Short-term and long-term outcomes of laparoscopic colectomy with multivisceral resection for surgical T4b colon cancer: Comparison with open colectomy

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Abstract
Aim: In response to the rising use of laparoscopic surgery, recent studies have shown that laparoscopic multivisceral resections for locally advanced colon cancer are safe, feasible, and provide acceptable oncological outcomes. However, the usefulness of laparoscopic multivisceral resection remains controversial. Here, we aimed to compare short-term and long-term outcomes between laparoscopic and open multivisceral resection approaches for treating locally advanced colon cancer.

Methods: We retrospectively collected data on 1315 consecutive patients admitted to the National Hospital Organization, Osaka National Hospital, for surgical treatment of colorectal cancer between 2010 and 2017. We assessed invasiveness in terms of operating times, blood loss, and complications. Oncological outcomes included 5-year survival rates and recurrences.

Results: We included 85 patients that underwent a colectomy with a multivisceral resection for locally advanced colon cancer; of these, 38 were treated with a laparoscopic approach and 47 were treated with an open approach. Compared to the open surgery group, the laparoscopic group had significantly less blood loss (median volume: 25 vs 140 mL, \( P < 0.001 \)), a lower complication rate (10.5% vs 29.8%, \( P = 0.036 \)), and shorter hospital stays (12 vs 15 days, \( P = 0.028 \)). After excluding patients with stage IV colon cancer, the groups showed similar pathologic outcomes and no significant differences in 5-year disease-free survival (73.9% vs 67.4%; \( P = 0.664 \)) or 5-year overall survival (75.8% vs 67.7%; \( P = 0.695 \)).

Conclusion: A laparoscopic approach for locally advanced colon cancer could be less invasive than an open approach without affecting oncological outcomes in selected patients.

KEYWORDS
laparoscopic surgery, locally advanced colon cancer, long-term outcomes, multivisceral resection, short-term outcomes
1 | **INTRODUCTION**

Due to advances in surgical techniques, instrumentation, and knowledge of anatomy in the field of colorectal cancer, surgical invasiveness has been reduced by performing a laparoscopic colectomy. The laparoscopic colectomy can be applied to a wide range of conditions—from early to advanced cancer and from colon cancer to rectal cancer. Large-scale randomized controlled trials and meta-analyses have indicated that laparoscopic surgery for colon cancer is equivalent or better than open surgery with regard to safety, feasibility, blood loss, postoperative pain, cosmesis, length of hospital stay, and oncological outcomes.

Locally advanced colorectal cancers sometimes invade or adhere to adjacent organs. In those cases, it is often difficult to determine whether adhesions between the tumor and the adjacent organs are due to a malignant invasion or a benign inflammatory change; consequently, radical removal requires an en bloc multivisceral resection with a safe margin. In those cases, several guidelines recommend open surgery, including the European Association of Endoscopic Surgeons, the Society of American Gastrointestinal and Endoscopic Surgeons, and the French Society of Digestive Surgery, because serious complications are associated with extended en bloc multivisceral resections. Moreover, little evidence has been published on oncological outcomes with laparoscopic surgery. The Japanese Society for Cancer of the Colon and Rectum (JSCCR) also stated that the indications for performing laparoscopic surgery to treat locally advanced colorectal cancers should be carefully considered, based on findings from the open-label, multi-institutional, randomized, phase III trial, JCOG0404.

In response to the rising use of laparoscopic surgery, recent studies have demonstrated that laparoscopic multivisceral resections are safe, feasible, and provide acceptable oncological outcomes. However, the usefulness of laparoscopic multivisceral resections for treating locally advanced colorectal cancers remains controversial, and no reports have discussed long-term outcomes for laparoscopic multivisceral resections with a sufficient median follow-up period (i.e., >5 years). Currently, data are available for long-term outcomes, including survival and recurrence, from a sufficiently long follow-up. Therefore, the present study investigated patients with locally advanced colon cancer that underwent laparoscopic or open multivisceral resections and were followed for a median of >5 years. We compared the short- and long-term outcomes between the laparoscopic and open surgical approaches.

