

Core outcome set for surgical trials in gastric cancer (GASTROS study): international patient and healthcare professional consensus

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Abstract

Background: Surgery is the primary treatment that can offer potential cure for gastric cancer, but is associated with significant risks. Identifying optimal surgical approaches should be based on comparing outcomes from well designed trials. Currently, trials report different outcomes, making synthesis of evidence difficult. To address this, the aim of this study was to develop a core outcome set (COS)—a standardized group of outcomes important to key international stakeholders—that should be reported by future trials in this field.

Methods: Stage 1 of the study involved identifying potentially important outcomes from previous trials and a series of patient interviews. Stage 2 involved patients and healthcare professionals prioritizing outcomes using a multilingual international Delphi survey that informed an international consensus meeting at which the COS was finalized.

Results: Some 498 outcomes were identified from previously reported trials and patient interviews, and rationalized into 56 items presented in the Delphi survey. A total of 952 patients, surgeons, and nurses enrolled in round 1 of the survey, and 662 (70 per cent) completed round 2. Following the consensus meeting, eight outcomes were included in the COS: disease-free survival, disease-specific survival, surgery-related death, recurrence, completeness of tumour removal, overall quality of life, nutritional effects, and 'serious' adverse events.

Conclusion: A COS for surgical trials in gastric cancer has been developed with international patients and healthcare professionals. This is a minimum set of outcomes that is recommended to be used in all future trials in this field to improve trial design and synthesis of evidence.

Introduction

Gastric cancer is a significant global health burden which is associated with poor survival¹. Although the adoption of multimodal

therapy for the minority of patients who present with early-stage disease has improved prognosis, surgery remains the only treatment offering a potential cure². Identifying the optimum surgical

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approach involves balancing the benefits of a radical oncological resection against the risk and impact of associated complications and physiological consequences. The ability to compare outcomes from surgical trials in a clinically meaningful manner is crucial to this process.

Homogeneity in the selection and reporting of key outcomes between studies is necessary if useful synthesis of evidence is to be achieved. However, outcome reporting in surgical trials for gastric cancer is heterogeneous and not based on methodologically robust standards³. Even when similar outcomes are reported, different definitions, measurement instruments, and time points are used. Likewise, patient priorities and perspectives tend to be overlooked when outcomes are selected by researchers. This potentially limits the subsequent relevance of aspects of the research effort to the most important stakeholder group⁴. For example, quality of life, an area identified as vitally important to patients, is reported in less than 10 per cent of trials³.

To address this challenge, the GASTROS study (Gastric cancer Surgery Trials Reported Outcomes Standardization) was undertaken to develop a core outcome set (COS) for surgical trials in gastric cancer⁵. A COS is 'an agreed, standardized collection of outcomes which should be measured and reported, as a minimum, in all trials for a specific clinical area'⁶. Outcomes should be relevant to key stakeholders, who should contribute to the stages of COS development.

These challenges in outcome reporting are not limited to the field of gastric cancer and affect virtually all clinical areas. COMET (Core Outcome Measures in Effectiveness Trials; <https://www.comet-initiative.org>) is an initiative that aims to promote COS development⁷. Its registry database and up-to-date systematic reviews have comprehensively mapped COS projects across all disciplines^{8–12}. Although groups have developed COSs for different gastrointestinal cancers^{13–15}, there has yet to be one developed for gastric cancer. The global incidence of gastric cancer and differences in patient characteristics, management, and outcomes, necessitated an international approach to this COS^{16,17}. An International Working Group (IWG) of collaborators was set up to support this project, aided by a comprehensive network of patient organizations, charities, and professional bodies across six continents.

Methods

The GASTROS study conforms to standards established for the development of COS as outlined by COS-STAD (Core Outcome Set—STAndards for Development)¹⁸. This report uses the COS-STAR (Core Outcome Set—STAndards for Reporting) standards to describe the development of a COS for surgical trials in gastric cancer¹⁹. The checklist is provided in [Table S1](#).

