Evaluation of indications for and outcomes of elective surgery

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Abstract

Background: Wide small-area variations in the rates of elective surgical procedures and lack of systematic outcome measurement have raised questions about the appropriateness of such surgery. Our objective was to determine the feasibility of routine evaluation of indications for and outcomes of elective surgery.

Methods: Participants consisted of 138 surgeons and 5313 patients who underwent 1 or more of 6 specific surgical procedures (for a total of 6274 operations). Surgical indications were evaluated according to published guidelines. Patients’ self-reported health-related quality of life (HRQOL) before and at appropriate intervals after surgery was measured with standard, validated generic and disease-specific instruments. Patient-specific results were routinely sent to the surgeons, from whom feedback was requested.

Results: Surgeons provided information on the indications for surgery for 44% to 95% of the 6 procedures, and the indications matched the guidelines in 73% to 99% of cases. Completed HRQOL questionnaires were returned by 58% of the patients. Postoperative HRQOL scores were markedly improved in most patients, but in 2% to 26% of the various procedures, there was either no change or a deterioration in HRQOL. In most of the procedure groups a small proportion of patients had relatively minor symptoms and disability preoperatively, but in the cataract surgery group this proportion was large. Opinion among the participating surgeons was divided as to the potential value of this method of evaluation. The cost of the outcome evaluation program was about $12/patient.

Interpretation: Evaluation of indications for and outcomes of elective surgery could be implemented systematically at reasonable cost and could be included in an accountability framework for health services. Most surgeons were not enthusiastic about this kind of evaluation.

The appropriateness of indications for elective surgery has stimulated much debate. The large variations in age- and sex-adjusted surgical rates that are reported whenever the issue is examined raise questions about the adequacy of access in some areas and the possibility of inappropriate surgery in others, especially for procedures known to have highly subjective indications.

It is remarkable that in reports of surgical results so little attention has been paid to measuring the outcome for which most elective surgical procedures were designed, namely health-related quality of life (HRQOL). There have been sporadic reports of HRQOL results in a research context, but there has been no previous attempt to introduce systematic HRQOL measurement as a routine component of the evaluation of elective surgery.

Because indications for and outcomes of surgery are becoming important measures of quality, we undertook this project to assess the feasibility and potential value to providers and decision-makers in the health care system of measuring each of them routinely.

Methods

In Vancouver between November 1999 and September 2000, 5313 consecutive patients were booked for the following 6 high-volume elective surgical procedures: cataract replacement, cholecystectomy, hysterectomy, lumbar diskeectomy, prostatectomy and total hip re-
placement. Patients with cancer and those who underwent arthroplastic revision were excluded. All acute care hospitals in the region participated, namely Vancouver Hospital and Health Sciences Centre, Providence Health Care (consisting of St. Paul’s Hospital, St. Vincent’s Hospital and Mount St. Joseph’s Hospital) and Richmond Hospital. The specialty distribution of the 138 participating surgeons is displayed in Table 1. The project steering committee consisted of surgeons and administrators representing the BC Ministry of Health, the Vancouver/Richmond Health Board and the 6 surgical divisions within the 5 hospitals.

For each patient booked for surgery, the surgeon or a designate was asked to complete a clinical indications form based on a set of proprietary guidelines developed by InterQual (a division of McKesson Corporation, San Francisco), known as Indications for Surgery and Procedures. The information gathered from the clinical indications form was reviewed by one of the project nurses to determine if the case profile matched the guidelines. A sample of 900 hospital charts was reviewed to determine whether the information necessary to review the surgical indications could have been obtained from the hospital record rather than directly from the surgeon.

For the outcomes part of the project, we used the SF-36 generic HRQOL questionnaire and an appropriate disease-specific HRQOL questionnaire for each procedure (see below for more details). The questionnaires were formatted to allow optical scanning of the responses. Questionnaires were sent to all patients before surgery (except those in the hysterectomy group [the Menorrhagia Outcomes Questionnaire was designed for single postoperative application only]) and at appropriate postoperative intervals ranging from 3 months for cataract to 1 year for lumbar disectomy and total hip replacement. A custom-built computer program was used to track the approximately 13 000 patient surveys, flag the date for the postoperative mailing and scan the responses into a database. The per-patient cost of the outcomes measurement program was determined. The self-reported outcomes for each patient were sent to the relevant surgeon as soon as the postoperative surveys were received, and surgeons were asked to provide anonymous feedback about the value of this process.