2 | **MATERIALS AND METHODS**

2.1 | **Patients**

We retrospectively collected data on 1315 consecutive patients that were admitted to the National Hospital Organization, Osaka National Hospital, for surgical treatment of colorectal cancer between 2010 and 2017. We identified all patients that underwent a colectomy with a multivisceral resection for a locally advanced colon cancer that had invaded or adhered to adjacent organs. Of these, we included all patients that underwent emergency surgery or a primary tumor resection with distant metastases and patients with bowel obstructions that were palliated with a colonic stent, ileus tube, and stoma before surgery. We excluded patients with rectal cancer or a recurrence of colorectal cancer.

We reviewed medical and pathology reports to collect data on clinicopathological parameters, including: sex, age, body mass index (BMI), physical status according to American Society of Anesthesiologists classifications (ASA-PS), the diameter and location of the tumor, preoperative complications and treatment, the operative type, depth of tumor invasion, lymph node metastasis, lymphatic invasion, venous invasion, the pathologic stage, proximal and distal resection margins, the histological type, and the recurrence site. We also collected information on perioperative outcomes, including operating time, blood loss volume, removal of adjacent structures, conversion to open surgery, postoperative complications, length of postoperative hospital stay, and mortality. Complications were defined as those classified as grade II or higher in the Clavien–Dindo classification system. The follow-up included physical examinations and blood tests, performed every 3 months for 3 years after the operation, and every 6 months thereafter. Computed tomography (CT) was performed every 6 months. Overall survival (OS) was defined as the time from surgery to the date of death from any cause. Disease-free survival (DFS) was defined as the time from surgery to the date of recurrence or death from any cause. Previous abdominal surgery was categorized into major surgery (all resections of the gastrointestinal tract, with the exception of cholecystectomy, any kind of perforation, bleeding or peritonitis) or minor surgery (isolated abdominal wall procedures, cholecystectomy, appendectomy, and other limited intra-abdominal procedures), as described in the previous report by Neeff et al.

Written informed consent was provided by all patients that participated in this study. This study was approved by the Institutional Review Board for Studies in Humans (approval number 19-77).

2.2 | **Surgical techniques**

Radical resections were performed with either open or laparoscopic surgery. In both cases, a central vascular ligation was performed, and the entire mesocolon and all lymph nodes around the vessels supplying the tumor were removed. Standard laparoscopic surgery for colon cancer at our institution was described previously. Briefly, laparoscopic surgery was performed with five ports, including the first 12-mm trocar in the umbilicus as a camera port, another 12-mm trocar, and three 5-mm trocars. Basically, a right hemicolectomy was achieved via a retroperitoneal approach, and a left hemicolectomy, sigmoidectomy, and anterior resection were performed with a medial-to-lateral approach. However, the approach was changed, according to tumor status. The final
Incision was extended as little as possible to pull the specimen out through the umbilical incision. Which procedure would be performed depended on surgeon’s techniques and the demands of the patient. As our technical skills have matured, the range of application of laparoscopic surgery has been expanded, and cases with laparoscopic multivisceral resection have increased over time. In fact, until 2012, open multivisceral resections were more common than laparoscopic multivisceral resections in our hospital (percentage of laparoscopy: 34.5%). However, since 2013, the proportion of laparoscopies has increased (62.9%); indeed, the most recent multivisceral resections were performed with laparoscopic surgery, except for difficult cases, such as cases requiring a pancreatoduodenectomy or cases with tumors >10 cm or advanced peritoneal dissemination.