Scope

The scope, objectives, and methodological approaches of this study have been described previously in detail⁵. In summary, the COS developed in this study was aimed at all clinical effectiveness trials examining therapeutic surgical trials for patients with early-stage (potentially curable) gastric cancer. The GASTROS study used existing best-practice approaches as developed by the COMET Initiative, while adapting the methodological principles to the challenges of an international consensus exercise. An overview of the study stages is shown in [Fig. 1](#). This publication describes stages 1 and 2.

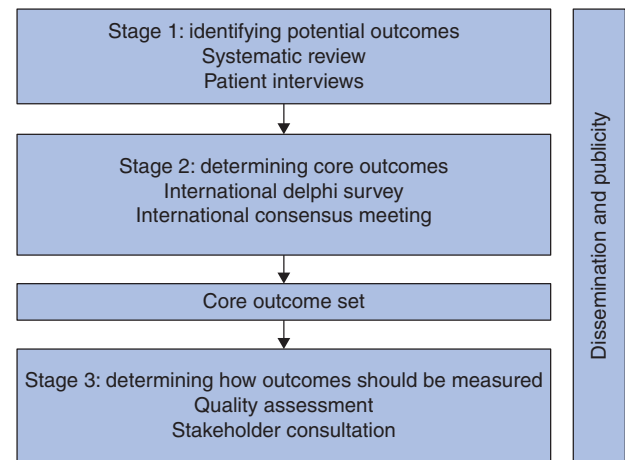


Fig. 1 Stages of the GASTROS study

Stakeholder participants and eligibility

The GASTROS study aimed to consider the views of key stakeholders during the process, namely patients, surgeons, and oncology nurses. The guiding principle was to promote the patient voice as they are the beneficiaries of trials in this field and have all-important lived experience. Surgeons provide a clinical perspective and the experience of treating large volumes of patients. Oncology nurses were invited to participate given their central roles as caregivers, patient advocates, and core members of the clinical team. Participation in the study was open to all interested stakeholders who fulfilled the following criteria: surgeons who had completed their training and routinely treat gastric cancer, oncology nurses with a recognized proportion of their role involved in the care and follow-up of patients with gastric cancer, and patients who have undergone surgical resection for gastric cancer with the intention of cure. Patients and healthcare professionals were identified through local, regional, and national clinical and research networks. Support from patient groups, charities, and professional societies was also key.

Stages of GASTROS study

Stage 1 ([Fig. 1](#)) identified outcomes that may be important to stakeholders and stage 2 subsequently prioritized outcomes for inclusion in the final COS (what to measure). In addition, the GASTROS study aimed to collate the corresponding outcome measurement instruments used in surgical trials and to determine the variability in measurement time points, for use in future outcomes research (to determine how and when to measure) in gastric cancer.

Stage 1: identifying outcomes

A systematic review of surgical trials for gastric cancer over two decades was undertaken from which all reported outcomes were extracted verbatim³. Patient-reported outcome measurement instruments used in these trials were broken down into their component parts to identify additional outcomes. To ensure that patients' perspectives were captured, a series of qualitative interviews exploring outcome prioritization was undertaken to identify potentially important outcomes not identified in the systematic review⁴. A subsequent long list of potentially important outcomes was compiled which underwent a process of rationalization, before being presented to the stakeholder groups for prioritization in stage 2.

The rationalization process (Table S2) was initiated through discussion within the Study Management Group (SMG) to merge closely related items and map them against a taxonomy developed for COSs²⁰. This process was assessed independently by an external methodologist with extensive experience in COS development. The resulting short list was presented to stakeholder representatives (patients, surgeons, and oncology nurses) comprising the Study Advisory Group (SAG). The SAG was tasked with ratifying the process so far, further merging of outcomes if required, developing plain-language descriptions of the outcomes, and identifying additional outcomes that they believed had not yet been identified. Following this, the short list of outcomes and corresponding plain-language descriptions were presented to a patient group as part of a cognitive debriefing exercise to ensure understanding and comprehensibility.

