Is Textbook Outcome a valuable composite measure for short-term outcomes of gastrointestinal treatments in the Netherlands using hospital information system data? A retrospective cohort study

Nèwel Salet,1,2 Rolf H Bremmer,2 Marc A M T Verhagen,3 Vivian E Ekkelkamp,4 Bettina E Hansen,5,6 Pieter J F de Jonge,6 Rob A de Man6

ABSTRACT

Objective To develop a feasible model for monitoring short-term outcome of clinical care trajectories for hospitals in the Netherlands using data obtained from hospital information systems for identifying hospital variation.

Study design Retrospective analysis of collected data from hospital information systems combined with clinical indicator definitions to define and compare short-term outcomes for three gastrointestinal pathways using the concept of Textbook Outcome.

Setting 62 Dutch hospitals.

Participants 45 848 unique gastrointestinal patients discharged in 2015.

Main outcome measure A broad range of clinical outcomes including length of stay, readmission, and doctor–patient counselling.

Results Patients undergoing endoscopic retrograde cholangiopancreatography (ERCP) for gallstone disease (n=4369), colonoscopy for inflammatory bowel disease (IBD; n=19 330) and colonoscopy for colorectal cancer screening (n=22 149) were submitted to five suitable clinical indicators per treatment. The percentage of all patients who met all five criteria was 54±9.6% (SD) for ERCP treatment. For IBD this was 47±7% of the patients, and for colon cancer screening this number was 85±14%.

Conclusion This study shows that using data obtained from hospital information systems combined with clinical indicator definitions can be used to express short-term outcomes using the concept of Textbook Outcome without any excess registration. This information can provide meaningful insight into the clinical care trajectory on the level of individual patient care. Furthermore, this concept can be applied to many clinical trajectories within gastroenterology and beyond for monitoring and improving the clinical pathway and outcome for patients.

INTRODUCTION

Background

Indicators for measuring healthcare quality can be divided into three main groups: structure indicators, clinical indicators and genuine health outcomes.1–3 Structure indicators focus on infrastructure and the presence of protocols and guidelines. Clinical indicators largely focus on the presence of evidence-based treatment and adverse events like infection or readmission rates directly impacting the individual patient. Finally, genuine health outcomes consist of patient-reported health and quality of life after receiving care. Although genuine health outcomes are considered to be the most valuable indicators, these health outcomes are however mostly unavailable for most diseases. Moreover, collecting data concerning genuine health outcomes requires substantial effort, is time consuming and usually not routinely part of standard care in most hospitals. Although clinical indicators provide inadequate information on long-term outcome, they can provide useful information on the clinical path of individual patients. Monitoring clinical indicators can be used to improve the quality of healthcare1 and bears most value when analysed in a combination of multiple indicators, due to the multidimensional nature of most diseases.5

Strengths and limitations of this study

► The data and analyses were obtained from existing data from hospital information systems.
► The approach used in this study is applicable for evaluating various interventional processes.
► The study does not provide insight into post-discharge outcomes.
Textbook Outcome (TO) is a composite measure of clinical process indicators. TO is realised for patients for whom all desired short-term health indicators are met. This approach enables a simple comprehensive summary of clinical care, and an in-depth analysis to get clinical insight into daily practice, per patient group and indicator, all the way down to the clinical pathway per individual patient. The approach of TO is particularly suited for clinical interventions (surgery, invasive diagnostics) and was previously used in a study performed in the Netherlands in patients undergoing colon resection due to colon cancer. The concept was also used in the form of a questionnaire in which patients reported their considerations in the choice of a hospital, in patients with oesophagogastric cancer in need of surgery, and elective aneurism surgery. However none of these studies used existing data primarily used for reimbursement.

**Objective**

The objective of this study is to develop a model for monitoring short-term outcome of clinical care trajectories for hospitals in the Netherlands using data obtained from hospital information systems. The model is expected to successfully identify hospital variation on short-term outcomes on a large scale in a feasible and reproducible manner. In order to assess the discriminative value of the indicators used, the specificity score per indicator will be calculated.

In order to establish these objectives we further elongate on previous approaches using TO, and apply this means of clinical pathway measurement on a larger scale based on clinical indicators for high-volume treatments. A TO was defined for three different treatments performed by gastroenterologists, consisting of at least five evidence-based indicators as reviewed by an undisclosed panel of physicians.