In cases with bowel obstruction, a colonic stent or ileus tube was used as a bridge to surgery because postoperative complications and mortality rates are lower with elective surgery than with emergency surgery.21 According to JSCCR guidelines, the decision to perform a radical resection of the primary tumor with distant metastasis was based on a comprehensive assessment of clinical conditions for each patient. These assessments included the symptoms related to the primary tumor, the metastatic status, the general condition of the patient, the prognosis, the risk of surgical complications, and the effect of resection.22

### 2.3 Statistical analysis

Statistical analyses were performed with JMP Pro 14 software (SAS Institute, Cary, NC). Significant differences between groups were evaluated with the Mann–Whitney test, χ² test, or Fisher’s exact test, as appropriate. OS and DFS were analyzed with the Kaplan–Meier method, and differences between the two groups were assessed with the log-rank test. Probabilities < 0.05 were considered statistically significant. Propensity scores were calculated for each patient with bivariate logistic regression on the basis of the following covariates: preoperative abscess/perforation, depth of tumor invasion, and pathologic stage. These propensity scores were

| Characteristic                          | Lap (N = 38) | Open (N = 47) | p‡  |
|-----------------------------------------|--------------|---------------|-----|
| Sex, male/female                        | 19/19        | 24/23         | 1.000† |
| Age, years                              | 70 (45-90)   | 70 (39-94)    | 0.863 |
| BMI, kg/m²                              | 22.7 (16.1-30.2) | 21.9 (16.2-31.2) | 0.155 |
| ASA-PS                                  |              |               | 0.505† |
| ≤2                                      | 25 (65.8%)   | 27 (57.5%)    |     |
| >2                                      | 13 (34.2%)   | 20 (42.5%)    |     |
| Previous major abdominal surgery        | 4 (10.5%)    | 3 (6.4%)      | 0.695‡ |
| Previous minor abdominal surgery        | 10 (26.3%)   | 10 (21.3%)    | 0.616‡ |
| Maximum diameter of tumor, mm           | 61.5 (25-150) | 71.5 (25-140) | 0.395 |
| Tumor location                          | 1.000†       |               |     |
| Right side                              | 16 (42.1%)   | 19 (40.4%)    |     |
| Left side                               | 22 (57.9%)   | 28 (59.6%)    |     |
| Preoperative complication               |              |               | 0.274‡ |
| Bowel obstruction                       | 4 (10.5%)    | 5 (10.6%)     |     |
| Abscess/perforation                     | 1 (2.6%)     | 6 (12.8%)     |     |
| Preoperative treatment                  |              |               | 1.000‡ |
| Stent/ileus tube                        | 4 (10.5%)    | 5 (10.6%)     |     |
| Stoma creation                          | 3 (8.1%)     | 4 (8.5%)      |     |
| Preoperative therapy                    | 0            | 2 (4.3%)      | 0.500 |
| Chemotherapy                            | 0            | 1 (2.1%)      |     |
| Chemoradiotherapy                       | 0            | 1 (2.1%)      |     |

**Note:** Data are expressed as the median (range) or n (%), as indicated.

**Abbreviations:** ASA-PS, American society of Anaesthesiologists - Physical Status; BMI, body mass index.

†P-values were determined with the Mann–Whitney test or

‡Fisher’s exact test.
scores (caliper = 0.2) were used to match patients in the laparoscopic surgery group 1:1 with those in the open surgery group.

3 | RESULTS

3.1 | Patient characteristics

This study included a total of 85 patients that underwent a colectomy with a multivisceral resection for locally advanced colon cancer. Of these patients, 38 underwent laparoscopy and 47 underwent open surgery. The patient characteristics between the laparoscopy and open surgery groups were not significantly different with regard to sex, age, BMI, ASA-PS, or rates of previous abdominal surgery (Table 1). The two groups were similar, in terms of the distributions of tumor diameters and tumor locations and the proportions of preoperative complications, preoperative treatments, and preoperative therapies. The patients in the open surgery group tended to be in poorer condition compared with those in the laparoscopic surgery group, because the proportion of patients with ASA-PS of >2 and with abscess/perforation was high in the open surgery group.

