Robotic or laparoscopic surgery for rectal cancer - which is the best answer? A comprehensive review of oncological outcomes

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Abstract

Treatment of rectal cancer is ever evolving with the introduction of newer surgical technologies and multimodal treatment approach. The literature evaluating the various surgical treatment options with regards to operative and nonoperative outcomes is abundant. This is a comprehensive review focused on oncological outcomes of rectal cancer resection performed robotically or laparoscopically. Based on the current literature available, there is no significant difference in total mesorectal excision completeness, lymph node harvest, positive circumferential resection margin, or proximal resection margin between robotic and laparoscopic approaches for rectal resection. Selection of surgical approach should not be based on pathological outcomes as they are equivalent.

Keywords: Robotic, robotics, laparoscopic, laparoscopy, rectal cancer, rectal carcinoma, total mesorectal excision

INTRODUCTION

The treatment of rectal cancer has evolved during the last several decades into a multidisciplinary model of care. During this time, surgical innovations continued to revolutionize treatment and improve patient outcomes, most notably the introduction of total mesorectal excision (TME) by Heald et al. [1]. This landmark discovery changed the trajectory of rectal cancer resections and greatly improved patient outcomes by reducing pelvic recurrences. Since that time, laparoscopic TME was introduced and has
now become standard of care after several large randomized controlled trials (RCTs) assessed oncologic outcomes and early postoperative recovery\[2-8\].

Performing a laparoscopic TME is not without its challenges, especially in a deep narrow pelvis with the two-dimensional view and limited dexterity. Robotic TME was introduced to overcome some of these challenges. The theoretical technical advantages of robotic TME include a stable camera platform, three-dimensional view, and better articulation of the surgical instruments\[9\]. Although this technology has gained widespread popularity, it is not without its own set of challenges, including higher cost, longer operative time, and loss of tactile sensation.

Surgical innovation continues to play a vital role in the multimodal treatment of rectal cancer. Examining the pathologic outcomes is important to ensure appropriate care is provided when introducing new technologies. To date, the largest RCT available to compare laparoscopic and robotic rectal resections is the ROLARR trial\[10\]. Several other RCTs are now available, along with numerous meta-analyses to further evaluate the literature on pathologic outcomes with robotic compared to laparoscopic rectal cancer resections, which are discussed in this review. This review is Part 2 of a two-part series, in which the non-oncologic outcomes and learning curve are separately discussed.

Pathologic outcomes

**Total mesorectal excision grade**

When assessing pathologic outcomes for rectal cancer, the completeness of the TME is one of the important oncologic factors to consider. It is also a useful marker to compare the effectiveness and safety of the various surgical techniques, such as laparoscopic and robotic. The three RCTs discussed below on robotic vs. laparoscopic approach for the treatment of rectal cancer have assessed TME grade or completeness and none have shown a significant difference in the quality of TME specimen\[10-12\]. Furthermore, multiple meta-analyses have also shown no significant differences.

The ROLARR RCT trial included TME pathology specimen grading using the method of Quirke and Dixon for completeness and found complete TME in 77.6% of laparoscopic specimens vs. 76.4% of robotic specimens ($P = 0.14$)\[10\]. A phase II open label prospective RCT also assessed the quality of TME by a pathologist, as the primary outcome and found similar results: complete TME 78.1% (laparoscopic) vs. 80.3% (robotic) and near complete in 21.9% (laparoscopic) vs. 18.2% (robotic) ($P = 0.599$)\[11\]. They did, however, note one incomplete TME (1.5%) in the robotic group and none in the laparoscopic group. Lastly, a smaller pilot RCT also found no difference in macroscopic judgement of the TME specimen with complete TME noted in 17 of 18 robotic samples and 1 nearly complete vs. 13 of 16 complete TME and 3 nearly complete ($P = 0.323$)\[12\].

A recent meta-analysis by Eltair et al.\[13\] assessed the robotic and laparoscopic approaches for the treatment of rectal cancer within nine RCTs that included 1463 patients (728 robotic vs. 735 laparoscopic)\[13\]. Four RCTs were included in the macroscopic assessment of complete TME, including the three discussed above. They found no statistically significant difference in complete resection with zero heterogeneity in their assessment. Simillis et al.\[14\] compared open vs. laparoscopic vs. robotic vs. transanal mesorectal excision for rectal cancer in their meta-analysis and included 29 RCTs. The authors reported significantly higher incomplete or nearly complete TME in the laparoscopic vs. open group (odds ratio of 1.52), but no differences in the laparoscopic vs. robotic (odds ratio of 0.98) approaches. A recent network meta-analysis assessed the quality of TME and reported no difference in complete mesorectum excision in the pooled analysis of 11 studies\[15\]. Nine studies were included in the pooled analysis examining near-complete mesorectal excision and also reported no difference when comparing laparoscopic vs. robotic methods.
One meta-analysis that included 12 studies (11 case-control and only 1 RCT) did find a higher complete TME in robotic vs. laparoscopic surgery (odds ratio of 1.83, \( P = 0.03 \)), however there was significant heterogeneity noted in the analysis (\( I^2 = 47\% \))\cite{16}. Furthermore, this analysis included a majority of case-control studies, which are of lower level of evidence, while other meta-analyses have included more RCTs and prospective studies.

**Lymph node harvest**

The current guidelines, including those of the American Joint Committee on Cancer (AJCC) and College of American Pathologists (CAP), recommend a minimum of 12 lymph nodes be examined to accurately stage rectal cancer in order to aid in the decision for adjuvant treatment\cite{17-19}. The reasons for low lymph node harvest can include neoadjuvant treatment, lack of high ligation of the vessels, and potentially poor surgical or pathologic technique. When comparing surgical approaches for rectal cancer, it is important to evaluate lymph node harvest with each technique.