![Flow chart of included hospitals and corresponding patient trajectories.](Image)

*Hospitals affiliated with LOGEX are included in this study. ERCP, endoscopic retrograde cholangiopancreatography; IBD, inflammatory bowel disease.*
Dutch gastroenterologists. In order to give a valid representation of the care patients received in any included hospital, these indicators should cover as many stages of care as the data allow. This study aims to include indicators covering preprocedural, procedural and postprocedural care. The value of clinical indicators for patients in need of an endoscopic retrograde cholangiopancreatography (ERCP) due to gallstone disease will serve as an example in this study, focusing on the treatment trajectory prior, during and after ERCP. Furthermore, clinical indicators will be applied on two other major gastrointestinal treatments based on the registration of available operational care activities. While analysing a great number of treatments and providing a clear but comprehensive measure of the proportion of patients who have reached a TO, a new approach of assimilating existing data is exerted.

**METHODS**

**Study design**

This study was reported in accordance with the Strengthening the Reporting of Observational Studies in Epidemiology statement for reporting observational studies.10

The potential of TO was assessed by choosing three high-prevalent gastrointestinal diagnoses requiring endoscopic intervention. The first trajectory included patients with the diagnosis of ‘gallstone disease’ who underwent ERCP for stone extraction. The second trajectory included patients who underwent at least one colonoscopy for ‘colorectal cancer screening’. The third trajectory included patients with ‘Inflammatory bowel disease’ (IBD), who also underwent a colonoscopy.

**Setting and data sources**

The data set was retrieved from the benchmark database owned by LOGEX (Amsterdam, The Netherlands). This database contains specific care activities and treatment characteristics registered within the hospital information system from hospitals associated with LOGEX. Each of these data sets is carefully validated in cooperation with hospital information technology specialists, medical specialists and LOGEX. This validation process includes comparison with previous data deliveries (to identify unexpected outliers) and comparison of outpatient contacts, inpatient contact and operations with the electronic patient records of the hospital. The retrieved benchmark database includes a wide variety of information, such as, but not limited to: start and end dates of treatment, doctor–patient contacts, performed endoscopic, radiologic or laboratory diagnostics, surgical intervention, time of admission and days of inpatient stay. These activities per patient combine into care products (so called ‘DBC-DOT Zorgproducten’), comparable with diagnosis-related groups, which are primarily used for structuring and reimbursement of delivered care to healthcare providers. A recent study has shown that administrative data are a valid venue of data, and can be used for quality assessment of healthcare in cardiac patients.11 In 2015, the total number of hospitals in the Netherlands was 83, of which 62 were included in this study (75%); academic hospitals were excluded as will be discussed in the Discussion section, as well as four hospitals without a gastrointestinal department. An overview of the selection of included hospitals per treatment is illustrated in figure 1.

**Study size and participants**

All patients (n=45 848) with one of the three defined trajectories discharged between 1 January 2015 and 31 December 2015 were evaluated. By definition, all patients were discharged within 42 days after intervention and inpatient stay (related to reimbursement regulations). All analysed patient trajectories were required to have at least one ‘core’ care intervention registered for their diagnosis; for gallstone disease this was the ERCP activity, and for IBD and colon cancer screening this core activity was the colonoscopy. For example, to be included in the analysis of ‘gallstone disease’, patients were required to fulfill the following inclusion criteria: (1) patients were diagnosed with gallstone disease, (2) were treated by a gastroenterologist and (3) underwent at least one ERCP for stone extraction (being the core activity), and (4) patient was discharged in 2015; there were no

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Overview of the criteria per treatment structured by preprocedural, procedural and postprocedural indicators</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Diagnosis</strong></td>
<td><strong>Treatment/core activity</strong></td>
</tr>
<tr>
<td>Choledocholithiasis (Gallstone disease)</td>
<td>ERCP</td>
</tr>
<tr>
<td>Colorectal cancer screening</td>
<td>Colonoscopy</td>
</tr>
<tr>
<td>Morbus Crohn and colitis ulcerosa (IBD)</td>
<td>Colonoscopy</td>
</tr>
</tbody>
</table>

ER, emergency room; ERCP, endoscopic retrograde cholangiopancreatography; IBD, inflammatory bowel disease.
additional exclusion criteria for patients. An overview of the selection of included patients per treatment is illustrated in figure 1; all diagnoses and used indicators are shown in table 1.