### Table 2 Perioperative outcomes of patients with colon cancer treated with a laparoscopic (Lap) or open (Open) multivisceral resection

| Outcome                              | Lap (N = 38) | Open (N = 47) | P† |
|--------------------------------------|---------------|---------------|----|
| Operating time, min                  | 208 (108-995) | 180 (94-561)  | 0.155 |
| Blood loss volume, mL                | 25 (0-3170)   | 140 (0-2780)  | <0.001 |
| Conversion to open surgery           | 2 (5.3%)      |               |    |
| Complications                        |               |               | 0.036‡|
| Pneumonia                            | 1 (2.6%)      | 4 (8.5%)      |    |
| Wound infection                      | 0             | 4 (8.5%)      |    |
| Urinary tract infection              | 0             | 3 (6.4%)      |    |
| Ileus                                | 1 (2.6%)      | 1 (2.1%)      |    |
| Anastomotic leakage                  | 2 (5.3%)      | 0             |    |
| Bleeding                             | 0             | 1 (2.1%)      |    |
| Other                                | 0             | 1 (2.1%)      |    |
| Mortality                            | 0             |               |    |
| Length of postoperative hospital stay, days | 12 (6-47) | 15 (6-64) | 0.028 |

Note: Data are expressed as the median (range) or n (%), as indicated. †P-values were determined with the Mann-Whitney test or ‡Fisher’s exact test.

### Table 3 Adjacent structures removed in patients with colon cancer treated with a laparoscopic (Lap) or open (Open) multivisceral resection

| Structure                        | Lap (N = 38) | Open (N = 47) |
|----------------------------------|--------------|---------------|
| Abdominal wall                   | 17           | 18            |
| Retropertitoneum                 | 6            | 14            |
| Small intestine                  | 6            | 14            |
| Bladder                          | 6            | 6             |
| Omentum                          | 4            | 4             |
| Other parts of colorectum        | 3            | 5             |
| Ovary                            | 3            | 3             |
| Gonadal vessels                  | 1            | 4             |
| Seminal vesicle                  | 1            | 2             |
| Prostate                         | 1            | 2             |
| Pancreas                         | 0            | 3             |
| Ureter                           | 0            | 2             |
| Spleen                           | 0            | 2             |
| Uterus                           | 2            | 0             |
| Iliac vessels                    | 0            | 1             |
| Iliopsoas                        | 0            | 1             |
| Duodenum                         | 0            | 1             |
| Stomach                          | 0            | 1             |
| Kidney                           | 0            | 1             |
| Liver                            | 1            | 0             |

Note: Values are the number of structures removed. In some patients, more than one structure was removed.

3.2 | Perioperative outcomes

The two groups had similar median operating times, but the laparoscopy group experienced significantly less blood loss (Table 2). Similar results were obtained only in patients who required removal of solid organs (Table S1). Two cases (5.3%) required conversion to open surgery due to uncontrolled bleeding and tumor invasion into the trigone of the bladder that made it difficult to determine whether partial cystectomy or total cystectomy should be performed. Compared to the open surgery group, the laparoscopy group showed a lower rate of complication and shorter hospital stays. There were no deaths within 30 days of surgery in either group. The most common adjacent structures removed in the laparoscopy and open surgery groups, respectively, were the abdominal wall (17 vs 18), the retroperitoneum (6 vs 14), the small intestine (6 vs 14), and the bladder (6 vs 6) (Table 3). The number of patients who required removal of solid organs was 18 (47.4%) in the laparoscopic surgery group and 28 (59.6%) in the open surgery group. Two or more structures were removed in nine patients (23.7%) in the laparoscopic surgery group and 24 patients (51.1%) in the open surgery group. These data suggested that more advanced cases were included in the open surgery group.
Pathologic and oncological outcomes

Each group included 32 patients, after excluding stage IV cases (Table 4). We did not observe any significant differences in pathologic parameters between the two groups, in terms of the depth of tumor invasion, lymph node metastasis, venous invasion, pathologic stage, lymphatic invasion, venous invasion, proximal margin, distal margin, resection margin, or histological type. The two groups included similar proportions of patients that received adjuvant chemotherapy (Table 5). Recurrence rates were similar between the two groups (25.0% vs 28.1%; \( P = 1.000 \)), and the most common recurrence site was the liver in both groups.