The SF-36 was chosen as the generic HRQOL instrument for all procedures. The following disease-specific instruments were also used: the Visual Function Assessment (based on the VF-14 index) for cataract extraction, the Gastrointestinal Quality of Life Index for lumbar disectomy and total hip replacement. A custom-built computer program was used to track the approximately 13 000 patient surveys, flag the date for the postoperative mailing and scan the responses into a database. The per-patient cost of the outcomes measurement program was determined. The self-reported outcomes for each patient were sent to the relevant surgeon as soon as the postoperative surveys were received, and surgeons were asked to provide anonymous feedback about the value of this process.

Table 1: Number of surgeons and response rate to request for clinical information

<table>
<thead>
<tr>
<th>Procedure</th>
<th>No. of surgeons</th>
<th>Total no. of procedures</th>
<th>No. (and %) of procedures with response</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cataract replacement</td>
<td>31</td>
<td>4239</td>
<td>2840 (67)</td>
</tr>
<tr>
<td>Cholecystectomy</td>
<td>27</td>
<td>673</td>
<td>477 (71)</td>
</tr>
<tr>
<td>Hysterectomy</td>
<td>40</td>
<td>488</td>
<td>239 (49)</td>
</tr>
<tr>
<td>Lumbar disectomy</td>
<td>11</td>
<td>122</td>
<td>115 (94)</td>
</tr>
<tr>
<td>Prostatectomy</td>
<td>13</td>
<td>264</td>
<td>195 (74)</td>
</tr>
<tr>
<td>Total hip replacement</td>
<td>16</td>
<td>488</td>
<td>214 (44)</td>
</tr>
<tr>
<td>Total</td>
<td>138</td>
<td>6274</td>
<td>4080 (65)</td>
</tr>
</tbody>
</table>

North American Spine Society lumbar spine outcome assessment instrument for lumbar disk surgery, the American Urological Association symptom index for benign prostatic hyperplasia for prostatectomy, and the Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) for total hip replacement. Because of the culturally diverse population in Vancouver, we ensured that staff were available to communicate with patients who spoke English, Mandarin, Cantonese or Punjabi. Chinese-language versions of the SF-36 and the Visual Function Assessment were also available.

The project raised controversy among the members of the steering committee about the distinction between clinical research and quality management, but the Behavioural Research Ethics Board of the University of British Columbia determined that patient consent was not required, because these activities were fundamentally related to the management and reporting of quality of care.

Results

Only one department in one of the hospitals declined to participate in the study. The response rates to the requests for clinical information varied widely by surgeon, by procedure and by institution. The response rate for individual surgeons ranged from 0% to 100% of procedures. No obvious patterns of response in terms of type of procedure or volume of practice were evident. The overall collection rate of clinical indication forms varied from 44% to 94%, depending on the procedure (Table 1), with the lowest response rates for hysterectomy and total hip replacement and the highest response rate for lumbar disectomy. For patients for whom a clinical indications form was submitted by the surgeon, the indications for surgery matched the guidelines for 73% to 99% of cases, depending on the procedure (Table 2).

The response rate for the HRQOL survey packages mailed directly to patients (and reminder postcards and telephone calls where necessary) ranged from 52% to 77%, depending on the procedure. In the returned surveys, 95% of the questions were answered.

For each procedure, information was compiled on changes in all 8 domains of the SF-36 score and in the disease-specific instrument scores, as well as the distribution of preoperative (except for hysterectomy) and postoperative symptom and disability scores (Table 2). The complete data sets and graphs of results for all 6 procedures are available online.

The most striking improvements in HRQOL were seen after lumbar disk surgery and total hip replacement; for these procedures, very large positive changes occurred in the disease-specific instruments and in almost all of the SF-36 domains (Table 2). The disease-specific instruments showed that most patients in all procedure groups experienced benefit from surgery, but for a certain proportion (ranging from 2% for total hip replacement to 26% for cataract replacement) HRQOL scores after the operation were worse than before. From the large data sets available for each procedure 2 illustrative examples were chosen for presentation here: cataract replacement and total hip replacement.
Of the patients booked for cataract surgery, 32% scored higher than 90 points on the 100-point Visual Function Assessment before surgery (Fig. 1). The mean scores before and after surgery for all patients were 79 (standard deviation [SD] 17) and 88 (SD 15) respectively. Seventy percent of the patients who underwent cataract surgery experienced improvement, as recorded by the Visual Function Assessment, but for 27% the score was the same or worse after the procedure (Fig. 2). As expected (because of the general nature of the questions), there was no change in any of the domains of the generic HRQOL instrument, the SF-36.

For the patients who underwent total hip replacement, there was a large mean positive change in HRQOL as measured by both the SF-36 and the WOMAC scales (Table 2). For more than 90% of patients, the WOMAC scale indicated improvement after the procedure, whereas only 2% of patients reported deterioration (Fig. 3).

