Comparison of patient-reported quality of life and functional outcomes following laparoscopic and transanal total mesorectal excision of rectal cancer

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INTRODUCTION

For colorectal cancer, several randomized controlled trials have suggested similar oncological outcomes between open and laparoscopic (La) total mesorectal excision (TME) [1-3]. In laparoscopic surgery for low rectal cancer, however, several factors, including obesity, advanced T stage, and narrow pelvis, result in longer operative times and increase the risk of anastomotic leakage [4], which in turn potentially increase the risk of incomplete tumor resection and local recurrence [5].

Transanal TME (TaTME) was proposed to overcome technical limitations. The oncological safety and effectiveness of TaTME...
have been described by the International TaTME registry [6]. For postoperative functions, a more precise dissection may be achieved with the transanal approach than with the laparoscopic approach; thus, leading to improvements in the urogenital function and continence [7]. Several studies have evaluated the quality of life (QoL) or functional outcomes following TaTME, with some of them comparing these parameters with those following LaTME. However, most were retrospective studies, and the administration times of the questionnaires were not consistent and the populations were small. In addition, most dealt with some functions and excluded various affected outcomes. In this study, we hypothesized that TaTME would be an alternative to the laparoscopic approach in technically difficult cases. Thus, we previously reviewed the oncological outcomes after LaTME and TaTME [8], and this study focused on comparing the postoperative QoL and functional outcomes, including bowel, anorectal, and urogenital functions, between LaTME and TaTME.

METHODS

Study population
In our early experience, we published prospective, single-arm studies, in which TaTME was performed for rectal cancer located 3–12 cm from the anal verge (AV), excluding cases with a body mass index greater than 30 kg/m², circumferential resection margin measured less than 1 mm on rectal magnetic resonance imaging, or clinical T4 stage [9,10]. However, recently, we assessed the feasibility of TaTME in challenging cases including obese patients, bulky tumors, or threatened mesorectal fascias in a prospective, single-arm setting [11]. Excluding cases with abdominoperineal resection, recurrent cancers, stage IV, or other malignancies, the oncological outcomes after LaTME and TaTME among patients with pathologically confirmed rectal adenocarcinoma treated between January 2014 and December 2017 at the National Cancer Center, Korea were reviewed retrospectively in the propensity score-matched population [8]. Different variables, including age, tumor distance from the AV, primary tumor size, neoadjuvant chemoradiotherapy, and lateral lymph node dissection, were matched during propensity score-matching analysis, and evaluation of the total cohort revealed that both groups were similar. For the same population, we performed this study involving comparisons of postoperative QoL and functional outcomes between LaTME and TaTME. This study was approved by the Institutional Review Board of National Cancer Center, Korea (No. NCC2019-0247). Informed consent was obtained from all the participants.

Preoperative evaluations, neoadjuvant therapy, and operative techniques for both LaTME and TaTME have previously been reported in detail [11]. TME was performed 4–8 weeks after completion of neoadjuvant chemoradiotherapy. A diverting ileostomy was created in case of lower anastomosis or neoadjuvant chemoradiotherapy. Stoma reversal was performed at 3 months postoperatively or 1 month after adjuvant therapy, if indicated, after evaluating the anastomosis with loopogram and sigmoidoscopy.

The outcomes were assessed with validated questionnaires and anorectal manometry at the following 2 time points: before treatment (i.e., neoadjuvant chemoradiotherapy, if indicated, or TME) and at 1 year after TME. The questionnaires were administered by a single study coordinator, and she recorded the answers directly.

Only data of patients who completed the questionnaire at the 2 time points of evaluation were analyzed. Moreover, only the results of patients who had an anorectal manometry at the 2 time points of evaluation were analyzed. Patients without stoma reversal were excluded from the analysis of bowel and anorectal function.

Questionnaires
The outcomes for all patients were assessed using the European Organization for Research and Treatment of Cancer (EORTC) Core Quality of Life (QLQ-C30). Colorectal Cancer-Specific Quality of Life (QLQ-CR38), low anterior resection syndrome (LARS) score. Fecal Incontinence Severity Index (FISI), and International Prostate Symptom Score (IPSS) questionnaires.

The EORTC QLQ-C30 was used for measuring the QoL of cancer patients and comprised 5 functional scales (including social functioning), 3 symptom scales (including nausea and vomiting), 6 single items (including appetite loss), and a global health status scale [12].

