Diagnostic criteria and symptom grading for delayed gastric conduit emptying after esophagectomy for cancer: international expert consensus based on a modified Delphi process

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SUMMARY. Delayed gastric conduit emptying (DGCE) after esophagectomy for cancer is associated with adverse outcomes and troubling symptoms. Widely accepted diagnostic criteria and a symptom grading tool for DGCE are missing. This hampers the interpretation and comparison of studies. A modified Delphi process, using repeated web-based questionnaires, combined with live interim group discussions was conducted by 33 experts within the field, from Europe, North America, and Asia. DGCE was divided into early DGCE if present within 14 days of surgery and late if present later than 14 days after surgery. The final criteria for early DGCE, accepted by 25 of 27 (93%) experts.

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were as follows: > 500 mL diurnal nasogastric tube output measured on the morning of postoperative day 5 or later or > 100% increased gastric tube width on frontal chest x-ray projection together with the presence of an air–fluid level. The final criteria for late DGCE accepted by 89% of the experts were as follows: the patient should have ‘quite a bit’ or ‘very much’ of at least two of the following symptoms; early satiety/fullness, vomiting, nausea, regurgitation or inability to meet caloric need by oral intake and delayed contrast passage on upper gastrointestinal water-soluble contrast radiogram or on timed barium swallow. A symptom grading tool for late DGCE was constructed grading each symptom as: ‘not at all’, ‘a little’, ‘quite a bit’, or ‘very much’, generating 0, 1, 2, or 3 points, respectively. For the five symptoms retained in the diagnostic criteria for late DGCE, the minimum score would be 0, and the maximum score would be 15. The final symptom grading tool for late DGCE was accepted by 27 of 31 (87%) experts. For the first time, diagnostic criteria for early and late DGCE and a symptom grading tool for late DGCE are available, based on an international expert consensus process.

KEY WORDS: consensus, esophagectomy, gastric emptying, malnutrition.

INTRODUCTION

Multimodal treatment including esophagectomy with gastric conduit reconstruction is the treatment of choice for esophageal cancer. Recent improvements in survival after curative treatment put additional focus on short- and long-term morbidity and functional outcome. Delayed gastric conduit emptying (DGCE) is recognized as one of the most important ostectomy problems. DGCE is associated with short-term adverse outcomes including anastomotic leakage, pneumonia, increased ICU, and total hospital stay. DGCE may also lead to nutritional problems and a reduced quality of life.

The pathophysiology of DGCE is not fully understood. Proposed contributing causes are pyloric and gastric dysmotility due to vagal and sympathetic denervation, mobilization of the conduit from a positive pressure compartment to a negative pressure compartment, and disruption of the native antireflux mechanisms. Also other factors, such as conduit size or reconstructive route, relate to variations in clinical presentation. No widely accepted diagnostic criteria and tools to evaluate the presence and severity of DGCE are available. This has led to limitations in the assessment of the current literature on the incidence of DGCE and the effect of preventive and therapeutic measures.

To establish diagnostic criteria and symptom grading for DGCE, we conducted an international expert consensus process, based on modified Delphi methodology. This was combined with live interim group discussions. The Delphi process is a questionnaire-based method that enables experts to express their opinion independently, avoiding the risk of dominant speakers influencing consensus work in open sessions. The Delphi process methodology is well established and has previously been used to achieve expert consensus within various medical fields.

The aims of this expert consensus process were (i) to reach international agreement regarding diagnostic criteria for DGCE after esophagectomy with gastric conduit reconstruction, both in the immediate postoperative phase and in a more long-term perspective and (ii) to reach international agreement regarding a symptom grading tool for DGCE after esophagectomy with gastric conduit reconstruction.

MATERIALS AND METHODS

Modified Delphi expert consensus process

Expert group

This international expert consensus process was a collaboration between experts in Europe, North America, and Asia, using repeated online questionnaires and interim live group discussions. The consortium of 33 experts included a previously assembled collaborative group of surgeons in Europe within the field of minimally invasive esophagectomy, the European Minimally Invasive Osophagectomy Think Tank, supplemented by a number of international leading esophageal surgeons listed in Table 1. A modified Delphi process with live interim group discussions was used to gradually achieve consensus on the exact combination of symptoms, clinical findings, and diagnostic modalities required for the diagnosis of DGCE. The same method was used to reach expert consensus on symptoms and methods used for severity grading of DGCE.