The staffing cost for producing, tracking and scanning the surveys for one patient was estimated as $2.69 on the basis of a workload estimation model. Supply costs including postage (mail-out and stamped self-addressed envelope) were $2.75 per survey, and the cost of producing the surgeon’s report was approximately $0.50. Therefore, the total costs were approximately $6 per outcome survey or $12 for a preoperative and postoperative protocol. This estimate does not include the costs of nurses identifying patients, which could be eliminated by implementing an automated procedure in the booking offices, or the costs of surgeons’ time, project management, data analysis and interpretation.

Just over half (21 or 53%) of the 40 surgeons who responded to our request for feedback had positive comments about the project and the information it produced, but the remainder felt it was of little value and indicated that they did not wish to receive such information on their patients in the future.

Interpretation

These results suggest that it is feasible to carry out a guideline-based review of the indications for elective surgery and to measure HRQOL outcomes routinely at reasonable cost.

Surgical indications

Given that the project involved 138 surgeons from 6 disciplines in 5 hospitals, it is unsurprising that the level of cooperation was variable, with data on indications reported for 0% to 100% of procedures by individual surgeon, and one division in one of the hospitals declining to participate at all. It is difficult to interpret the data on the rate of compliance with the guidelines (73% to 99%) for several reasons: not all of the surgeons agreed with the proprietary guidelines used, secondary review of individual cases by a physician was not done systematically, and information was incomplete or missing for more than one-third of patients.

### Table 2: Changes in health-related quality of life after elective surgery

<table>
<thead>
<tr>
<th>Procedure</th>
<th>No. of procedures</th>
<th>Mean age (and SD), yr</th>
<th>% matching guidelines*</th>
<th>Mean change (and SD) in SF-36 component score, %</th>
<th>Mean change (and SD) in disease-specific score, %</th>
<th>% of patients with positive change ‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cataract replacement</td>
<td>423</td>
<td>73 (10)</td>
<td>87</td>
<td>PF: -1 (10)</td>
<td>RP: -1 (10)</td>
<td>V: -1 (10)</td>
</tr>
<tr>
<td>Cholecystectomy</td>
<td>67</td>
<td>50 (15)</td>
<td>99</td>
<td>G: +18 (10)</td>
<td>H: +12 (10)</td>
<td>P: +18 (10)</td>
</tr>
<tr>
<td>Hysterectomy</td>
<td>488</td>
<td>47 (9)</td>
<td>97</td>
<td>L: +36 (10)</td>
<td>N: +14 (10)</td>
<td>O: +21 (10)</td>
</tr>
<tr>
<td>Lumbar discectomy</td>
<td>122</td>
<td>44 (9)</td>
<td>97</td>
<td>M: +36 (10)</td>
<td>I: +14 (10)</td>
<td>N: +21 (10)</td>
</tr>
<tr>
<td>Prostatectomy</td>
<td>264</td>
<td>72 (9)</td>
<td>73</td>
<td>T: +36 (10)</td>
<td>R: +14 (10)</td>
<td>S: +21 (10)</td>
</tr>
<tr>
<td>Total hip replacement</td>
<td>488</td>
<td>64 (14)</td>
<td>93</td>
<td>SF: +36 (10)</td>
<td>V: +21 (10)</td>
<td>W: +21 (10)</td>
</tr>
</tbody>
</table>

*Percentage of procedures (for patients for whom clinical information was submitted) for which indications matched the InterQual guidelines on indications for surgery.
†Mean postoperative change in score for each of the 8 domains in the SF-36 questionnaire: PF = physical functioning, RP = role physical, BP = bodily pain, GH = general health, V = vitality, SF = social functioning, RE = role emotional, MH = mental health.
‡Proportion of procedures for which the patient’s disease-specific score improved by any amount after surgery.
enrolled, which suggests the possibility of selection bias by the surgeons. However, the data suggest that local practices often vary from the guidelines. For cholecystectomy, lumbar discectomy and total hip replacement, the rates of criteria match were consistently high.

Our review of hospital records demonstrated that these records are not adequate sources of information on indications for surgery. For systematic implementation of a review of the indications for surgery, the necessary clinical information would have to be collected from the physicians’ offices. Most of the project’s personnel costs arose from time spent in hospital surgical booking offices to identify patients for the study and in attempts to obtain the necessary information from surgeons’ offices. As with an outcomes management system, automation of the surgical booking process would be necessary for successful implementation on a large scale.