Stage 2: prioritizing outcomes

To prioritize which items to include in the COS, patients, surgeons, and oncology nurses were invited to participate in an international, multilanguage Delphi survey. The methodological approach used to translate the surveys and recruit participants has been described separately²¹. Although there is no formal sample size requirement for Delphi surveys, recruitment was facilitated with the support of a large network of professional bodies, patient groups, and charities, to help ensure that a representative spectrum of opinion was captured for each stakeholder group.

Participants were invited to score outcomes in terms of importance on a Likert-type scale ranging from 1 to 9 (1–3, not important; 4–6, important; 7–9, critically important) in a two-round online Delphi survey. Participants were given the opportunity to add outcomes they considered were missing, for consideration by participants in round 2. Suggested additional outcomes were considered by the SMG and reviewed independently by an external methodologist with experience of cancer-related surgical COSs. The scores of each stakeholder group were collated and summarized separately to ensure an equal voice among stakeholder groups. Participants who had completed 50 per cent of the first survey were included in the round 1 analysis and invited to participate in round 2. They were then presented with group scores (presented as score distribution charts) for each stakeholder group from round 1, and given the opportunity to reflect on the opinions of others before deciding whether to change their scores for each outcome in round 2. Those who changed scores between rounds were able to provide a reason for this¹⁸. After two rounds of voting, outcomes were categorized according to predetermined criteria for inclusion in, or exclusion from, the COS. Participants who had completed at least 50 per cent of the survey in round 2 were included in the final analysis.

Any outcome scored as critically important (7–9) by more than 70 per cent and not important (1–3) by less than 15 per cent in all three stakeholder groups was categorized for inclusion. Any outcome scored as critically important (7–9) by less than 50 per cent in all three stakeholder groups was categorized for exclusion. These criteria were adapted from established COS methodology⁶. Outcomes achieving any other combination of scores were categorized as not having reached consensus (no consensus) and presented for further discussion at a consensus meeting.

Survey participants were invited to attend a consensus meeting in Manchester (UK) during March 2020. The aim of the meeting was to review the results of the Delphi survey and consider the outcomes for which no consensus was reached before finalizing the COS. Participants could take part by attending the

meeting venue in person, or through an online platform. The meeting was undertaken in English and chaired by a clinical academic from the SMG with experience in COS development, and with no clinical expertise in the management of gastric cancer.

Following discussion, stakeholders were asked to score outcomes using the same criteria as was set out in the Delphi survey. Similarly, scores from each stakeholder group were considered separately to mitigate for imbalance in the numbers of each participant type. Turning Point software (Turning Technologies, Youngstown, OH, USA) was used to support voting at the venue and online simultaneously. Participants were also asked to complete an online voting form to mitigate against software malfunction. Outcomes reaching the original consensus criteria for inclusion in the final COS were to be added to those included from the Delphi survey.

Assessing bias

To assess the impact of attrition bias between survey rounds, mean scores of participants completing both rounds of the Delphi survey were compared against those of participants completing round 1 alone. Mean scores of those who took part in the Delphi survey but did not attend the consensus meeting were compared against the mean scores of those who attended to assess the degree to which consensus meeting participants were representative of those who participated in the survey. Both analyses were undertaken using a *t* test to examine for statistically significant differences at the 0.05 level. Furthermore, the characteristics of stakeholders participating in both rounds were compared with those who only completed round 1. A descriptive analysis was undertaken, and the *c*² test applied to examine for statistically significant differences at the 0.05 level.