Variables
Extensive literature search was conducted prior to defining TO indicators. The indicators listed in table 1 show the selected criteria to assess the clinical outcomes of ERCP, colorectal cancer screening and IBD. The selection choice of the clinical indicators is described in the online supplementary appendix in more detail.

Outcome definition
This study’s primary goal was to determine the variation among hospitals’ treatment score. For each patient treated, we determined if the treatment was considered to conform with TO—a binary outcome score (1/0). A patient was considered TO when all five indicators were met; if one or more of the five indicators were not met, the treatment was not considered TO. The selected set of indicators was applied to all patients, regardless of their background or medical complexity. The hospital score per treatment consists of every individual patient accumulated into a total score illustrated in a percentage of patients who have reached TO in the corresponding hospital: the TO score (%) is the quotient of total number of patients treated while fulfilling all five indicators (numerator) and the total number of treated patients in that hospital for that intervention (denominator). The indicators in the TO scope range from the first outpatient contact with the gastroenterologist up to the last registered care activity in the care trajectory, usually being a consult to check-up on the patient after the treatment to conclude the care cycle. If no new care activity is registered for a patient related to this intervention after patient discharge, the care trajectory closes automatically.

Statistical methods
The clinical indicators were assessed for each patient and the product of all clinical indicators resulted in the number and proportion of patients for whom all desired outcomes were realised and thereby a ‘Textbook Outcome’ was achieved. Per treatment and for each hospital, the proportion of patients with a ‘Textbook Outcome’ was calculated.

Figure 2 Distribution of the scores per indicator shown for 53 hospitals: (A) distribution of scores on doctor–patient contact prior to ERCP, (B) no second ERCP, (C) inpatient stay, (D) no CT scan after ERCP, (E) no readmission within 30 days, and (F) product of all criteria, defined as Textbook Outcome. ERCP, endoscopic retrograde cholangiopancreatography.
For colon cancer screening with a total of 22, 45.0% were male; the average age was 48±17 years (SD).

54% of 4369 unique patient trajectories have met all five criteria. ERCP, endoscopic retrograde cholangiopancreatography; IBD, inflammatory bowel disease; TO, Textbook Outcome.

To assess the impact of clinical indicators where the total TO was not met (TO=0), the specificity of each indicator was determined. The mean specificities across hospitals were depicted along with the percentage of TO that was not met (score=0) in order to provide increasing discriminative value of each singular indicator.

Second, a pairwise comparison between TO score on hospital level and score per indicator was performed per treatment to assess the relationship between reported score per hospital on individual indicators and total TO score. The relation is expressed in Pearson’s correlation. Additionally, pairwise comparison between clinical indicators on hospital level was assessed. Pearson’s correlation coefficient ≥0.7 was considered a strong correlation. The variation in score among hospitals is displayed by the SD. Statistical analyses were performed in Excel and SPSS V.25.

RESULTS
Descriptive data
In total, 62 of the LOGEX-affiliated hospitals were included in this study (figure 1). For the ERCP trajectory a total number of 4369 ERCP patients treated in 51 hospitals were included, of which 41.5% were male; and the average age was 66±18 years (SD). For the IBD colonoscopy subgroup a total of 19 330 patients were included, 45.0% were male; the average age was 48±17 years (SD). For colon cancer screening with a total of 22 149 patients, 60.4% were male; the average age was 67±4 years (SD).

Outcome data and main result
The average TO score (score=1) for ERCP due to gallstone disease was 54%, with an SD of 9%. Accordingly, 54% of 4369 unique patient trajectories have met all five criteria: doctor–patient contact prior to ERCP, not more than one ERCP, inpatient stay equal or shorter than 7 days, no CT scan after ERCP and no readmission within 30 days. Individual scores per indicator are illustrated in figure 2A–F; average score per indicator ranged from 96% (no readmission within 30 days after ERCP) to 79% (length of stay does not exceed 7 days).

Each indicator was assessed in closer detail with regard to specificity in order to assess discriminative value. Figure 3A–C illustrates the specificity and variance on the TO score and each individual indicator. For patients who underwent an ERCP, indicator 5 (no readmission within 30 days) shows the lowest variance, and therefore is influencing the hospital’s total TO score the least. Indicator 3 (inpatient stay <7 days), however, shows the largest variance.