The EORTC QLQ-CR38 was used for measuring the QoL of colorectal cancer patients and comprised 4 functional scales (including body image) and 8 symptom scales (including micrurition problems and weight loss) [13]. In both the EORTC QLQ-C30 and QLQ-CR38, higher scores on the functional scales and global health status scale represent better health-related QoL. Higher scores on symptom scales/single items represent worse health-related QoL. A difference of ≥10 points was considered clinically significant [3].

The LARS score was used to evaluate the bowel and anorectal function after rectal cancer surgery. It comprises 5 items, including flatus incontinence, liquid stool incontinence, bowel frequency, clustering of stools, and urgency. On the basis of the total score, LARS is categorized as no (0–20), minor (21–29), or major (30–42) LARS [14].

The FISI was used for measuring the severity of fecal incontinence and associated factors. Each item with its different frequencies was assigned a weighted score. The total score ranged from 0 to 61. A higher score was indicative of more
severe fecal incontinence [15,16].

The IPSS was used for evaluating the factors describing the urinary function. It comprises 7 items, including incomplete emptying, frequency, intermittency, urgency, weak stream, straining, and nocturia. On the basis of the total score, the dysfunction was categorized as mild (0–7), moderate (8–19), or severe (20–35). The IPSS was also used to measure the QoL related to urinary symptoms, and the scores ranged from 0 (delighted) to 6 (terrible) [17].

Anorectal manometry
Anorectal manometry was performed to assess the anal sphincter functions. Bowel preparation was not routinely required. Patients were positioned in the left lateral decubitus position with the hips flexed at 90°. After perianal inspection and digital rectal examination, patients were evaluated using an eight-channel, water-perfused manometry system, and a standard catheter (Medical Measurement Systems, Enschede, the Netherlands) by trained technical personnel.

We evaluated 2 parameters to assess internal and external anal sphincter functions (resting pressure and squeezing pressure of voluntary contraction), and the highest value recorded was considered valid.

Statistical analysis
To compare the characteristics between the 2 groups, the t-test or Wilcoxon rank-sum test was used to analyze the continuous variables, and Pearson chi-square test or Fisher exact test was used to analyze the categorical variables. The scores for each questionnaire were summarized as the median with range or interquartile range (Q1–Q3) for the continuous variables and the frequency with proportion for the categorical variables. Differences between the 2 groups were tested using Pearson chi-square test, Fisher exact test, or Wilcoxon rank-sum test. Wilcoxon signed-rank test was used to evaluate significant changes in continuous scores over the study period in each

| Characteristic                                      | LaTME          | TaTME          | P-value |
|-----------------------------------------------------|----------------|----------------|---------|
| No. of patients                                     | 202            | 202            | 0.362   |
| Age (yr)                                            | 61.46 ± 11.24  | 62.43 ± 9.98   |         |
| Sex                                                 |                |                |         |
| Male                                                | 131 (64.9)     | 129 (63.9)     | 0.835   |
| Female                                              | 71 (35.1)      | 73 (36.1)      |         |
| Body mass index (kg/m²)                             | 24.06 ± 3.42   | 24.02 ± 3.12   | 0.915   |
| ASA PS classification                               |                |                |         |
| I or II                                             | 188 (93.1)     | 192 (95)       | 0.400   |
| III                                                 | 14 (6.9)       | 10 (5)         |         |
| Comorbidity                                         | 121 (59.9)     | 123 (60.9)     | 0.839   |
| Previous abdominal open surgery                     | 43 (21.3)      | 42 (20.8)      | 0.903   |
| Tumor distance from the anal verge                  |                |                |         |
| ≤5 cm                                               | 98 (48.5)      | 83 (41)        | 0.238   |
| ≤10 cm                                              | 94 (46.5)      | 111 (55)       |         |
| >10 cm                                              | 10 (5)         | 8 (4)          |         |
| Tumor size (cm)                                     | 2.4 (0–8.5)    | 2.3 (0–14.0)   | 0.914   |
| Preoperative T stage                                 |                |                |         |
| T1                                                  | 24 (11.9)      | 24 (11.9)      | 0.960   |
| T2                                                  | 24 (11.9)      | 27 (13.4)      |         |
| T3                                                  | 136 (67.3)     | 135 (66.8)     |         |
| T4                                                  | 18 (8.9)       | 16 (7.9)       |         |
| Preoperative N stage                                 |                |                |         |
| N–                                                  | 61 (30.2)      | 66 (32.7)      | 0.592   |
| N+                                                  | 141 (69.8)     | 136 (67.3)     |         |
| Neoadjuvant chemoradiotherapy                        | 118 (58.4)     | 129 (63.9)     | 0.262   |
| Operative time (min)                                | 205 (90–605)   | 215 (90–705)   | 0.838   |
| Open conversion                                     | 0 (0)          | 5 (2.5)        | 0.061   |
| Diverting ileostomy                                 | 151 (74.8)     | 168 (83.2)     | 0.038   |
| Lateral lymph node dissection                       | 28 (13.9)      | 19 (9.4)       | 0.163   |