Patient and public involvement

A guiding principle of the GASTROS study was that patients' voices should be represented at each stage of the project. Patient representatives were integral with membership in the SAG and support from international charities. The dissemination of results from this study will be supported by a network of international charities and patient support groups. The patient representatives opted to participate in the patient-focused study report of results rather than the scientific dissemination.

Study registration and protocol

The GASTROS study was registered in the COMET database (<http://www.comet-initiative.org/studies/details/764?result=true>) before commencement. The study protocol has been described previously⁵.

Ethical approvals and portfolio adoption

Ethical approvals were required for the qualitative patient interviews and international Delphi surveys. The qualitative interview study was given ethical approval by the National Research Ethics Service North West—Cheshire (11/NW/0739) and governance approvals by Central Manchester University Hospital National Health Service (NHS) Foundation Trust. The Delphi survey was given ethical approval by the North West—Greater Manchester East Research Ethics Committee (18/NW/0347) and governance approvals by Manchester University Hospitals NHS Foundation Trust. Both the patient interviews and Delphi survey were adopted on to the National Institute for Health Research (NIHR) Clinical Research Network Portfolio (IDs 33312 and 38318). Ethical approval for international participants was sought and obtained locally by IWG collaborators.

Results

The results for each stage of the study are summarized in Fig. 2. The 498 outcomes identified from the systematic review, patient-reported outcome measures, and patient interviews were rationalized by the SMG into 58 items, which were presented to the SAG. The SAG merged chyle leak, nutritional complications, respiratory function, surgical-site infection, and time to ambulation into other existing outcomes. Bleeding, anaesthetic complications, and destination on discharge were expanded or added as separate outcomes, which meant that a total of 56 items were presented to participants in the Delphi survey (Table S3).

Delphi survey

A total of 1021 patients, surgeons, and oncology nurses registered for the Delphi survey, of whom 952 (268 patients, 445 surgeons, 239 nurses) from 55 countries across six continents fulfilled the criteria for inclusion in the round 1 analysis. Table 1 summarizes the characteristics of those included in the analyses. Table S4 details the results of voting in both rounds. One additional outcome (duration of stay in an intensive care ward) suggested by participants in round 1 was presented in round 2 along with the original 56 outcomes for rescoring (a total of 57 outcomes in round 2). Although other outcomes were suggested by participants in round 1, these were deemed by the SMG and an external reviewer as either direct duplication of outcomes already included or not sufficiently unique that they warranted being presented separately (Table S5). Scores from 662 participants in round 2 were included in the final analysis representing an attrition rate of 30 per cent. Some 557 participants (84 per cent) changed the score of least one answer from round 1, with 191 (29 per cent) participants changing a score to cross a boundary (for example, from 1–3 to 4–6 or 7–9). A detailed analysis exploring the reasons for changing scores has been reported previously¹⁸.

Consensus was reached to include 13 outcomes: disease-free survival, disease-specific survival, surgery-related death, recurrence of cancer, completeness of tumour removal, overall quality of life, nutritional effects, all-cause complications, intraoperative complications, anaesthetic complications, anastomotic complications, multiple organ failure, and bleeding.

Thirteen outcomes were categorized for exclusion, and no consensus was reached for 31 outcomes, which were subsequently discussed at the consensus meeting. An analysis exploring the relationship between participant characteristics (such as regional and demographic differences) and their impact on how outcomes were scored is under review²².

There was no statistically significant difference between the mean scores of participants completing both survey rounds and those completing round 1 only (mean(s.d.) difference 0.17(0.1), largest difference 0.4; $P = 0.76$).

Consensus meeting

Forty-three Delphi survey participants (7 patients, 29 surgeons, 7 nurses) attended the consensus meeting in person (18) or using the online platform (25). Fourteen countries from four continents (South America, North America, Europe, Asia) were represented. A full breakdown of the regional origin of participants is described in Appendix S1. The difference in mean scores between consensus meeting participants and those completing round 2 of the survey was statistically significant (mean(s.d.) difference 0.3(0.23), largest difference 1.16; $P < 0.001$).