**Patient-reported outcomes**

The preoperative HRQOL information indicated that patients booked for these 6 elective surgical procedures had a wide range of symptoms and disability. There was evidence that the more pain and disability experienced before operation, the more likely that there would be benefit from surgery. For most of the procedures, the proportion of patients reporting minimal symptoms and disability preoperatively was small, but of those scheduled for cataract surgery 32%, 15% and 4% had a preoperative visual function score of greater than 90, greater than 95 and 100 respectively (on a scale of 100). These findings suggest that the threshold indications for cataract surgery are now very low. The mean preoperative Visual Function Assessment score of 79 is comparable to the range of 64 to 77 reported in the literature.

The disease-specific instruments showed substantial improvement in patients’ self-reported HRQOL after surgery for all of the procedures, but in a certain proportion the postoperative score was unchanged or worse. The wide range in the proportion with poorer outcomes (2% for total hip replacement to 26% for cataract extraction) may reflect differences in surgical complication rates, severity of disease or appropriateness of the indications for surgery in the first place. For the generic HRQOL instrument, the changes were strongly positive after lumbar discectomy and total hip replacement, moderately positive for cholecystectomy and hysterectomy, minimal for prostatectomy and absent for cataract surgery; these changes might be expected, given the likely overall health effects of the diseases studied.

We were disappointed by the lack of interest expressed by almost half of the surgeons in receiving patients’ self-reported outcome information. This lack of interest suggests...
Implications for health services and policy

The appropriateness and outcomes of health care interventions are important quality-of-care issues that should be considered in the accountability framework for health services now under consideration by the provinces. The results of this study suggest that clinicians, managers and trustees must all be involved if indications and outcomes are to be evaluated. For systematic implementation of such evaluation, it would be effective and less costly to target specific areas in a structured sampling program, but strong incentives for participation by surgeons would be necessary.

The wide range of severity of symptoms and disability for which elective surgery was recommended raises questions about the appropriateness of some procedures. With systematic implementation of an outcomes assessment program and better understanding of the validity of patients’ self-reported outcomes, surgeons and referring physicians may eventually use such information to investigate the reasons for poor outcomes in individual patients and thus refine the surgical selection process. Since all surgery carries risk, the potential benefits and harms of a procedure must be examined for patients with non-life-threatening conditions causing minimal symptoms. In elective surgery with high demand and limited resources, it seems reasonable to ask surgeons to document their reasons for operating and patients to assess their condition before and after surgery. Furthermore, making such information available to surgeons and referring physicians could improve the quality of care and clinical decision-making.

A detailed report of this project, including all the data, is available in PDF format at www.resio.org.

Competing interests: None declared.

Contributors: Charles J. Wright was the principal investigator and mainly responsible for the design of the protocol. He submitted the application for funding, chaired the steering committee and was involved at all stages of the project. G. Keith Chambers contributed to the design of the project, sat on the steering committee throughout the project’s execution, and played a major role in the data analysis and interpretation. Yovel Robens-Paradies was the project director with the major responsibility for all aspects of the running of the project.

Acknowledgements: We offer our warm appreciation to all of the members of the steering committee (Dr. Ron Calderisi, Associate Head of Surgery, Mount Saint Joseph Hospital and St. Vincent’s Hospital, Providence Health Care, Vancouver; Dr. Keith Chambers, Assistant Director, Centre for Clinical Epidemiology, Vancouver Hospital and Health Sciences Centre, Vancouver; Mark Chase, Vice President, Planning and Evaluation, Vancouver/Richmond Health Board, Vancouver; Sue Emmons, Director, QUEST, Vancouver Hospital and Health Sciences Centre, Vancouver; Dr. Charles Fisher, Clinical Assistant Professor, Vancouver General Hospital, University of British Columbia, Vancouver; Dr. Ken Hughes, Head, Division of Orthopaedic Surgery, The Richmond Hospital, Richmond, BC; Dr. David Manson, Head, Department of Surgery, St. Paul’s Hospital, University of British Columbia, Vancouver; Dr. David Rollins, Clinical Professor, Department of Ophthalmology, Medical Director, Eye Centre Operational Rooms, and Head, Division of Comprehensive Ophthalmology, Department of Ophthalmology, University of British Columbia, Vancouver Hospital and Health Sciences Centre, Vancouver; Dr. Barry Sanders, Clinical Professor, Vancouver Hospital and Health Sciences Centre and BC Women’s Hospital, Vancouver; Dr. Allan Thompson, Executive Director, Information Support, BC Ministry of Health, Victoria; and Barbara Terese, Leader, Evaluation and Utilization Management, Providence Health Care, Vancouver), the project staff (Gwyneth Aikens, Robert Chernuk, Tejinder J. Jagpal, Drew Mildon, Rena O’Neil, Pauline Sumeil and Elsie Taylor) and the participating surgeons. This project was supported by a financial contribution from the Health Transition Fund, Health Canada, and was sponsored by the Vancouver/Richmond Health Board.

References


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