Values are presented as number only, mean ± standard deviation, number (%), or median (range). ASA, American Society of Anesthesiologists; PS, physical status; LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision.
The LARS score was assessed using univariable and multivariable logistic regression models with the binary 1-year LARS score as a dependent variable (no vs. minor/major LARS). All results were considered statistically significant when the 2-sided P-value was <0.05. Statistical analyses were performed with SAS software ver. 9.4 (SAS Institute, Cary, NC, USA) and R software ver. 3.6.2 (R Foundation for Statistical Computing, Vienna, Austria).

RESULTS

During the study period, 514 patients underwent LaTME, and 208 underwent TaTME. After matching, 202 patient pairs were included. Both groups showed similar baseline characteristics except for a diverting ileostomy (74.8% in the laparoscopic group vs. 83.2% in the transanal group, P = 0.038) (Table 1). Stoma reversal was not performed in 15 patients; 7 in the LaTME group and 8 in the TaTME group.

Quality of life based on the EORTC QLQ-C30 and QLQ-CR38

A total of 113 patients completed the EORTC QLQ-C30 at both time points of evaluation: 60 patients in the LaTME group and 53 patients in the TaTME group. Before treatment, the global health status scale score was significantly worse in the TaTME group than in the LaTME group (P = 0.004), but it was comparable between the groups at 1 year after TME (Table 2). Thus, we did a pairwise comparison of the scale at different times in each group. In the LaTME group, the scale showed no significant changes at 1 year postoperatively. However, in the TaTME group, the scale at 1 year postoperatively showed improvement relative to that before treatment (P = 0.024). Otherwise, both groups were comparable (Table 2).

A total of 115 patients (62 patients in the LaTME group and 53 patients in the TaTME group) answered the EORTC QLQ-CR38 at both time points of evaluation. The scales for chemotherapy side effects and stoma-related problems were omitted before treatment. The scale for chemotherapy side effects was calculated for patients who had received neoadjuvant chemoradiotherapy or adjuvant chemotherapy, and the result was comparable between the groups. For 1 patient, a stoma reversal was not performed and questions for stoma-related problems 1 year postoperatively were not answered. The questions about sexual enjoyment, male sexual problems, and female sexual problems, which were applicable to sexually active patients, were completed by 6, 7, and zero patients, respectively, at 1 year. At 1 year, the score for micturition problems was worse in the LaTME group than in the TaTME group (P = 0.016). Both groups showed comparable results for the remaining scales (Table 3).

Bowel and anorectal functions based on the LARS score, FISI, and anorectal manometry

Table 4 shows a comparison of the LARS score, FISI, and the

| Table 2. Results of the European Organization for Research and Treatment of Cancer Core Quality of Life Questionnaire |
|------------------|------------------|------------------|------------------|------------------|
| Variable         | Before treatment | 1 Year after surgery |
|                  | LaTME (n = 60)   | TaTME (n = 53)   | P-value          | LaTME (n = 60)   | TaTME (n = 53)   |
| Global health status scale scorea | 62.5 (8.33–100) | 50 (50–83.33) | 0.004 | 66.67 (33.33–100) | 66.67 (33.33–100) | 0.456 |
| Functional scale scoreb | | | | | |
| Physical functioning | 100 (66.67–100) | 100 (66.67–100) | 0.523 | 100 (46.67–100) | 100 (33.33–100) | 0.937 |
| Role functioning | 100 (66.67–100) | 100 (66.67–100) | 0.274 | 100 (33.33–100) | 100 (33.33–100) | 0.280 |
| Emotional functioning | 91.67 (41.67–100) | 91.67 (33.33–100) | 0.984 | 100 (58.33–100) | 100 (33.33–100) | 0.368 |
| Cognitive functioning | 100 (66.67–100) | 100 (66.67–100) | 0.779 | 100 (50–100) | 100 (83.33–100) | 0.304 |
| Social functioning | 100 (33.33–100) | 100 (33.33–100) | 0.230 | 100 (66.67–100) | 100 (66.67–100) | 0.464 |
| Symptom scale scoreb | | | | | |
| Fatigue | 0 (0–55.56) | 0 (0–55.56) | 0.713 | 0 (0–55.56) | 0 (0–22.22) | 0.684 |
| Nausea and vomiting | 0 (0–16.67) | 0 (0–16.67) | 0.940 | 0 (0–16.67) | 0 (0–0) | 0.357 |
| Pain | 0 (0–83.33) | 0 (0–33.33) | 0.607 | 0 (0–33.33) | 0 (0–16.67) | 0.491 |
| Dyspnea | 0 (0–33.33) | 0 (0–33.33) | 0.642 | 0 (0–33.33) | 0 (0–100) | 0.489 |
| Insomnia | 0 (0–100) | 0 (0–100) | 0.044 | 0 (0–66.67) | 0 (0–66.67) | 0.300 |
| Appetite loss | 0 (0–66.67) | 0 (0–66.67) | 0.882 | 0 (0) | 0 (0–33.33) | 0.295 |
| Constipation | 0 (0–100) | 0 (0–100) | 0.236 | 0 (0–66.67) | 0 (0–66.67) | 0.491 |
| Diarrhea | 0 (0–100) | 0 (0–66.67) | 0.993 | 0 (0–33.33) | 0 (0–33.33) | 0.861 |
| Financial difficulties | 0 (0–66.67) | 0 (0–66.67) | 0.292 | 0 (0–66.67) | 0 (0–33.33) | 0.286 |