Delphi study

A literature search was performed in April 2017 to scan the published literature for possibly relevant symptoms and diagnostic modalities suggested to represent DGCE. The search was performed in PubMed using the following MESH terms: esophagectomy, gastric emptying, and gastric outlet obstruction; additionally, we complemented the search with the following terms: delayed gastric emptying and gastric conduit dysfunction. The Delphi round 1 questionnaire included multiple choice questions with options graded on a 5-point Likert scale evaluating the experts’ opinion on both frequency of each symptom (symptom present in 0–20%, 21–40%,
Table 1 Participating experts

| Name                  | Country | Institution                                                                 |
|-----------------------|---------|-----------------------------------------------------------------------------|
| Mark I van Berge Henegouwen | The Netherlands | Amsterdam UMC, location AMC, University of Amsterdam, Cancer Center Amsterdam |
| Christiane Bruns     | Germany | University Hospital Cologne                                                  |
| Asif Chaudry          | UK      | The Royal Marsden Cancer Centre, London                                      |
| Edward Cheong         | UK      | Norfolk & Norwich University Hospital                                         |
| Miguel Cuesta         | The Netherlands | Free University Medical Center Amsterdam                                   |
| Gail E. Darling       | Canada  | University Health Network                                                    |
| Suzanne S Gisberz     | The Netherlands | Amsterdam UMC, location AMC, University of Amsterdam, Cancer Center Amsterdam |
| Michael Griffin       | UK      | Royal Victoria Infirmary, Newcastle                                          |
| Christian Gutschow    | Switzerland | University Hospital Zurich                                                   |
| Richard van Hillegersberg | The Netherlands | Johns Hopkins Cancer Center, Houston                                           |
| Wayne Hofstetter      | USA     | Agaplesion Markus Hospital Frankfurt                                        |
| Arnulf Hölscher       | Germany | Keio University Hospital, Tokyo                                               |
| Yuko Kitagawa         | Japan   | Landspathal National University Hospital                                     |
| Magnus Konradsson     | Sweden/Iceland | Erasmus Medical Center, Rotterdam                                          |
| Jan JB van Lanschot   | The Netherlands | University Medical Center Utrecht                                               |
| Lorenzo Ferri         | Canada  | Toronto General Hospital                                                    |
| Donald Low            | USA     | Queen Alexandra Hospital Portsmouth                                          |
| Misha D Luyer         | The Netherlands | St. Mary's Hospital, London                                                  |
| Stuart Mercer         | UK      | Massachusetts General Hospital, Boston                                         |
| Krishna Moorhy        | UK      | Leuven University Hospital                                                   |
| Christopher Morse     | USA     | Catharina Hospital, Eindhoven                                                |
| Philippe Nafteux      | Belgium | Karolinska Institutet, Stockholm                                              |
| Girard A Niuenhuijzen | The Netherlands | Ghent University                                                            |
| Magnus Nilsson        | Sweden  | Radboud University Medical Center                                             |
| Piet Pattyn           | Belgium | Utrecht Medical Center                                                       |
| Camiel Rosman         | The Netherlands | Helsinki University Hospital                                                |
| Jelle P Ruurda        | The Netherlands | Hirslanen Medical Center, Zürich                                           |
| Jari Räsänen          | Finland | University of Cologne                                                      |
| Paul M Schneider      | Switzerland | Oxford University Hospitals                                                  |
| Wolfgang Schröder     | Germany | Leuven University                                                           |
| Bruno Sgromo          | UK      | Erasmus Medical Center, Rotterdam                                            |
| Hans Van Veer         | Belgium |                                                                      |
| Bas PL Wijnhoven      | The Netherlands |                                                                      |

41–60%, 61–80%, or 81–100% of cases of DGCE) and specificity of each symptom (not at all specific, slightly specific, moderately specific, specific, or very specific) for DGCE. For diagnostic modalities, experts were asked to grade the strength of support for the diagnosis of DGCE that each diagnostic modality would provide on a 5-point Likert scale. Opportunities were given in each Delphi round to suggest additional items, and fields for additional comments were available throughout the questionnaires. Repeated Delphi rounds were planned until consensus was achieved on which symptoms and diagnostic modalities should be kept to form diagnostic criteria, based on successive exclusions during the Delphi process. Repeated Delphi rounds were planned until consensus was reached on diagnostic criteria and a symptom grading tool for DGCE. After each Delphi round, a live interim group discussion was planned, until the last Delphi round would test the final product of the consensus process. In the last Delphi round, basic demography of participants was gathered. The Delphi process was performed using an online survey-system (SurveyMonkey, Palo Alto, CA). The complete Delphi survey questionnaires are provided in the Supplementary Appendix 1.

