001 | 0.516 | < 0.001 | |
| pMx | 0.9 | 14.3 | 1.1 | | | | | |
| pM– | 96.8 | 83.9 | 97.4 | | | | | |
| pM+ | 2.3 | 1.8 | 1.5 | | | | | |
| Resection margin [°] | | | | _1 | _1 | _1 | 0.002 [§] | |
| RO | 81.8 | 93.4 | 95.9 | | | | | |
| R1 | 18.2 | 6.1 | 4.1 | | | | | |
| R2 | 0.0 | 0.5 | 0.1 | | | | | |

Data are reported only as percentages, in order to simplify the table; the associated numerators and denominators are reported in Table S2. The Esophagectomy Complications Consensus Group (ECCG) data exclude 119 patients who did not have cancer. OGAA, Oesophago-Gastric Anastomosis Audit; DUCA, Dutch Upper Gastrointestinal Cancer Audit; CRT, chemoradiotherapy MI, minimally invasive. $^{\dagger}\chi^2$ test; ‡ Bonferroni-corrected χ^2 test, except $^{\$}$ Bonferroni-corrected Mann–Whitney U test for ordinal variables. The OGAA used a different definition of margin involvement from the other cohorts, so comparisons were not meaningful.

surgery in the OGAA cohort suggesting that the minimally invasive approach may have offset the risk of pulmonary complications in the DUCA cohort.

CRT was also used more frequently in the DUCA cohort, where 87.8 per cent of patients received this treatment. Global variations in neoadjuvant treatment options are largely explained by centres that favour neoadjuvant chemotherapy based on evidence from the MAGIC, OE02, and OE05 trials^{34–36}. High uptake of CRT in the Netherlands is likely to have been influenced by the success of the CROSS trial³⁷. The absence of increased rates of overall complications in the DUCA cohort compared with the other studies supports existing evidence that neoadjuvant CRT does not increase overall complications³⁸.

Anastomotic leakage is generally regarded as a serious complication of oesophagectomy because of the risk of associated sepsis. Leak rates were highest in the DUCA group probably reflecting the high rates of anastomoses performed in the neck, and the highest rates of minimally invasive surgery, both of which are recognized to contribute to higher leak rates^{26,38-41}. The extent to which neoadjuvant CRT contributes to anastomotic leak is controversial^{38,42,43}, including whether the anastomosis lies within the radiation field or whether the stomach has been irradiated^{44,45}. The combination of chronic pulmonary disease and CRT has been shown to potentially double rates of anastomotic leakage⁴⁶, so these features could also have contributed to the higher leak rates in the DUCA cohort. Despite having the highest rate of anastomotic leak, the DUCA cohort had the lowest 30-day mortality rate, suggesting that anastomotic leaks *per se* are not critical determinants of mortality, or that a cervical leak is less likely to result in death than an intrathoracic leak. The higher readmission rate in the DUCA cohort may indicate a lower threshold for readmission that may itself have influenced outcomes.

A specific focus of the OGAA study was capturing clinically relevant data at a patient-, disease- and operation-specific level

Table 4 Intraoperative and postoperative outcomes across the OGAA, ECCG, and DUCA cohort studies