Values are presented as median (range); the scale ranges from 0 to 100.
LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision.

aA higher score represents better health-related quality of life. bA higher score represents worse health-related quality of life.
Table 3. Results of the European Organization for Research and Treatment of Cancer Colorectal Cancer-Specific Quality of Life Questionnaire

| Variable                        | Before treatment |                        | P-value |                        |                        | P-value |
|---------------------------------|------------------|------------------------|---------|------------------------|------------------------|---------|
|                                 | LaTME (n = 62)   | TaTME (n = 53)         |         | LaTME (n = 62)         | TaTME (n = 53)         |         |
|                                 | n                | Median (range)         |         | n                      | Median (range)         |         |
|                                 | P-value          |                        |         |                        |                        |         |
| Functional scale score          |                  |                        |         |                        |                        |         |
| Body image                      | 62               | 100 (66.67–100)        | 0.611   | 62                     | 100 (55.56–100)        | 0.921   |
| Sexual functioning              | 62               | 0 (0–100)              | 0.026   | 62                     | 0 (0–50)               | 0.844   |
| Sexual enjoyment                | 11               | 66.67 (33.33–100)      | >0.999  | 3                      | 33.33 (33.33–66.67)    | 0.505   |
| Future perspective              | 62               | 33.33 (0–100)          | 0.774   | 62                     | 66.67 (33.33–100)      | 0.821   |
| Symptom scale score             |                  |                        |         |                        |                        |         |
| Micturition problems            | 62               | 0 (0–44.44)            | 0.076   | 62                     | 11.11 (0–44.44)        | 0.016   |
| Gastrointestinal symptoms       | 62               | 0 (0–66.67)            | 0.491   | 62                     | 0 (0–26.67)            | 0.228   |
| Chemotherapy side effects       | NA               | NA                     | NA      | NA                     | NA                     | NA      |
| Defecation problems             | 62               | 9.52 (0–66.67)         | 0.185   | 61                     | 14.29 (0–52.38)        | 0.773   |
| Stoma-related problems          | NA               | NA                     | NA      | NA                     | NA                     | NA      |
| Male sexual problems            | 9                | 0 (0–33.33)            | 0.705   | 4                      | 25 (0–100)             | >0.999  |
| Female sexual problems          | 3                | 16.67 (0–16.67)        | 0.617   | NA                     | NA                     | NA      |
| Weight loss                     | 62               | 0 (0–100)              | 0.834   | 62                     | 0 (0–33.33)            | 0.662   |

Values are presented as the number of responders for each scale and median (range) of the score; the scale ranges from 0 to 100.
LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision; NA, not available.