Live interim group discussions
After analysis of the results of each Delphi round, live interim group discussions were planned as a part of the consensus process, until the last Delphi round. The group discussions were planned to provide opportunities to raise safety concerns, discuss practical issues, and provide suggestions for further developments to the diagnostic criteria and symptom severity tool to be considered in following Delphi rounds on the basis of the results of previous rounds.

Statistics and consensus algorithms
Consensus to exclude any symptom from the diagnostic criteria was reached when at least 50% of the experts considered frequency or specificity of the symptom to be in the lower two levels of a 5-level Likert scale. Consensus to exclude any diagnostic modality was reached when at least 50% of the experts considered the strength of the diagnostic modality supporting the diagnosis to be of the two lowest levels of a 5-level Likert scale.

In Delphi round 2, consensus was considered to be achieved if 50% agreed on 1–2 of the peripheral levels of a 5-level Likert’s scale, and in binary
questions or multiple choice questions not compatible with a Likert’s scale, if 70% of experts agreed on one answer. Regarding diagnostic criteria and symptom grading, consensus was considered to be achieved if 80% agreed upon one option.

We classified questions regarding diagnostic criteria and symptom score for DGCE into two domains depending on whether they occurred during (domain 1) or after (domain 2) the first 14 days postesophagectomy. We calculated Cronbach alpha for each of the domains to validate internal consistency.

Cronbach alpha was calculated using Stata 14 (StataCorp. 2015. Stata Statistical Software: Release 14; StataCorp LP, College Station, TX).

RESULTS

The Delphi questionnaires were sent to all 33 experts for completion. Table 1 lists all participating experts, and Table 2 shows an overview of the modified Delphi expert consensus process.

Expert demographics and study overview

The mean (range) number of esophagectomies performed by the responding 27 of 33 experts’ institutions was 81 (30–180) per year. The mean time the experts had been actively engaged in the surgical treatment of patients with esophageal cancer was 18 (6–35) years.

Relevance of time point of DGCE diagnosis

The Delphi round 1 questionnaire addressed whether a proposed early–late DGCE dichotomy should be established. Consensus was reached that DGCE should be classified as early if diagnosed within the first 14 days of surgery and late if diagnosed later.

Early DGCE

Frequency and specificity of symptoms typical for early DGCE

Based on the literature search 17 symptoms were evaluated. In Delphi round 1, 14 of the 17 symptoms for early DGCE were excluded according to the criteria of the consensus algorithm (see Table 3). At the first live interim group discussion held on December 8, 2017, concern was raised that even the symptoms retained after the first Delphi session would be too unspecific and rather represents general postoperative symptoms. Consensus to abort both further attempts to incorporate symptoms in the early DGCE diagnostic criteria and to design a symptom grading scale for early DGCE was reached in Delphi round 2 by 29 of 32 (91%) and 26 of 31 (84%) experts, respectively.

Diagnostic criteria for early DGCE

Six diagnostic modalities were included based on the literature search. After Delphi round 1, all diagnostic modalities (chest x-ray, upper gastrointestinal [GI] water-soluble contrast radiogram, upper GI endoscopy, gastric scintigram, computerized tomography with oral contrast, and timed barium swallow) remained as candidates for further evaluation (Table 4). At the first live interim group discussion after Delphi round 1, a unanimous suggestion regarding early DGCE diagnostic criteria emerged as follows: large amount of nasogastric tube drainage fluid or dilated gastric conduit on frontal chest x-ray and upper GI water-soluble contrast radiogram showing delayed contrast passage to the duodenum.