| $ \begin{array}{ c c c c c c c c c c c c c c c c c c c$ | | % of patients | | | Overall P^{\dagger} | P for pairwise comparisons $^{\mathbb{S}}$ | | |
|---|-----------------------------------|---------------|------|------|-----------------------|--|------------------|--------------------|
| | | OGAA | ECCG | DUCA | | OGAA versus ECCG | OGAA versus DUCA | ECCG versus DUCA |
| Highest Clavien-Dindo grade 0.752 ⁴ - | Intraoperative complications | 2.5 | _ | 5.5 | < 0.001 | _# | _# | _# |
| No complication36.441.037.8I12.07.59.4II26.220.423.7IIIA9.714.212.0IIB7.1668.0IVA4.36.46.9IVB1.21.30.7V3.22.61.7Gastrointestinal complications11.522.424.2Volta0.001<0.001 | Highest Clavien–Dindo grade | | | | 0.752 [‡] | - | - | - |
| $\begin{array}{cccccccccccccccccccccccccccccccccccc$ | No complication | 36.4 | 41.0 | 37.8 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | I | 12.0 | 7.5 | 9.4 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | II | 26.2 | 20.4 | 23.7 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | IIIA | 9.7 | 14.2 | 12.0 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | IIIB | 7.1 | 6.6 | 8.0 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | IVA | 4.3 | 6.4 | 6.9 | | | | |
| $ \begin{array}{c c c c c c c c c c c c c c c c c c c $ | IVB | 1.2 | 1.3 | 0.7 | | | | |
| Gastrointestinal complications11.522.424.2< 0.001< 0.001< 0.0010.501Thrombotic complications2.95.22.8< 0.001< 0.001< 0.0011.000< 0.001Anastomotic leak85.888.981.1< 0.001*< 0.008*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001*< 0.001* </td <td>V</td> <td>3.2</td> <td>2.6</td> <td>1.7</td> <td></td> <td></td> <td></td> <td></td> | V | 3.2 | 2.6 | 1.7 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | Gastrointestinal complications | 11.5 | 22.4 | 24.2 | < 0.001 | < 0.001 | < 0.001 | 0.501 |
| Anastomotic leak $< 0.001^{\ddagger}$ $0.008^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ $< 0.001^{\$}$ < 0 | Thrombotic complications | 2.9 | 5.2 | 2.8 | < 0.001 | < 0.001 | 1.000 | < 0.001 |
| No leak85.888.981.1Type 17.03.35.7Type 23.44.88.1Type 33.83.05.1Conduit necrosis $< 0.001^{4}$ $< 0.001^{1}$ $< 0.001^{1}$ 0.702^{1} No necrosis97.398.899.2 $< 0.001^{1}$ $< 0.001^{1}$ 0.702^{1} Type 11.20.10.1 $< 0.001^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ Type 30.80.90.6 $< 0.001^{1}$ $< 0.002^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ necrosis rate $< 0.8^{1}$ 0.4.1 0.043^{1} 0.061^{1} 0.203^{1} 1.000^{1} Vocal cord injury4.64.7 7.2^{2} $< 0.001^{1}$ 0.002^{2} 0.002^{2} Respiratory complications35.927.8 32.7^{2} $< 0.001^{1}$ 0.002^{2} 0.002^{2} Diaphragmatic complications1.8 2.9^{1} 1.9^{2} 0.001^{1} $< 0.001^{1}$ $< 0.001^{1}$ Diaphragmatic complications1.8 2.9^{1} 1.9^{2} 0.046^{1} 0.001^{2} $< 0.001^{1}$ Diaphragmatic complications 5.9^{1} 8.3^{1} 4.1^{2} $< 0.001^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ Urological complications 5.9^{1} 8.3^{1} 4.1^{2} $< 0.001^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ Urological complications 5.9^{1} 8.3^{1} 4.1^{2} $< 0.001^{1}$ $< 0.001^{1}$ $< 0.001^{1}$ <td>Anastomotic leak</td> <td></td> <td></td> <td></td> <td>< 0.001[‡]</td> <td>0.008[¶]</td> <td>< 0.001 1</td> <td>< 0.001¶</td> | Anastomotic leak | | | | < 0.001 [‡] | 0.008 [¶] | < 0.001 1 | < 0.001¶ |
| $\begin{array}{c ccccccccccccccccccccccccccccccccccc$ | No leak | 85.8 | 88.9 | 81.1 | | | | |
| $\begin{array}{cccccccccccccccccccccccccccccccccccc$ | Type 1 | 7.0 | 3.3 | 5.7 | | | | |
| Type 3 3.8 3.0 5.1 < 0.001 [‡] < 0.001 [¶] < 0.001 [¶] 0.702 [¶] No necrosis 97.3 98.8 99.2 | Type 2 | 3.4 | 4.8 | 8.1 | | | | |
| $\begin{array}{c c c c c c c c c c c c c c c c c c c $ | Type 3 | 3.8 | 3.0 | 5.1 | | | | |
| No necrosis 97.3 98.8 99.2 Type 1 1.2 0.1 0.1 Type 2 0.7 0.3 0.2 Type 3 0.8 0.9 0.6 Combined anastomotic leak/conduit necrosis rate 14.6 11.4 19.0 < 0.001 0.002 < 0.001 < 0.001 Nonecrosis rate V V < 0.001 0.002 < 0.001 < 0.001 Vocal cord injury 4.6 4.7 7.2 < 0.001 0.002 0.002 0.002 Respiratory complications 35.9 27.8 32.7 < 0.001 0.011 0.002 1.000 Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Infective complications 1.8 2.9 1.9 0.020 0.046 1.000 0.033 Urological complications 5.9 8.3 4.1 < 0.001 < 0.001 < 0.001 < 0.001 Urological complications 5.9 8 | Conduit necrosis | | | | < 0.001 [‡] | < 0.001 ¶ | < 0.001 1 | 0.702 [¶] |
| Type 11.20.10.1Type 20.70.30.2Type 30.80.90.6Combined anastomotic leak/conduit14.611.419.0< 0.0010.002< 0.001< 0.001necrosis rateChyle leak5.44.04.10.0430.0610.2031.000Vocal cord injury4.64.77.2< 0.0011.0000.0020.002Respiratory complications35.927.832.7< 0.001< 0.0010.1170.002Cardiac complications13.116.817.1< 0.001< 0.0010.0021.000Diaphragmatic complications1.82.91.90.0200.0461.0000.108Infective complications1.82.91.90.0200.0461.0000.013Urological complications1.82.91.90.021< 0.001< 0.001< 0.001Urological complications1.9.414.27.4< 0.001< 0.001< 0.001< 0.001Olday mortality3.22.41.70.0130.3150.0110.31830-day mortality4.54.5-0.967-##### | No necrosis | 97.3 | 98.8 | 99.2 | | | | |