aA higher score represents better health-related quality of life. bA higher score represents worse health-related quality of life.
Table 4. Results of the LARS score, FISI, and manometric evaluations

| Variable | Before treatment | 1 Year after surgery |
|----------|-----------------|---------------------|
|          | LaTME (n = 44)  | TaTME (n = 42)      | P-value | LaTME (n = 44)  | TaTME (n = 42) | P-value |
| LARS score | 11 (0–23)       | 13 (0–25)           | 0.874   | 13 (0–22.5)     | 25 (15–32)     | <0.001  |
| LARS score category |  |  |  |  |  |  |
| No LARS  | 33 (75.0)       | 31 (73.8)           | 0.720   | 28 (63.6)       | 14 (33.3)      | 0.004   |
| Minor LARS | 5 (11.4)       | 3 (7.1)             |         | 13 (29.6)       | 15 (35.7)      |         |
| Major LARS | 6 (13.6)       | 8 (19.1)            | 0.001   | 3 (6.8)         | 13 (31.0)      | 0.001   |
| FISI | 0 (0–0) | 0 (0–0) | 0.961 | 0 (0–10) | 6 (0–12) | 0.162 |
| Manometric evaluation |  |  |  |  |  |  |
| Maximal resting pressure (mmHg) | 65 (54–84) | 69 (51–86) | 0.719 | 46 (30–66) | 45 (34–57) | 0.979 |
| Maximal squeezing pressure (mmHg) | 228 (150–256) | 220 (140–266) | 0.929 | 199 (143–263) | 185 (134–244) | 0.889 |

Values are presented as median (interquartile range) or number (%).

LARS, low anterior resection syndrome; FISI, Fecal Incontinence Severity Index; LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision.

Fig. 1. Scores measured before treatment (Pre) and at 1 year postoperatively (Post 1Y). (A) Lower anterior resection syndrome (LARS) score, (B) Fecal Incontinence Severity Index (FISI), (C) maximal resting pressure on anorectal manometry, and (D) maximal squeezing pressure on anorectal manometry. LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision.
results of anorectal manometry between the groups. Fig. 1 shows the change in each index before and after treatment in both groups.

LARS score
Eighty-seven patients completed the LARS questionnaire at both time points of evaluation. Excluding 1 patient for whom stoma reversal was not noted, 86 patients were included in the analysis: 44 patients in the LaTME group and 42 patients in the TaTME group. At 1 year postoperatively, the LARS score was significantly higher in the TaTME group than in the LaTME group (P < 0.001). A greater number of patients were classified as having major LARS at 1 year (P = 0.004) after TaTME than after LaTME. In the TaTME group, the scores did not significantly differ between before treatment and at 1 year postoperatively. However, in the TaTME group, the score at 1 year was significantly higher than that before treatment (P < 0.001).

When comparing the baseline characteristics of the LARS questionnaire responders (Table 5), the TaTME group showed a significantly shorter tumor distance from the AV (P = 0.005), shorter anastomosis distance from the AV (P < 0.001), higher frequency of hand-sewn colorectal anastomosis (P = 0.006), and longer operative time (P < 0.001) than did the LaTME group. Thus, we examined the difference in 1-year LARS (no vs. minor/major) between the LaTME and TaTME groups in a multivariable model that was adjusted for 2 clinical factors, neoadjuvant chemoradiotherapy and anastomosis distance from the AV, which statistically affect 1-year LARS (no vs. minor/major) (Table 6). The results showed that there was no statistically significant difference between the groups (odds ratio, 2.30; 95% confidence interval, 0.79–6.72; P = 0.127).

FISI
A total of 118 patients completed the FISI questionnaire at both time points of evaluation. Excluding a patient for whom stoma reversal was not noted, 117 patients were included in the analysis: 62 patients in the LaTME group and 55 patients in the TaTME group. Significant differences were not observed between the groups. In both groups, the FISI at 1 year was significantly increased (P = 0.003 in the LaTME group and P < 0.001 in the TaTME group) relative to the score before treatment.