The median follow-up periods were similar between the laparoscopy group (61.5 months, range: 3.0-111.1 months) and the open surgery group (65.2 months, range: 7.5-119.4 months; \( P = 0.347 \)). The laparoscopic and open surgery groups did not differ significantly in the 5-year DFS (73.9% vs 67.4%; \( P = 0.664 \)) or OS (75.8% vs 67.7%; \( P = 0.695 \); Figure 1A,B). No difference between the two groups was observed even in patients who required removal of solid organs in the 5-year DFS (64.6% vs 62.1%; \( P = 0.915 \)) or OS (70.1% vs 63.6%; \( P = 0.969 \); Figure S1A,B). We applied propensity score matching considering the effect of several biases that might result from the retrospective nature of this study. After propensity score matching, there was no significant difference of oncologic outcomes between the two groups (Figure S2 and Table S2).

### 3.3 Pathologic and oncological outcomes

| Outcome | Lap (N = 32) | Open (N = 32) | \( p \)† |
|---------|-------------|--------------|-------|
| **Depth of tumor invasion** | | | 0.313‡ |
| T3 | 16 (50.0%) | 11 (34.4%) | |
| T4a | 5 (15.6%) | 4 (12.5%) | |
| T4b | 11 (34.4%) | 17 (53.1%) | |
| **Lymph node metastasis** | | | 1.000 |
| Positive | 10 (31.3%) | 11 (34.4%) | |
| Negative | 22 (68.7%) | 21 (65.6%) | |
| **Pathologic stage** | | | 1.000 |
| II | 22 (68.8%) | 21 (65.6%) | |
| III | 10 (31.2%) | 11 (34.4%) | |
| **Lymphatic invasion** | | | 0.203 |
| Positive | 22 (68.8%) | 16 (50.0%) | |
| Negative | 10 (31.2%) | 16 (50.0%) | |
| **Venous invasion** | | | 1.000 |
| Positive | 19 (59.4%) | 19 (59.4%) | |
| Negative | 13 (40.6%) | 13 (40.6%) | |
| **Proximal margin** | | | 0 |
| Negative | 32 (100%) | 32 (100%) | |
| Positive | 0 | 0 | |
| **Distal margin** | | | 0 |
| Negative | 32 (100%) | 32 (100%) | |
| Positive | 0 | 0 | |
| **Resection margin** | | | 0.536 |
| Negative | 27 (84.4%) | 24 (75.0%) | |
| Positive | 5 (15.6%) | 8 (25.0%) | |
| **R0 resection rate** | | | 0.536 |
| tub1, tub2, pap | 29 (90.6%) | 29 (90.6%) | |
| por, muc | 3 (9.4%) | 3 (9.4%) | |

Note: Data are expressed as \( n(\%) \).
Abbreviations: muc, mucinous carcinoma; pap, papillary adenocarcinoma; por, poorly differentiated adenocarcinoma; tub1, well-differentiated adenocarcinoma; tub2, moderately differentiated adenocarcinoma.

†\( p \)-values were determined with Fisher’s exact test or \( \chi^2 \) test.

### 4 Discussion

Only a few previous reports have assessed the long-term outcomes for laparoscopic multivisceral resections of locally advanced colon cancer cured by a laparoscopic or open approach through a systematic review. However, the results of these assessments were not significantly different between the two groups, which is consistent with our findings. Thus, we believe that these findings further support the use of laparoscopic techniques for the management of locally advanced colon cancer. However, further studies are needed to confirm these findings and to establish the optimal surgical approach for these patients.
coloerectal cancers. Nishikawa et al reported that a laparoscopic approach was non-inferior to an open approach in terms of DFS (median 3-year DFS: 56.7% vs 62.7%; \( P = 0.578 \)). In another study, Takahashi et al reported that OS and DFS were comparable between the laparoscopic and open surgery groups (median 3-year OS: 92.8% vs 79.8%). To our knowledge, no reports on laparoscopic multivisceral resections have discussed long-term outcomes with a sufficient follow-up period (>5 years). The present study was the first to present detailed oncologic outcomes with a sufficient median follow-up period of 61.5 months.