These diagnostic criteria were subsequently tested in Delphi round 2 and were accepted by 29 of 32 (91%) experts, in effect achieving formal consensus. However, at the second live interim discussion held on September 20, 2018, several participating experts questioned the lack of standardization in the performance and interpretation of upper GI water-soluble contrast radiogram in the diagnosis of early DGCE. In addition, some experts expressed concern regarding risk of aspiration associated with this diagnostic procedure in the immediate postoperative phase. This eventually led to a revised suggestion, excluding upper GI water-soluble contrast radiogram, which was unanimously supported by the live meeting participants. The revised diagnostic criteria for early DGCE were subsequently tested in Delphi round 3. The final early DGCE criteria, accepted by 25 of 27 (93%) experts in Delphi round 3 were as follows: >500-mL diurnal nasogastric tube output measured...
### Table 3  Suggested symptoms of early and late DGCE and results of Delphi round 1

| Suggested symptoms of DGCE | Early DGCE | Late DGCE |
|---------------------------|-----------|-----------|
|                           | Frequency of occurrence considered 40% or lower‡ | Specificity considered none or very low§ | Action after Delphi round 1 | Frequency of occurrence considered 40% or lower‡ | Specificity considered none or very low§ | Action after Delphi round 1 |
| Vomiting                  | 53.6%     | 21.4%     | Excluded | 25.0%     | 17.9%     | FE         |
| Nausea                    | 42.9%     | 53.6%     | Excluded | 32.1%     | 42.9%     | FE         |
| Dysphagia to solids       | 53.6%     | 60.7%     | Excluded | 53.6%     | 57.1%     | Excluded   |
| Dysphagia to liquidized/soft food | 64.3% | 53.6%     | Excluded | 71.4%     | 57.1%     | Excluded   |
| Dysphagia to liquids      | 82.1%     | 53.6%     | Excluded | 78.6%     | 67.9%     | Excluded   |
| Oral intake intolerance†  | 50.0%     | 42.9%     | Excluded | 42.9%     | 28.6%     | FE         |
| Early satiety/fullness    | 10.7%     | 28.6%     | FE       | 10.7%     | 25.0%     | FE         |
| Regurgitation              | 17.9%     | 32.1%     | FE       | 25.0%     | 25.0%     | FE         |
| Heart burn (cervical)     | 60.7%     | 82.1%     | Excluded | 60.7%     | 64.3%     | Excluded   |
| Bloating                  | 53.6%     | 67.9%     | Excluded | 60.7%     | 64.3%     | Excluded   |
| Chest pressure            | 60.7%     | 78.6%     | Excluded | 53.6%     | 67.9%     | Excluded   |
| Pain                      | 82.1%     | 96.4%     | Excluded | 85.7%     | 89.3%     | Excluded   |
| Coughing                  | 35.7%     | 57.1%     | Excluded | 32.1%     | 50.0%     | Excluded   |
| Recurring pneumonia       | 78.6%     | 46.4%     | Excluded | 67.9%     | 42.9%     | Excluded   |
| Loss of appetite          | 39.3%     | 67.9%     | Excluded | 35.7%     | 67.9%     | Excluded   |
| Inability to meet caloric need by oral intake | 17.9% | 57.1% | Excluded | 21.4% | 35.7% | FE |
| Large amount of gastric tube drainage fluid (only suggested in early DGCE) | 39.3% | 14.3% | FE | NA | NA | NA |

†Excluded in Delphi 2 due to redundancy.
‡Percentage of experts considering symptom occurring in 40% of DGCE cases or less in early and late DGCE, respectively.
§Percentage of experts considering symptom specificity very low or none in early and late DGCE, respectively.
FE, further evaluation in the consensus process.

### Table 4  Suggested diagnostic modalities for early and late DGCE and results of Delphi round 1

| Diagnostic modality | Early DGCE | Late DGCE |
|---------------------|------------|-----------|
|                     | Support of diagnosis considered slight or none† | Action after Delphi round 1 | Support of diagnosis considered slight or none† | Action after Delphi round 1 |
| Delayed contrast passage to the duodenum on upper gastrointestinal water-soluble contrast radiogram | 3.6% | FE | 10.7% | FE |
| Wide gastric tube with minimal contrast passage below the pylorus on CT thorax–abdomen with oral contrast | 3.6% | FE | 3.6% | FE |
| Dilated gastric tube on chest x-ray | 3.6% | FE | 25.0% | FE |
| Delayed esophagogastric emptying on timed barium swallow | 10.7% | FE | 10.7% | FE |
| 50% gastric emptying time (T50) > 180 minutes on gastric scintigraphy Marked retention of food on upper gastrointestinal endoscopy despite >4 hours fasting | 21.4% | FE | 17.9% | FE |
| Marked retention of food on upper gastrointestinal endoscopy despite >4 hours fasting | 14.3% | FE | 10.7% | FE |

†Percentage of experts considering the strength of support to the diagnosis slight or none, in early and late DGCE, respectively.

on the morning of postoperative day 5 or later (but within 14 days of surgery) or > 100% increased gastric tube width on frontal chest x-ray projection (in comparison to baseline chest x-ray taken on the day of surgery) together with the presence of an air–fluid level within 14 days of surgery (Fig. 1).