Table 5. Baseline characteristics of the LARS questionnaire responders

| Characteristic                          | LaTME            | TaTME            | P-value |
|----------------------------------------|------------------|------------------|---------|
| No. of responders                      | 44               | 42               |         |
| Age (yr)                               | 62.2 ± 10.4      | 65.6 ± 8.8       | 0.103   |
| Body mass index (kg/m²)                | 23.6 ± 3.4       | 25.1 ± 3.5       | 0.049   |
| Sex                                    |                  |                  |         |
| Male                                   | 30 (68.2)        | 26 (61.9)        | 0.542   |
| Female                                 | 14 (31.8)        | 16 (38.1)        |         |
| Previous abdominal open surgery        | 6 (13.6)         | 6 (14.3)         | 0.931   |
| Tumor distance from the AV (cm)        | 8.0 (1.0–15.0)   | 6.0 (1.0–10.0)   | 0.005   |
| Tumor size (cm)                        | 2.9 (0.0–8.5)    | 2.3 (0.4–6.0)    | 0.414   |
| Preoperative T stage                   |                  |                  |         |
| T1                                     | 6 (13.6)         | 6 (14.3)         | 0.907   |
| T2                                     | 7 (15.9)         | 8 (19.0)         |         |
| T3                                     | 27 (61.4)        | 26 (61.9)        |         |
| T4                                     | 4 (9.1)          | 2 (4.8)          |         |
| Preoperative N stage                   |                  |                  |         |
| N–                                     | 16 (36.4)        | 16 (38.1)        | 0.868   |
| N+                                     | 28 (63.6)        | 26 (61.9)        |         |
| Neoadjuvant chemoradiotherapy          | 20 (45.5)        | 22 (52.4)        | 0.521   |
| Anastomosis distance from the AV (cm)  | 4.0 (1.0–12.0)   | 3.0 (1.0–6.0)    | <0.001  |
| Anastomosis method                     |                  |                  |         |
| Stapled CRA                            | 37 (84.1)        | 24 (57.1)        | 0.006   |
| Hand-sewing CAA                        | 7 (15.9)         | 18 (42.9)        |         |
| Lateral lymph node dissection          | 3 (6.8)          | 4 (9.5)          | 0.710   |
| Diverting ileostomy                    | 26 (59.1)        | 33 (78.6)        | 0.052   |
| Open conversion                        | 0 (0)            | 1 (2.4)          | 0.488   |
| Operative time (min)                   | 170 (100–425)    | 220 (135–490)    | <0.001  |

Values are presented as number only, mean ± standard deviation, number (%), or median (range).
LARS, lower anterior resection syndrome; LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision; AV, anal verge; CRA, colorectal anastomosis; CAA, coloanal anastomosis.
Anorectal manometry

A total of 136 patients underwent anorectal manometry at both time points of evaluation; 54 patients in the LaTME group and 82 patients in the TaTME group. Stoma reversal was successful in all patients. While no significant differences were observed between the groups, both the maximal resting pressure (P < 0.001 in the LaTME group and P < 0.001 in the TaTME group) and maximal squeezing pressure (P = 0.002 in the LaTME group and P < 0.001 in the TaTME group) at 1 year were significantly decreased relative to the values noted before treatment.

Urinary function based on IPSS

A total of 115 patients, 62 patients in the LaTME group and 53 patients in the TaTME group, completed the International Prostate Symptom Score (IPSS) before treatment and 1 year after surgery. The results are presented in Table 7.

Table 6. Results of the univariable and multivariable logistic regression analyses for the LARS score at 1 year postoperatively

| Variable | No LARSc | Minorb/major LARSd | Univariable | Multivariabled |
|----------|-----------|-------------------|-------------|----------------|
|          | (n = 42), n (%) | (n = 44), n (%) | OR (95% CI) P-value | OR (95% CI) P-value |
| Age      | 1.06 (1.01–1.11) 0.023 |         |             |               |
| Sex      |          |                   |             |               |
| Male     | 30 (53.6) | 26 (46.4) | 1           |               |
| Female   | 12 (40.0) | 18 (60.0) | 1.73 (0.70–4.26) 0.232 |               |
| Body mass index | 1.00 (0.88–1.12) 0.931 |         |             |               |
| Tumor distance from the AV | 0.77 (0.66–0.91) 0.002 |         |             |               |
| Surgical method |          |                   |             |               |
| LaTME    | 28 (63.6) | 16 (36.4) | 1           |               |
| TaTME    | 14 (33.3) | 28 (66.7) | 3.50 (1.44–8.51) 0.006 | 2.30 (0.79–6.72) 0.127 |
| Anastomosis distance from the AV | 0.59 (0.44–0.78) <0.001 | 0.64 (0.47–0.88) 0.006 |               |
| Anastomosis method |          |                   |             |               |
| Stapled CRA | 36 (59.0) | 25 (41.0) | 1           |               |
| Hand-sewing CAA | 6 (24.0) | 19 (76.0) | 4.56 (1.60–13.03) 0.005 |               |
| Lateral lymph node dissection | 1 (14.3) | 6 (85.7) | 6.47 (0.75–56.22) 0.091 |               |
| Diverting ileostomy | 22 (37.3) | 37 (62.7) | 4.81 (1.75–13.19) 0.002 |               |
| Operative time | 1.01 (1.01–1.02) 0.001 |         |             |               |
| Pretreatment LARS category |          |                   |             |               |
| No LARSc | 34 (53.1) | 30 (46.9) | 1           |               |
| Minorb/major LARSd | 8 (36.4) | 14 (63.6) | 1.98 (0.73–5.38) 0.179 |               |

LARS, low anterior resection syndrome; OR, odds ratio; CI, confidence interval; AV, anal verge; LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision; CRA, colorectal anastomosis; CAA, coloanal anastomosis.