| $\begin{array}{c ccccccccccccccccccccccccccccccccccc$ | Type 1 | 1.2 | 0.1 | 0.1 | | | | |
| Type 3 0.8 0.9 0.6 Combined anastomotic leak/conduit necrosis rate 14.6 11.4 19.0 < 0.001 0.002 < 0.001 < 0.001 Chyle leak 5.4 4.0 4.1 0.043 0.061 0.203 1.000 Vocal cord injury 4.6 4.7 7.2 < 0.001 1.000 0.002 0.002 Respiratory complications 35.9 27.8 32.7 < 0.001 < 0.001 0.117 0.002 Cardiac complications 13.1 16.8 17.1 < 0.001 < 0.001 0.002 1.000 Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Infective complications 1.8 2.9 1.9 0.020 0.046 1.000 0.018 Urological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Urological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Solday readmission 11.5 10.2 14.4 <td>Type 2</td> <td>0.7</td> <td>0.3</td> <td>0.2</td> <td></td> <td></td> <td></td> <td></td> | Type 2 | 0.7 | 0.3 | 0.2 | | | | |
| Combined anastomotic leak/conduit 14.6 11.4 19.0 < 0.001 0.002 < 0.001 < 0.001 necrosis rate Chyle leak 5.4 4.0 4.1 0.043 0.061 0.203 1.000 Vocal cord injury 4.6 4.7 7.2 < 0.001 1.000 0.002 0.002 Respiratory complications 35.9 27.8 32.7 < 0.001 < 0.001 0.117 0.002 Cardiac complications 13.1 16.8 17.1 < 0.001 < 0.001 0.002 1.000 Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.018 Infective complications 19.4 14.2 7.4 < 0.001 < 0.001 < 0.001 < 0.001 Urological complications 5.9 8.3 4.1 < 0.001 < 0.001 < 0.001 < 0.001 Urological complications 5.9 8.3 4.1 < 0.001 < 0.003 0.038 < 0.001 Outogat a c | Type 3 | 0.8 | 0.9 | 0.6 | | | | |
| necrosis rate Chyle leak 5.4 4.0 4.1 0.043 0.061 0.203 1.000 Vocal cord injury 4.6 4.7 7.2 < 0.001 | Combined anastomotic leak/conduit | 14.6 | 11.4 | 19.0 | < 0.001 | 0.002 | < 0.001 | < 0.001 |
| Chyle leak 5.4 4.0 4.1 0.043 0.061 0.203 1.000 Vocal cord injury 4.6 4.7 7.2 < 0.001 | necrosis rate | | | | | | | |
| Vocal cord injury4.64.77.2< 0.0011.0000.0020.002Respiratory complications 35.9 27.8 32.7 < 0.001 | Chyle leak | 5.4 | 4.0 | 4.1 | 0.043 | 0.061 | 0.203 | 1.000 |
| Respiratory complications 35.9 27.8 32.7 < 0.001 0.117 0.002 Cardiac complications 13.1 16.8 17.1 < 0.001 < 0.001 0.002 1.000 Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Infective complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Urological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Vological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Vological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Return to theatre 12.0 15.7 12.9 < 0.001 0.423 0.022 < 0.001 30-day mortality 3.2 2.4 1.7 0.013 0.315 0.011 0.318 90-day mortality 4.5 4.5 - | Vocal cord injury | 4.6 | 4.7 | 7.2 | < 0.001 | 1.000 | 0.002 | 0.002 |
| Cardiac complications 13.1 16.8 17.1 < 0.001 < 0.001 0.002 1.000 Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Infective complications 19.4 14.2 7.4 < 0.001 | Respiratory complications | 35.9 | 27.8 | 32.7 | < 0.001 | < 0.001 | 0.117 | 0.002 |
| Diaphragmatic complications 1.8 2.9 1.9 0.020 0.046 1.000 0.108 Infective complications 19.4 14.2 7.4 < 0.001 < 0.001 < 0.001 < 0.001 Urological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Return to theatre 12.0 15.7 12.9 < 0.001 0.423 0.022 < 0.001 30-day readmission 11.5 10.2 14.4 < 0.001 0.423 0.022 < 0.001 30-day mortality 3.2 2.4 1.7 0.013 0.315 0.011 0.318 90-day mortality 4.5 4.5 - 0.967 -# -# -# | Cardiac complications | 13.1 | 16.8 | 17.1 | < 0.001 | < 0.001 | 0.002 | 1.000 |
| Infective complications 19.4 14.2 7.4 < 0.001 < 0.001 < 0.001 < 0.001 Urological complications 5.9 8.3 4.1 < 0.001 | Diaphragmatic complications | 1.8 | 2.9 | 1.9 | 0.020 | 0.046 | 1.000 | 0.108 |
| Urological complications 5.9 8.3 4.1 < 0.001 0.003 0.038 < 0.001 Return to theatre 12.0 15.7 12.9 < 0.001 | Infective complications | 19.4 | 14.2 | 7.4 | < 0.001 | < 0.001 | < 0.001 | < 0.001 |
| Return to theatre 12.0 15.7 12.9 < 0.001 < 0.001 1.000 0.033 30-day readmission * 11.5 10.2 14.4 < 0.001 | Urological complications | 5.9 | 8.3 | 4.1 | < 0.001 | 0.003 | 0.038 | < 0.001 |
| 30-day readmission * 11.5 10.2 14.4 < 0.001 | Return to theatre | 12.0 | 15.7 | 12.9 | < 0.001 | < 0.001 | 1.000 | 0.033 |
| 30-day mortality 3.2 2.4 1.7 0.013 0.315 0.011 0.318 90-day mortality 4.5 4.5 - 0.967 -# -# -# | 30-day readmission | 11.5 | 10.2 | 14.4 | < 0.001 | 0.423 | 0.022 | < 0.001 |
| 90-day mortality 4.5 4.5 – 0.967 – [#] – [#] – [#] | 30-day mortality | 3.2 | 2.4 | 1.7 | 0.013 | 0.315 | 0.011 | 0.318 |
| | 90-day mortality | 4.5 | 4.5 | - | 0.967 | _# | _# | _# |