Figure 4A–F illustrates the association between the total TO score on hospital level and the hospital score per individual indicator. The calculated Pearson’s correlation coefficient depicts the correlation between the score per individual indicator on hospital level and the total TO score. The statistical correlation for the score on doctor–patient contact with the total TO score proved weak-moderate positive (r=0.38). The correlation for scores on readmission (r=0.56) and no CT scan (r=0.62) were higher than with patient contact, being considered as a moderate positive linear relationship. The correlation for scores on the indicators’ inpatient stay ≤7 days (r=0.74) and no second ERCP (r=0.80) proved strong positive. Pairwise comparison of the two indicators with highest correlation with total TO score on hospital level gives a weak correlation of r=0.55 (figure 4F).

The average TO score for IBD was 47%, with an SD of 7%. Accordingly, 47% of 19 330 unique patient trajectories in 62 hospitals met all five criteria: time between first consult and colonoscopy does not exceed 56 days, the amount of colonoscopies is not more than one, inpatient stay equal or shorter than 3 days, no emergency room (ER) admission after colonoscopy and doctor–patient counselling afterwards. Individual scores per indicator are shown...
Figure 4  Correlation between the total Textbook Outcome score and the individual indicators (A) doctor–patient contact prior to ERCP, (B) no second ERCP, (C) inpatient stay, (D) no CT scan after ERCP, (E) no admission within 30 days, and (F) the relation between hospital scores on no second ERCP and inpatient stay ≤7 days. ERCP, endoscopic retrograde cholangiopancreatography.

In figure 5A–F, average score per indicator ranged from 68% (time between first consult and colonoscopy does not exceed 56 days) to 97% (no second colonoscopy in clinical pathway). Indicator R values ranged from a weak r=0.02 (no CT scan after colonoscopy) to a moderate r=0.57 (doctor–patient contact prior to colonoscopy) positive correlation.

The average TO score for colorectal cancer screening was 85%, with an SD of 14%. Accordingly, 85% of 22,149 unique patient trajectories in 53 hospitals met all five criteria: doctor–patient contact prior to colonoscopy, no CT colon and no laboratory tests indicating complications, no inpatient admission after colonoscopy and no ER admission afterwards. Individual scores per indicator are shown in figure 6A–F average score per indicator range from 93% (doctor–patient contact prior to colonoscopy) to 100% (no ER admission after colonoscopy). Again, indicator R values ranged from a weak r=0.17 (no second colonoscopy) to a moderate r=0.68 (maximum 56 days waiting period) positive correlation.
The total number of 4369 ERCPs performed for gallstones included in this research covers 87% of the total of 5000 reimbursed ERCPs performed in the Netherlands in 2015*. With 19 330 patients, the total amount of colonoscopies for IBD covers 75% of the total in the Netherlands, while the 22 149 patients for screening colon cancer IBD cover 76% of the Netherlands in 2015.12

**DISCUSSION**

**Key results**

With the use of TO, departments and physicians will be able to evaluate and compare their clinical outcomes with their peers throughout the entire country. Reporting the composite measure of TO shows added value with regard to points of interest for the total clinical pathway. The composition of TO adds most value when chosen indicators do not overlap and add discriminative value, as is depicted in figure 3. With this model, a representable benchmark can be compiled for meaningful comparison between medical centres to monitor improvement over the years. Pinpointing underperforming segments of clinical care in comparison to their peers is among the possibilities.

To the same extent, it is possible as well to identify ‘best in class’ departments who might serve as an example for horizontal improvement. TO scores can be cross-referenced against produced volume in order to analyse the influence of volume per hospital on the score in terms of clinical outcomes. In this study, we show that available and existing registration data for declaration purposes can be used for monitoring and evaluation of clinical pathways in high-prevalent treatments. While this study does not investigate a relation between volume per hospital and total score in the investigated diagnosis, these results can be integrated in future studies on volume quota per treatment.13 For improving local TO scores, the Pearson’s
correlation coefficient can assist in determining which indicator is most dominant for defining the total TO score; for ERCP this would be focusing on reducing reintervention (figure 4).