Table 7. Results of the IPSS

| Variable | Before treatment | 1 Year after surgery |
|----------|------------------|----------------------|
|          | LaTME (n = 62)   | TaTME (n = 53)       | LaTME (n = 62)   | TaTME (n = 53)       |
| IPSS     | 2 (0–4)          | 2 (0–7)              | 3 (1–6)          | 3 (2–5)              | 0.723 |
| IPSS categories |          |                      |                    |                      |
| Mild dysfunctiona | 54 (87.1) | 40 (75.5) | 0.144 | 51 (82.3) | 45 (84.9) | 0.523 |
| Moderate dysfunctionb | 8 (12.9) | 11 (20.7) | 0.049 | 11 (17.7) | 7 (13.2) |
| Severe dysfunctionc | 0 (0) | 2 (3.8) | 0 (0) | 1 (1.9) |
| Urinary symptom-related QoL scale | 0 (0–1) | 0 (0–1) | 0.457 | 0 (0–1) | 0 (0–1) | 0.892 |

Values are presented as median (interquartile range) or number (%).

IPSS, International Prostate Symptom Score; LaTME, laparoscopic total mesorectal excision; TaTME, transanal total mesorectal excision; QoL, quality of life.

a)0–7, b)8–19, c)20–35.
patients in the TaTME group completed the IPSS questionnaire at both time points of evaluation. Significant differences between the groups were not observed with respect to the total score, severity score, and QoL score related to urinary symptoms (Table 7). In the LaTME group, the IPSS at 1 year was significantly increased (P = 0.011) relative to the scores noted before treatment. However, in the TaTME group, significant changes in the IPSS were not observed at 1 year postoperatively (Fig. 2).

**DISCUSSION**

This study evaluated and compared QoL and functional outcomes following LaTME and TaTME in rectal cancer patients. The health-related QoL scores and IPSS were similar between the groups. Although the FISI and anorectal manometry results were comparable between the groups, the postoperative LARS score was worse in the TaTME group than in the LaTME group. However, multivariable analysis showed no statistically significant difference in the LARS score between the groups at 1 year postoperatively (P = 0.127).

For low rectal cancer, sphincter-preserving surgery with intersphincteric resection may prevent a permanent stoma, but lower anastomosis consequently results in LARS in the long term [18] and affects QoL [19]. Previous studies reported comparable LARS scores between patients who underwent LaTME and those who underwent TaTME [20,21]; however, the sample populations in these studies were small. In this study, a greater number of patients were classified as having major LARS at 1 year postoperatively after TaTME than after LaTME. Endoanal instrumentation during the transanal approach may have caused anal sphincter damage and aggravated LARS [22]; this notion was supported by Keller et al. [23] who reported that longer perineal operation time was associated with worsening of LARS. However, owing to the significantly lower anastomosis level among the LARS questionnaire responders in the TaTME group, we performed a multivariable analysis and found that TaTME was not a significant risk factor for LARS. In addition, the EORTC QoL scale score related to gastrointestinal symptoms, FISI, and results of manometric evaluations were comparable between the groups.

In this study, as in previous studies, neoadjuvant chemoradiotherapy and lower anastomosis were identified as significant predictors of LARS. Among patients who underwent curative low anterior resection for nondisseminated rectal cancer, neoadjuvant therapy, short- or long-course (chemo) radiotherapy, and TME were strongly associated with major LARS [24]. The FOWARC (Neoadjuvant FOLFOX6 Chemotherapy With or Without Radiation in Rectal Cancer) trial suggested that long-course neoadjuvant radiation therapy and lower anastomosis were the independent risk factors for major LARS [25].

Previous studies that used the EORTC QLQ-C30 showed comparable outcomes of QoL between the groups [21,26]. However, in this study, the score of global health status scale before treatment was worse in the TaTME group than in the LaTME group. However, the analysis for each questionnaire was based on the number of subjects who responded to that questionnaire; therefore, differences in baseline characteristics may exist between the groups. In addition, we did a pairwise comparison of each group for the score of global health status scale, and the result showed that the TaTME group improved after surgery, while the LaTME group showed no significant changes over the study period. This result may suggest a more severe cancer status in the TaTME group than in the LaTME group. In addition, the global health status score was comparable between the groups at 1 year postoperatively, and the results for the remaining scales were comparable between the groups.