In preparation for the consensus meeting, the SMG reviewed and discussed the Delphi results. Of the 13 outcomes that reached consensus to be included, six related to perioperative complications. The SMG took the view that, as the outcome all-cause complications was voted for inclusion, by extension, all complications would need to be measured and reported by researchers as a minimum. However, 14 complication-type outcomes from the list of 57 did not reach consensus for inclusion and a further two outcomes reached consensus for exclusion from the COS. The SMG decided to present this seemingly contradictory position at the consensus meeting for further discussion and voting on a desired final position.

After an interactive debate, participants were asked to vote for one of five propositions: all complications to be reported individually as a minimum; all 'serious' (without defining the term serious) complications to be reported as a minimum; outcomes meeting the criteria as core as set out by the GASTROS study to be included; unsure; and other options. The result of this live vote was presented to participants, who were given an opportunity for further discussion ahead of a final vote. The result of the second vote is shown in Appendix S1. Votes were split between the first two options, with the lack of a clear consensus mandating the need for further work in this area. Consequently, all complication-type outcomes were excluded from further discussion.

Non-complication-type outcomes for which there was no consensus to include or exclude in the Delphi survey were then discussed. Results from the subsequent voting are presented in Appendix S1. No further outcomes from this no-consensus group were sufficiently prioritized for addition to the final COS. The final COS is listed in Table 2. Participants agreed that future work on complications, definitions, and when outcomes should be reported should involve both patients and healthcare professionals.

Protocol deviations

The original study protocol described a three-round Delphi survey. Based on emerging evidence at that time²³, several COS developers have demonstrated that consensus can be reached with a two-round survey, which was less resource- and time-intensive. Consequently, the approach was altered and a two-round Delphi adopted survey in the present study.

Discussion

The GASTROS study has developed the first COS for use in surgical trials for gastric cancer. This represents a significant step towards addressing the current challenges related to outcome reporting, synthesis of evidence, and research waste in this field. Outcomes in the set were identified as critically important through an inclusive international consensus process involving patients and healthcare professionals. Further work is required to develop the COS, in particular, finalizing definitions, seeking consensus on how complications should be included, and identifying appropriate outcome measurement instruments. In its present form, the COS will guide trialists to which outcome domains should be reported.

A COS can only achieve its stated aims if it is used by researchers. From the outset of the study, the SMG set out a clear strategy to ensure buy-in by researchers and professional bodies. This resulted in broad international support, and the development of a network that enabled the recruitment of over 1000 participants to the Delphi survey. These participants were well balanced in terms of regional origins and personal or professional experience