The 5-year DFS rates for the laparoscopic and open surgery groups in our study were 73.9% and 67.4%, respectively, comparable to those previously reported for open surgery (i.e., 56.9%-66.8%). However, it is difficult to compare our results to results from previous studies that performed multivisceral resections, due to differences between studies, including the depth of tumor invasion, the presence of lymph node metastasis, and the pathologic stage. The rates of pT4b were reported to be 28.2%-70.0%, among patients undergoing multivisceral resections for colorectal cancer. Our pT4b rates were similar, with 34.4% in the laparoscopy group and 53.1% in the open surgery group. Similar to previous studies, we found that laparoscopic surgery was associated with a lower pT4b rate than open surgery. However, this finding might have been affected by a selection bias, because more advanced cases were included in the open surgery group; this was one limitation in our retrospective study. Alternatively, the lower pT4b rate might have been due to the magnifying effect of laparoscopic surgery. Indeed, small inflammatory changes between the tumor and the adjacent organs were more likely to be detected during laparoscopic surgery compared to open surgery; thus, surgeons were more likely to resect these areas with the laparoscopic approach, compared to the open approach.

The R0 resection is the most important factor in curing colorectal cancer with a multivisceral resection. Previous studies on multivisceral resections reported R0 resection rates of 68.4%-100% with laparoscopy and 68.8%-98.5% with open surgery. Thus, our R0 resection rates were within the published range, but on the low end. Our results might have been affected by the pathologic diagnosis, because suspicious cases, for example, cases where cauterized cancer cells near the excised edge, were included in the positive resection margin group. Kim et al reported that the local recurrence rates of multivisceral resections in the laparoscopic and open surgery groups were 7.7% and 27.3%, respectively. In our study, local recurrences occurred in one patient in the laparoscopy group (3.1%) and in five patients in the open surgery group (15.6%), which suggested that our oncological clearance rate was acceptable. Liver recurrence rates in the laparoscopic surgery group seemed to be higher than that in the open surgery group (18.8% vs 6.3%; \( P = 0.257 \)). This might result from some bias induced by the small number of cases, considering the report by Hasegawa et al showing that laparoscopic and open colectomy demonstrated comparable overall colon cancer recurrence rates and recurrence sites. Our data supported the notion that our laparoscopic approach provided long-term outcomes similar to those provided with the open approach, but with less invasiveness.

Our study had several limitations. First, we studied a small number of cases and all patients were treated in a single institution. Some bias might be induced by the difference of characteristics in patients, including rates of previous major and minor abdominal surgery that were not significantly different between the laparoscopic and open surgery groups. How to deal with patients who underwent previous abdominal surgery in the prognosis analysis was debatable, because the extent of intraperitoneal adhesion might affect surgical difficulty. Second, the determination of the operation type, open or laparoscopic surgery, was inconsistent, because it was determined by the attending physician. The maturation of our technical skills has expanded the application of laparoscopic surgery to locally recurrent colorectal cancers that require a total pelvic exenteration or a sacral resection. We believe that laparoscopic surgery provides advantages over open surgery, but with less invasiveness.

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surgery, such as reduced blood loss, due to the pneumoperitoneum pressure applied, particularly in highly difficult surgeries, including multivisceral resections. Additional evidence is necessary to confirm the utility of laparoscopic surgery in this subset of patients with colon cancer that require a multivisceral resection.