With respect to the EORTC QLQ-CR38 or QLQ-CR29 (updated version), previous studies reported that symptoms, including fecal incontinence, were worse in the TaTME group than in the LaTME group [21,26]. In this study, the results for the related scales were comparable between the groups. In addition, both groups were not significantly different with respect to the LARS score and FISI, which are more specifically developed to assess bowel and anorectal functions than is EORTC QLQ-CR38 or QLQ-CR29.

The IPSS was comparable between the groups; this finding was in agreement with those reported in previous studies [20,21]. However, Bjoern et al. [26] reported that the QoL related to urinary symptoms was better for patients who underwent TaTME than for those who underwent LaTME. In this study, pretreatment IPSS was comparable between the
groups. Assuming that there were no significant differences in the incidence of symptomatic benign prostatic hyperplasia or degree of cancer progression between the groups. However, IPSS significantly worsened at 1 year postoperatively in the LaTME group but was comparable before and after treatment in the TaTME group. In addition, the EORTC QoL scale for micturition problems at 1 year was worse in the LaTME group than in the TaTME group. During LaTME, pelvic dissection and traction of the bladder near the Denovilliers’ fascia may result in damage to the autonomic nerves [27]. Thus, TaTME may aid in the preservation of the autonomic nerves.

With respect to male sexual function, Pontallier et al. [20] reported that patients in the TaTME group (71%) and LaTME group (39%) in their study maintained sexual activity postoperatively, and the laparoscopic approach was a predictive factor for the loss of sexual activity. They also reported better postoperative erectile and ejaculatory functions in the TaTME group than in the LaTME group, although the difference was not statistically significant. With respect to female sexual function, previous studies reported low completion rates of the questionnaires and low rate of sexual activity as limitations [23,28]. In this study, the sexual function of 260 male patients was assessed with the International Index of Erectile Function questionnaire, which was applicable to sexually active patients: 104 patients completed the questionnaire before treatment, and 25 reported sexual activity (18 patients in the LaTME group and 7 patients in the TaTME group). Among the patients who were sexually active before treatment, 6 patients in the LaTME group and 2 patients in the TaTME group maintained sexual activity at 1 year postoperatively. We also assessed female sexual function using the Female Sexual Function Index questionnaire; however, only 2 women responded at 1 year after TME. In addition, the response rates for the EORTC QoL sexual scales were low. Owing to the small size of the sample population, valid statistical analysis for sexual function could not be performed. Thus, larger-scale studies are required to evaluate sexual function following TaTME.

Compared with previous studies, the present study holds strength in the fact that we attempted to evaluate QoL and the varied spectra of the affected functions following TaTME. In addition, previous relevant studies evaluated the questionnaires not at a designated time point, but for a certain period after TME. However, we evaluated the questionnaires at designated time points, before treatment and at 1 year after TME when patients visited the outpatient clinic. The consistency of the study design, such that only patients with available data for the 2 time points assessed were included, is another strength of the study. Furthermore, this study is the first to analyze the results of anorectal manometry with respect to the evaluation of the functional outcomes of TaTME. The procedure may be examiner-dependent, but it provides digitized information and complements the subjectivity of the questionnaires. In contrast, the subjectivity of the questionnaires may lead to differences between the self-assessed functional outcomes and real functions.

In addition, only data of patients who completed the questionnaires or manometric evaluations at the 2 designated time points were analyzed in each evaluation. Therefore, differences in baseline characteristics may exist between the groups, and this is a major limitation of our study. This was a retrospective study, and the response rate was low. Nevertheless, we tried to evaluate QoL and the varied spectra of the affected functions in 1 population. Thus, we used the previous propensity score-matched population and hypothesized that the 2 groups of each evaluation would be balanced in baseline characteristics. In addition, this was a single-institution study, which may not be representative of general outcomes. Another study limitation includes the preoperative LARS evaluation. The LARS score is not validated for use in a preoperative group, but we applied it before TME to compare it with that after TME.

In conclusion, LaTME and TaTME were comparable in terms of QoL and bowel, anorectal, and urinary functions. The rate of LARS was higher in the TaTME group at 1 year postoperatively; however, large prospective studies with long-term follow-up are required to clarify whether the higher rate of LARS following TaTME is the result of selecting patients with tumors located close to the AV or whether it is attributable to the TaTME technique itself.

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Conflicts of Interest
No potential conflict of interest relevant to this article was reported.

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