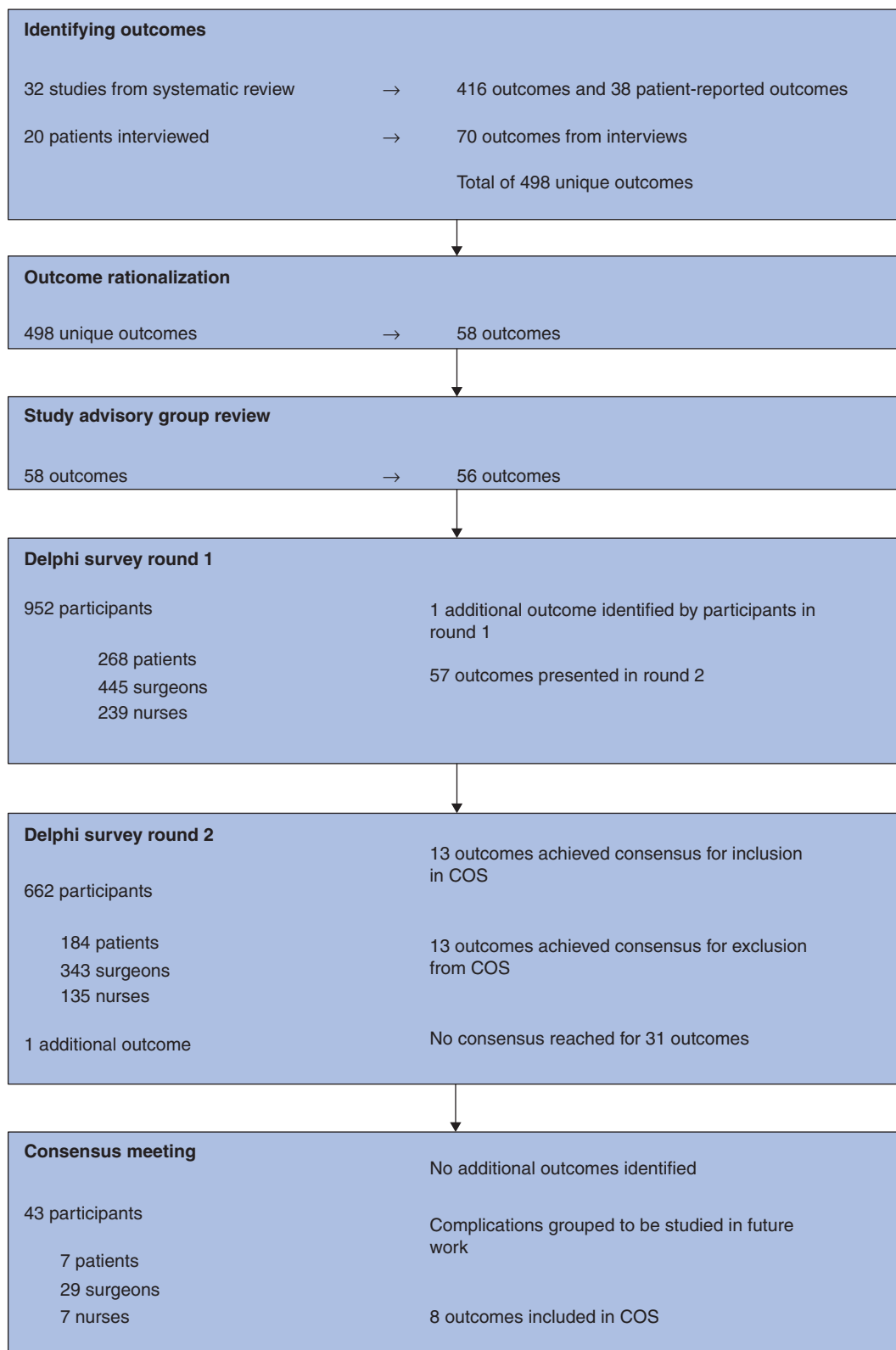


Fig. 2 Results from different stages of the GASTROS study

COS, core outcome set.

of gastric cancer. A strength of the study is that the methodology used is based on consensus guidelines and has been transparently reported in detail at each stage⁵. Researchers can therefore be reassured that the COS has been through a robust

development process and is a valid framework on which to base their research, regardless of where it takes place.

A COS is a minimum set of critically important outcomes. It does not limit trialists in their reporting of other outcomes of

Table 1 Demographic characteristics of participants included in analysis of round 1 and 2 scores