**DISCLOSURE**
Conflict of Interest: Masaaki Miyo and other co-authors declare no conflict of interest.

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**REFERENCES**

1. Buunen M, Veldkamp R, Hop WC, Kuhry E, Jeekel J, Haglind E, et al. Survival after laparoscopic surgery versus open surgery for colon cancer: long-term outcome of a randomised clinical trial. Lancet Oncol. 2009;10:44–52.
2. Green BL, Marshall HC, Collinson F, Quirke P, Guillou P, Jayne DG, et al. Long-term follow-up of the Medical Research Council CLASICC trial of conventional versus laparoscopically assisted resection in colorectal cancer. Br J Surg. 2013;100:75–82.
3. Theophilos M, Platell C, Spilsbury K. Long-term survival following laparoscopic and open colectomy for colon cancer: a meta-analysis of randomized controlled trials. Colorectal Dis. 2014;16:75–81.
4. Yamamoto S, Inomata M, Katayama H, Mizusawa J, Etoh T, Konishi F, et al. Short-term surgical outcomes from a randomized controlled trial to evaluate laparoscopic and open D3 dissection for stage II/III colon cancer: Japan Clinical Oncology Group Study JCOG 0404. Ann Surg. 2014;260:23–30.
5. Veldkamp R, Kuhry E, Hop WC, Jeekel J, Kazemier G, Jaap Bonjer H, et al. Laparoscopic surgery versus open surgery for colon cancer: short-term outcomes of a randomised trial. Lancet Oncol. 2005;6:477–84.
6. Bonjer HJ, Deijen CL, Abis GA, Cuesta MA, van der Pas MHGM, de Lange-de Klerk ESM, et al. A randomized trial of laparoscopic versus open surgery for rectal cancer. N Engl J Med. 2015;372:1324–32.
7. Kang SB, Park JW, Jeong SY, Nam BH, Choi HS, Kim D-W, et al. Open versus laparoscopic surgery for mid or low rectal cancer after neoadjuvant chemoradiation therapy (COREAN trial): short-term outcomes of an open-label randomised controlled trial. Lancet Oncol. 2010;11:637–45.
8. Jeong SY, Park JW, Nam BH, Kim S, Kang S-B, Lim S-B, et al. Open versus laparoscopic surgery for mid-rectal or low-rectal cancer after neoadjuvant chemoradiotherapy (COREAN trial): survival outcomes of an open-label, non-inferiority, randomised controlled trial. Lancet Oncol. 2014;15:767–74.
9. Fuji S, Akagi T, Inomata M, Katayama H, Mizusawa J, Ota M, et al. Transitional impact of short- and long-term outcomes of a randomized controlled trial to evaluate laparoscopic versus open surgery for colorectal cancer from Japan Clinical Oncology Group Study JCOG0404. Ann Gastroenterol Surg. 2019;3:301–9.
10. Hojo S, Kawahara H, Ogawa M, Suwa K, Eto K, Yanaka H. Laparoscopic surgical challenge for T4a colon cancer. Ann Gastroenterol Surg. 2017;1:69–74.
11. Nelson H, Petrelli N, Carlin A, Couture J, Fleshman J, Guillem J, et al. Guidelines 2000 for colon and rectal cancer surgery. J Natl Cancer Inst. 2001;93:583–96.
12. Miyake Y, Nishimura J, Takahashi H, Haraguchi N, Hata T, Takemasa I, et al. The short-term outcomes of laparoscopic multivisceral resection for locally advanced colorectal cancer: our experience of 39 cases. Surg Today. 2017;47:575–80.
13. Veldkamp R, Gholghesaei M, Bonjer HJ, Meijer DW, Buunen M, Jeekel J, et al. Laparoscopic resection of colon cancer: consensus of the European Association of Endoscopic Surgery (EAES). Surg Endosc. 2004;18:1163–85.
14. Zerey M, Hawver LM, Awad Z, Stefanidis D, Richardson W, Fanelli RD. SAGES evidence-based guidelines for the laparoscopic resection of curable colon and rectal cancer. Surg Endosc. 2013;27:1–10.