The Cronbach alpha level of internal consistency for domain 1 with 47 items was 0.89.
## Late DGCE

**Frequency and specificity of symptoms of late DGCE**

Based on the literature search, 16 symptoms were included for evaluation. In Delphi round 1, 10 out of the 16 symptoms were excluded according to the criteria of the consensus algorithm (Table 3). At the first live interim group discussion, removal of one of the remaining symptoms, ‘oral intake intolerance’ was suggested due to redundancy. This suggestion was subsequently accepted by 28 of 31 (90%) experts in Delphi round 2.

### Diagnostic criteria for early DGCE†

One of the following criteria should be fulfilled:

- >500 ml diurnal nasogastric tube output measured on the morning of postoperative day five or later (but within 14 days of surgery)
- >100% increased gastric tube width on frontal chest X-ray projection (in comparison to baseline chest X-ray taken on the day of surgery) together with the presence of an air-fluid level

### Diagnostic criteria for late DGCE‡

The patient should have “quite a bit” or “very much” of at least two of the following symptoms:

- Early satiety/fullness
- Vomiting
- Nausea
- Regurgitation
- Inability to meet caloric need by oral intake

Delayed contrast passage on upper GI water soluble contrast radiogram or on timed barium swallow (until precise evaluation criteria are available, relying on the verdict “delayed contrast passage” by an expert radiologist).

### Symptom grading tool for late DGCE

Each of the 5 symptoms of late DGCE, early satiety/fullness, vomiting, nausea, regurgitation or inability to meet caloric need by oral intake should be graded in a four-grade scale in the manner:

- “During the last week, have you had (the symptom; e.g., nausea)?”

| Grade | Symptom Description |
|-------|---------------------|
| 0     | Not at all          |
| 1     | A little            |
| 2     | Quite a bit         |
| 3     | Very much           |

The total symptom score for late DGCE will be the sum of the score for all individual symptoms, ranging between a minimum of 0 points and a maximum of 15 points.

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†Within 14 days of surgery, ‡Later than 14 days after surgery

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Fig. 1 Final Consensus Statement on DGCE.
patient should have a specified number of DGCE symptoms (selected in Delphi round 1) and DGCE on functional imaging. The framework design for diagnostic criteria of late DGCE was accepted by 30 of 32 (94%) experts in Delphi round 2. Opinion was divided in Delphi round 2 regarding the number of symptoms required for the diagnosis of late DGCE. Some 14 of 31 experts considered 2 symptoms appropriate and 14 of 31 three symptoms. The second criterion within the framework design was that diagnosis of DGCE should be supported by functional imaging. The preferred functional imaging was water-soluble contrast swallow for 48.4%, barium swallow for 32.3%, and scintigraphy for 19.3% of participants. At the second live interim group discussion, a first definition of the diagnostic criteria for late DGCE was proposed, unanimously suggesting a minimum number of symptoms of two and additionally a minimum severity of each of those symptoms of 2 points (presence of the symptom graded by the patient as ‘quite a bit’ or ‘very much’) on a 0–3 points scale. Furthermore, the functional imaging was unanimously suggested to include both the option of barium swallow and the option of upper GI water-soluble contrast radiogram. The revised diagnostic criteria for late DGCE were subsequently tested in Delphi round 3. The final late DGCE criteria accepted by 24 of 27 (89%) experts were as follows: the patient should have ‘quite a bit’ or ‘very much’ of at least two of the following symptoms: early satiety/fullness, vomiting, nausea, regurgitation or inability to meet caloric need by oral intake and delayed contrast passage on upper GI water-soluble contrast radiogram or on timed barium swallow (until precise evaluation criteria are available, relying on the verdict ‘delayed contrast passage’ by an expert radiologist) (Fig. 1).