| Stakeholder group | Variable | Completed round 1 only (%) | Completed both rounds (%) | P ^a |
|--------------------------|----------------------------|----------------------------|---------------------------|----------------|
| Patients (n = 268) | All | 84 | 184 | |
| | Age (years) | | | 0.602 |
| | <60 | 38 (45) | 77 (42) | |
| | ≥60 | 46 (55) | 107 (58) | |
| | Sex | | | 0.345 |
| | M | 52 (62) | 101 (55) | |
| | F | 32 (38) | 83 (45) | |
| | Region | | | 0.185 |
| | West | 53 (63) | 113 (61) | |
| | East | 23 (27) | 39 (21) | |
| | Other | 8 (10) | 32 (17) | |
| | Country income | | | 0.792 |
| | HIC | 53 (63) | 113 (61) | |
| | LMIC | 31 (37) | 71 (39) | |
| | Time since surgery (years) | | | 0.656 |
| | <1 | 15 (19) | 30 (17) | |
| | 1–3 | 34 (44) | 68 (39) | |
| >3 | 29 (37) | 75 (43) | | |
| Surgical approach | | | 0.850 | |
| Open | 70 (83) | 145 (78) | | |
| Minimally invasive | 14 (17) | 31 (22) | | |
| Type of gastrectomy | | | 0.503 | |
| Total | 40 (49) | 78 (44) | | |
| Partial | 42 (51) | 98 (56) | | |
| Treatment modality | | | 0.495 | |
| Surgery alone | 28 (34) | 69 (39) | | |
| Multimodal therapy | 54 (66) | 110 (61) | | |
| Surgeons (n = 445) | All | 102 | 343 | |
| | Region | | | 0.001 |
| | West | 33 (32) | 174 (51) | |
| | East | 53 (52) | 109 (32) | |
| | Other | 16 (16) | 60 (17) | |
| | Country income | | | 0.010 |
| | HIC | 45 (44) | 201 (57) | |
| | LMIC | 57 (56) | 142 (43) | |
| | Surgeon experience | | | 0.450 |
| | <50 | 21 (29) | 70 (23) | |
| 50–199 | 20 (27) | 103 (34) | | |
| ≥200 | 32 (44) | 127 (42) | | |
| Nurses (n = 239) | All | 104 | 135 | |
| | Region | | | 0.251 |
| | West | 22 (21) | 40 (30) | |
| | East | 57 (56) | 61 (45) | |
| | Other | 25 (24) | 34 (25) | |
| | Country income | | | 0.064 |
| | HIC | 24 (23) | 46 (34) | |
| LMIC | 80 (77) | 89 (66) | | |
| Nurse experience (years) | | | 0.056 | |
| 0–5 | 59 (57) | 59 (45) | | |
| >5 | 44 (43) | 73 (55) | | |

Values in parentheses are percentages. HIC, high-income country; LMIC, low–middle-income country. ^aχ² test.

interest. Furthermore, it should be noted that some grant-awarding bodies can make their own recommendations regarding which outcomes should be reported as a minimum. An example would be the recommendation to report overall survival, which was not prioritized through the present consensus process²⁴. As such, it is recommended that researchers ensure that additional outcomes selected in surgical trials for gastric cancer adequately reflect the opinions of patients and clinicians in their region, as well as taking into consideration other funding requirements. The authors also recommend that further work is done to examine the international applicability of the COS.

The study was unable to achieve agreement with respect to which complications should be included in the COS. The consensus meeting could not decide whether all complications or only

‘serious’ complications should be reported as a minimum. However, the overwhelming majority voted for one of these options, which will be the focus of future work in this area. Other surgical cancer-related COSs have differed in their recommendations for the reporting of complications. Some have included only a small number of ‘serious’ or ‘core’ complication-related outcomes^{15,25}, whereas others have recommended the reporting of a broader collection of complications²⁶. Based on discussions from the consensus meeting and the lack of agreement among other COS developers, the current recommendation is that all ‘serious’ adverse events should be reported as a minimum until this area is addressed further.

The term ‘serious’ was purposefully not defined at the consensus meeting so as not to remove focus from the discussions.

Table 2 Core outcome set for surgical trials in gastric cancer

| | Core outcome |
|----------------------------------|--|
| Surviving and controlling cancer | Disease-free survival Recurrence of cancer* Disease-specific survival Surgery-related death Completeness of tumour removal |
| Impact of surgery | Overall quality of life Nutritional effects |
| Complications | 'Serious' adverse events† |

*The outcome recurrence of cancer can be incorporated into the composite outcome disease-free survival. It is shown here as it separately reached consensus as a core outcome.†No consensus was reached with respect to which outcomes should be reported as a minimum. 'Serious' adverse events should be reported as a minimum while further work is undertaken in this area.