15. Peschau F, Alves A, Berdah S, Laurent C, Mabrut JY, Mariette C, et al. Indications of laparoscopic general and digestive surgery. Evidence based guidelines of the French society of digestive surgery. Ann Chir. 2006;131:125–48.
16. Kitano S, Inomata M, Mizusawa J, Katayama H, Watanabe M, Yamamoto S, et al. Survival outcomes following laparoscopic versus open D3 dissection for stage II or III colon cancer (JCOG0404): a phase 3, randomised controlled trial. Lancet Gastroenterol Hepatol. 2017;2:261–8.
17. Nagasue Y, Akiyoshi T, Ueno M, Fukunaga Y, Nagayama S, Fujimoto Y, et al. Laparoscopic versus open multivisceral resection for primary colorectal cancer: comparison of perioperative outcomes. J Gastrointest Surg. 2013;17:1299–305.
18. Nishikawa T, Nozawa K, Kawai K, Sasaki K, Otani K, et al. Short- and long-term outcomes of minimally invasive versus open multivisceral resection for locally advanced colorectal cancer. Dis Colon Rectum. 2019;62:40–6.
19. Neeff H, Mariaskin D, Spangenberg HC, Hopt UT, Makowiec F. Perioperative mortality after non-hepatic general surgery in patients with liver cirrhosis: an analysis of 138 operations in the 2000s using Child and MELD scores. J Gastrointest Surg. 2011;15:1–11.
20. Miyo M, Takemasa I, Ishihara H, Hata T, Mizushima T, Ohno Y, et al. Long-term outcomes of single-site laparoscopic colectomy with complete mesocolic excision for colon cancer: comparison with conventional multiport laparoscopic colectomy using propensity score matching. Dis Colon Rectum. 2017;60:664–73.
21. Saïda Y. Current status of colonic stent for obstructive colorectal cancer in Japan; a review of the literature. J Anus Rectum Colon. 2019;3:99–105.
22. Hashiguchi Y, Muro K, Saito Y, Ito Y, Ajioka Y, Hamaguchi T, et al. Japanese Society for Cancer of the Colon and Rectum (JSCCR) guidelines 2019 for the treatment of colorectal cancer. Int J Clin Oncol. 2020;25:1–42.
23. Takahashi R, Hasegawa S, Hirai K, Hisamori S, Hida K, Kawada K, et al. Safety and feasibility of laparoscopic multivisceral resection for surgical T4b colon cancers: Retrospective analyses. Asian J Endosc Surg. 2017;10:154–61.
24. Nakafusa Y, Tanaka T, Tanaka M, Kitajima Y, Sato S, Miyazaki K. Comparison of multivisceral resection and standard operation for locally advanced colorectal cancer: analysis of prognostic factors for short-term and long-term outcome. Dis Colon Rectum. 2004;47:2055–63.
25. Eveno C, Lefevre JH, Svrcel M, Bennis M, Chafai N, Tiret E, et al. Oncologic results after multivisceral resection of clinical T4 tumors. Surgery. 2014;155:669–75.
26. Hoffmann M, Phillips C, Oevermann E, Killaitis C, Roblick U-J, Hildebrand P, et al. Multivisceral and standard resections in colorectal cancer. Langenbecks Arch Surg. 2012;397:75–84.
27. Kim KY, Hwang DW, Park YK, Lee HS. A single surgeon’s experience for locally advanced colorectal cancer: analysis of prognostic factors for short-term and long-term outcome. Dis Colon Rectum. 2020;63:965–71.
29. Ichihara M, Ikeda M, Uemura M, Miyake M, Miyazaki M, Kato T, et al. Feasibility and safety of laparoscopic lateral pelvic lymph node dissection for locally recurrent rectal cancer and risk factors for re-recurrence. Asian J Endosc Surg. 2019.

SUPPORTING INFORMATION
Additional supporting information may be found online in the Supporting Information section.

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