**Limitations**

While facing a patient population with an above average medical complexity, it is more likely to result in longer overall inpatient care and/or a higher complication rate. Comparing hospital scores as in this study assumes comparable medical complexity among the analysed hospitals. Future studies have to investigate to what extent medical complexity and comorbidity (Charlson score) variation will influence TO scores. The possibility that patients receive care for the same condition (reintervention) in a different hospital cannot be ruled out. However, in the experience of the doctors who were involved in development of the indicators, the vast majority of these reinterventions take place in the same hospital. The current study does not include academic centres as there is an insufficient amount of academic hospitals in the database under study. Therefore, we are unable to compare the results of these hospitals with their peers or differentiate in scores between academic centres and non-academic centres.

While this study is based on indicator scores on patient level, we emphasise that TO focuses on clinical indicators and does not take patient-related outcome measurement (PROM) or patient-related experience measurement into account. Combining short and long-term outcomes is an interesting next step; however, studies show that consistently collecting patient-reported outcomes (PROMs) faces barriers, with the main issue being the technological limits of integrating an electronic health record on a platform that collects PROMs to rapidly analyse data. Hesitant healthcare providers may even be the largest operational barrier,
with the large amount of time-consuming tasks already being part of their daily routine.\textsuperscript{15} When taking these factors into consideration, the advantages of analysing clinical indicators over PROMs are evident. The suggested combination of indicators per trajectory is shown valuable on hand, but further research is necessary to evaluate the impact of patient characteristics including age, gender and comorbidity. By adding indicators concerning case mix, an even more proficient way of insight can be provided for physicians. We would like to stress that the results do not imply that patients who do not meet all indicators have been treated incorrectly. Certain medical complexity can be a valid reason to divert from TO or any other guidance protocols, if doing so benefits the individual patient. TO’s potential lies in identifying and interpreting significant differences on a group level, rather than advocating indicator-driven clinical decision-making.

Focusing on improving average score on TO will optimise patient care, and probably reduce healthcare costs.\textsuperscript{16} Cost-effectiveness of healthcare is an important debate in both the Dutch and worldwide healthcare.\textsuperscript{17,18} Unchecked expenses are to be increasing significantly in the upcoming years on the demand side due to the ageing of the population. On the supply side, new expensive medical technology and medication to treat the clinically ill patient, for example, with a malignancy, will be available. Although these developments are widely encouraged, it also faces economic and operational challenges. The healthcare sector can aggress these challenges when using advanced data analytics as portrayed before in proper-driven clinical decision-making.

The relative importance of quality of care information when choosing a hospital to improve healthcare can still be pursued.

CONCLUSION

This study shows that applying TO to existing data provides valuable insight into variance of daily clinical practice on a large scale, without additional time-consuming registration. This method of TO based on hospital information system data can be applied to many clinical trajectories for monitoring and improving the clinical pathway and outcome for patients.

REFERENCES


Author affiliations

1VU University Medical Center Amsterdam, Amsterdam, The Netherlands
2LOGEX, Amsterdam, The Netherlands
3Department of Gastroenterology and Hepatology, Diakonessenhuis Utrecht, Utrecht, The Netherlands
4Department of Gastroenterology and Hepatology, Reinier de Graaf Hospital, Delft, The Netherlands
5Institute of Health Policy Management and Evaluation, University of Toronto, Toronto, Ontario, Canada
6Department of Gastroenterology and Hepatology, Erasmus MC, University Medical Center Rotterdam, Rotterdam, The Netherlands

Contributors NS and RHB collected the data. NS, RHB, MV and RoM drafted the manuscript and contributed to all other quality aspects of the study. BH was involved in the data analyses. PJdJ and VE performed critical revision of the manuscript. All authors read and approved the final manuscript.

Funding This research received no specific grant from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests RHB has currently and NS had previously a relevant connection to LOGEX (Amsterdam, The Netherlands) as employees. LOGEX offers healthcare analytics to medical specialists. MV, RoM, PJdJ, BH and VE have no relevant connection to LOGEX.

Patient consent Not required.

Ethics approval No ethical approval was required in this study due to patient anonymity in the database.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The study brought together existing data obtained upon request and subject to license restrictions from a number of different sources. Due to the (commercially, politically, ethically) sensitive nature of the research, no source consented their data being retained or shared.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, provided the original work is properly cited and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/

© Article author(s) (or their employer(s) unless otherwise stated in the text of the article) 2018. All rights reserved. No commercial use is permitted unless otherwise expressly granted.