Others have already attempted to define this; the Gastrectomy Complications Consensus Group (GCCG) is a collaboration of European surgeons who have prioritized a list of 27 clearly defined complications which should be reported as a minimum in research, audits, and registries²⁷. They sought consensus through a Delphi process, although their methodology differs from that of the GASTROS study in that patients and non-European health-care professionals did not participate. Currently, it is the only substantial work available in this field addressing the reporting of complications, and will undoubtedly contribute to, and shape, future work in this field, which is being developed by the first author.

Defining outcomes is an area that deserves further consideration. The present approach was to use plain-language descriptions to define outcomes presented in the Delphi survey and consensus meetings. These were developed with the support of the SAG and an independent patient group. This was necessary to ensure that patients were engaged throughout the study and made translations easier. It is acknowledged that these may not be adequately detailed for use in trials, and more work is required with researchers and patients to address this for the outcomes included in the COS. Substantial work has already been undertaken by the StEP-COMPAC (Standardized Endpoints in Perioperative Medicine—Core Outcome Measures in Perioperative and Anaesthetic Care) group to identify available definitions for outcomes from several systematic reviews^{28–31}. As with the GCCG complication list, this process did not involve patients, which was contrary to the recommendation made by the GASTROS consensus meeting. Standardizing definitions for outcomes included in this COS will form part of stage 3 of the GASTROS study.

Identifying which outcomes to measure is the first step in standardizing outcome reporting in this field. Although many outcomes in the COS are event-type outcomes (such as complications, survival, and recurrence), some are composite outcomes which require the use of an instrument to measure (for example, quality of life and nutritional outcomes). There is currently no standardized approach to measuring these outcomes^{32–34}, and selecting the best tools to measure them requires a robust methodological approach similar to the one employed in this study^{35,36}. This will form the basis of future work.

There are limitations to the present study that require discussion. It could be argued that, given the multimodal nature of treatment for gastric cancer, the COS would be more relevant if it incorporated all therapies including chemotherapy, radiotherapy,

and endotherapy. However, at the time that GASTROS was conceived, there were 24 ongoing surgical trials planning to recruit 11 000 patients for whom non-surgical-related outcomes would not be applicable or relevant. The timing of support from Japanese and Korean collaborators meant that their participation was through the English-language Delphi survey which likely influenced the recruitment of patients from these countries. Nonetheless, an exploratory analysis of the Delphi survey results suggested that patients from Eastern countries did not prioritize outcomes differently from their Western counterparts²². As discussed above, it is recommended that researchers work with patients as partners to determine additional outcomes that are important locally at the trial design phase.

Another consideration relates to the type of stakeholder groups recruited to the study. It was agreed to limit participation to patients, nurses, and surgeons as this represented a balance of a broad spectrum of opinion but ensured that the study's coordination and data analysis were manageable. It should be acknowledged that other groups, such as caregivers, allied health professionals, regulators, policymakers, and grant awarding bodies, will also provide valuable opinion. Inclusion of these groups will be considered in future stages of the GASTROS study and when the COS is reviewed.

The consensus meeting was held in English, limiting participants to English speakers only. Although there was a broad spectrum of international representation from the surgeon group, this was not mirrored by the patient or nurse stakeholders who were primarily UK-based. That said, no further outcomes were added following discussions, supporting the validity of the Delphi process, which recruited widely in terms of regional origin and other demographic characteristics across all three stakeholder groups.

Collaborators

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Data availability statement

P.R.W. and I.A.B. are joint senior authors. The data sets analysed for the present study are available from the corresponding author on reasonable request.

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Supplementary material

Supplementary material is available at *BJS* online.

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