**Symptom grading tool for late DGCE**

The design of a symptom grading tool based on the symptom grading system used in the EORTC Health related Quality of Life Questionnaires was accepted by 26 of 28 (93%) experts in Delphi round 1. Consequently, when reporting symptom severity, the presence and severity of each symptom should be graded as follows: ‘not at all’, ‘a little’, ‘quite a bit’, or ‘very much’, generating 0, 1, 2, or 3 points, respectively. For the five symptoms retained after Delphi round 2 as presented above, the minimum score would be 0, and the maximum score would be 15. The final symptom grading tool for late DGCE presented in Figure 1 was accepted by 27 of 31 (87%) experts in Delphi round 2. A questionnaire for symptom grading of DGCE after esophagectomy is presented in Figure 2.

The Cronbach alpha level of internal consistency for domain 2 with 45 items was 0.85.

**DISCUSSION**

To date, the studies focusing on DGCE have relied on local practice or diagnostic criteria created for the purpose of individual study, making valid comparisons between studies and summaries of results difficult. The need for widely accepted diagnostic criteria has been mentioned in the conclusion of several trials, particularly in studies attempting to compare or summarize the results of multiple studies. In this study, a modified Delphi process was used based on the opinion of an international expert panel, to establish the diagnostic criteria for early and late DGCE, and a symptom severity grading tool for late DGCE.

These results can be used to compare the outcomes of DGCE in various studies determining post-esophagectomy morbidity and quality of life. In addition, the symptom grading tool for late DGCE provides a common symptom grading system.

A strength of this consensus process is the combination of the Delphi method and the live group discussions. The Delphi part allowed all participants to independently express their opinion without any peer pressure or influence of dominant speakers. Participants were prompted to comment specific parts of each Delphi round, and free opportunity for additional comments or proposals was given. The interim group discussions allowed for an open, structured discussion on relevant topics regarding safety, clinical relevance, and feasibility of specific parts of the diagnostic criteria and symptom score. Any suggestion, addition, or revision of former results provided in the interim group discussions was put to test in a subsequent Delphi round. A further strength of this study is the international participation of experts from three continents, indicating that the principal consensus statement on DGCE may be adapted in various countries despite local practice differences.

The live interim group discussions unavoidably carry a risk of strong opinion affecting the consensus process, and approximately ¼ of the expert group were not present at those meetings (Table 2). A majority of the experts did, however, participate in the live interim group discussions, and care was taken that any suggestions formed at those meetings, such as the final statement, would be tested in a subsequent survey. A weakness of this study is that available scientific evidence is limited, and the statement is thus based only on expert opinion. Further studies are mandatory to validate the diagnostic criteria and symptom score against diagnostic modalities. Despite the lack of validation, it may be argued that the diagnostic criteria and symptom score carry at least the same validity as criteria singularly created for the purpose of solitary studies and furthermore carries the strength of a
Please answer these questions by circling the answer below (0-3) that best describes your situation during the past week.

Name: 

Birth date: 

Today’s date: (day, month, year) 

## During the past week: 

| Question                                                                 | Not at all | A little | Quite a bit | Very much |
|-------------------------------------------------------------------------|------------|----------|-------------|-----------|
| 1. Have you felt full up too quickly while having your meal?             | 0          | 1        | 2           | 3         |
| 2. Have you vomited?                                                   | 0          | 1        | 2           | 3         |
| 3. Have you felt nausea?                                                | 0          | 1        | 2           | 3         |
| 4. Have you had acid, bile or food coming up into your throat or mouth? | 0          | 1        | 2           | 3         |
| 5. Have you been unable to eat or drink enough to meet your daily need for energy? | 0 | 1 | 2 | 3 |

Fig. 2 DGCE questionnaire after esophagectomy with gastric conduit reconstruction.

consensus reached within an international group of experts.

This study sought to provide diagnostic criteria for early and late DGCE and a symptom severity score for DGCE regardless of whether a whole stomach or a gastric tube was used as a conduit. To avoid disparity in the consensus process regarding the use of chest x-ray in the early DGCE diagnostic criteria, all items concerning functional radiology in the questionnaires and during interim group discussions were focused on the use of a gastric tube conduit. Whether the diagnostic criteria for early DGCE are less applicable in the presence of a whole stomach conduit remains to be evaluated.

In conclusion, this modified Delphi consensus process provides diagnostic criteria for early and late DGCE and a symptom grading tool supported by a group of international experts. This will be an important tool that can be used for future studies and allows defining a benchmark on DGCE.

**SUPPLEMENTARY DATA**

Supplementary data are available at DOTESO online.

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