Data are reported only as percentages, in order to simplify the table; the associated numerators and denominators are reported in *Table S3*. The Oesophago-Gastric Anastomosis Audit (OGAA) data exclude 79 patients who either died in hospital, or for whom follow-up was not available. ECCG, Esophagectomy Complications Consensus Group; DUCA, Dutch Upper Gastrointestinal Cancer Audit; $^{+}\chi^{2}$ test, except [‡]Kruskal–Wallis test for ordinal variables. [§]Bonferroni-corrected χ^{2} test, except [‡]Bonferroni-corrected Mann–Whitney U test for ordinal variables. [‡]Pairwise comparisons not applicable as data available for only two cohorts.



Fig. 1 Complication rate by highest Clavien–Dindo grade in OGAA, ACCG, and DUCA studies

OGAA, Oesophago-Gastric Anastomosis Audit; ECCG, Esophagectomy Complications Consensus Group; DUCA, Dutch Upper Gastrointestinal Cancer Audit. P = 0.752 (Kruskal–Wallis test).

to minimize reporting bias in the study. The short duration of data collection was designed to minimize effects due to changes in practice. Despite the overall success in achieving these goals, there are limitations. ECCG and DUCA data were extracted from relevant publications. Data were not available at a patient level; therefore, statistical adjustment for differing preoperative and intraoperative factors was not possible when evaluating outcomes. The inclusion of patients without cancer in the ECCG and DUCA cohorts should be noted, and it was not possible to comment on the success of reported evaluation of neoadjuvant treatments and how this may have influenced short-term outcomes. Different pathological classifications were used to determine margin positivity, precluding comparisons between all three studies^{20,21,46}. For the OGAA, data were verified by each unit's lead investigator, although no specific data verification process was undertaken. Previous data verification in national and international observational studies has shown high accuracy^{4,14,47–49}. A standardized internationally agreed data set covering not only complications, but demographic, oncological, surgical, and pathological data, as developed for pancreatoduodenectomy⁵⁰, seems desirable to make fair comparisons that can result in quality improvements.

The outcome data presented by both the DUCA and ECCG represent high-quality care in centralized and selected settings. The OGAA sought to identify whether these outcomes were achievable in an unselected global cohort. Despite variations in patient demographics, resources, and surgical volumes, the similarity in overall complication rates in all three studies suggests that oesophagectomy can be performed safely at an international level. The present study has also highlighted fundamental shortcomings when comparing international outcome data for oesophagectomy. The development of a standardized data set for future studies should be considered.

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Acknowledgements

The authors are grateful to the Birmingham Surgical Trials Consortium at the University of Birmingham for use of its servers for secure online data collection.

Disclosure. The authors declare no conflict of interest.

Supplementary material

Supplementary material is available at BJS Open online.

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