The ROLARR RCT performed an intention to treat analysis in which one of the outcomes measured was median lymph nodes retrieved\cite{10}. They reported no differences; both groups yielded a high number of lymph nodes: 24.1 (laparoscopic) vs. 23.2 (robotic), almost double the minimal requirement. Kim et al.\cite{11} noted a higher number of lymph nodes in the robotic (median 18) compared to the laparoscopic group (median 15) (\( P = 0.04 \)) in their RCT. They also examined the rate of 12 or more lymph nodes retrieved in their groups and found 90.9% of patients achieved this benchmark in the robotic group compared to 74% of patients in the laparoscopic group. Of note, the majority of patients in this single-center RCT received preoperative chemoradiation (77.3% in robotic vs. 77.5% in laparoscopic), which might have led to the lower number of lymph nodes.

A seven-institution multicenter study examined consecutive patients who underwent robotic or laparoscopic intersphincteric resection for low rectal cancer\cite{20}. Propensity score analysis was performed with 1:1 case-match, in which no difference was found in the number of lymph nodes retrieved (\( P = 0.126 \)) or the number of positive lymph nodes (\( P = 0.712 \)). Kim et al.\cite{21} also used propensity score matching to analyze their retrospective cohort and, after matching, found no difference in the number of harvested lymph nodes (\( P = 0.44 \)). Furthermore, a propensity score match study was performed in consecutive obese patients who underwent laparoscopic or robotic rectal resection at three centers, and no difference was noted in the mean lymph node yield (17 in robotic vs. 16 in laparoscopic, \( P = 0.639 \))\cite{22}. A single-center study examined their prospectively collected database of mid to distal rectal cancers and found a higher median number of lymph nodes harvested (12 in laparoscopic vs. 14 in robotic, \( P = 0.002 \))\cite{23}. However, the groups however were not matched between the median tumor distance of 8 cm in laparoscopic vs. 7 cm in robotic. Moreover, there were more male patients, more comorbidities, and preoperative radiation in the robotic surgery group.

Multiple meta-analyses examining the highest level of evidence available in the form of RCTs have found no difference in the number of lymph nodes retrieved when comparing laparoscopic and robotic surgery for rectal cancer\cite{13-15,24,25}.

**Margins**

Rectal cancer specimen margins assessed are circumferential radial (CRM), proximal, and distal. Ensuring negative margins is of utmost importance in reducing local recurrence rates. Margin assessment is used as a marker to examine and compare surgical techniques. The literature on robotic vs. laparoscopic rectal resection for each margin status is discussed below.

Circumferential Radial Margin: The largest RCT to date on robotic vs. laparoscopic resection for rectal cancer is the ROLARR trial\cite{10}. In total, 237 patients were randomized to robotic, of whom the CRM
status was available for 235, and 234 to laparoscopic with CRM status available for 224 patients. The CRM positivity rate was 6.3% in laparoscopic vs. 5.1% in the robotic group ($P = 0.56$). Kim et al.\textsuperscript{[11]} also found similar CRM positivity rates in their RCT with no difference in robotic (6.1%) compared to laparoscopic (5.5%) ($P = 0.999$). Eltair et al.\textsuperscript{[13]} also confirmed no difference in positive CRM in their pooled analysis of three RCTs in their meta-analysis, but high heterogeneity was noted ($I^2 = 57\%$). Several meta-analyses that included retrospective studies along with the available RCTs have also shown no difference in positive CRM\textsuperscript{[14,24-26]}.

Proximal Resection Margin: Adequate mobilization of the colon, including splenic flexure mobilization, should allow for sufficient proximal resection margins in rectal cancer surgery. The advantages of laparoscopic and robotic rectal cancer resection with this regard pertain to the smaller incisions required for sufficient mobilization compared to open surgery. The three RCTs examined in this review by Jayne et al.\textsuperscript{[10]}, Kim et al.\textsuperscript{[11]}, and Baik et al.\textsuperscript{[12]} reported no difference in proximal resection margins when comparing robotic to laparoscopic rectal cancer operations. None of the meta-analyses examined in this review reported a difference in proximal margins\textsuperscript{[13,14,27,28]}.

Distal Resection Margin: The ROLARR RCT did not compare length of distal margin between the two surgical groups but did note one patient had a positive distal margin in the laparoscopic group\textsuperscript{[16]}. Kim et al.\textsuperscript{[11]} reported median distal resection margins and noted no statistical difference between robotic (1.5 cm) and laparoscopic (0.7 cm) ($P = 0.11$). Baik et al.\textsuperscript{[12]} also noted no difference in mean or median distal resection margins in their groups ($P = 0.467$). Eltair et al.\textsuperscript{[13]} examined five RCTs, which included 455 patients, in their meta-analysis for distal resection margins and found slightly longer distal margins in the robotic group compared to the laparoscopic one with a mean difference of 0.8 cm ($P = 0.004$). There was significantly high heterogeneity ($I^2 = 75\%$) observed in this pooled analysis. A meta-analysis by Liao et al.\textsuperscript{[27]} included five RCTs, with 340 patients, and also found longer distal margin in the robotic group compared to the laparoscopic one ($P = 0.003$), but again high heterogeneity was noted ($I^2 = 75\%$). Simillis et al.\textsuperscript{[14]} also found the robotic surgical approach to have higher distal resection margins when compared to open (7.6 mm), laparoscopic (6.8 mm), and transanal (6.8 mm) techniques. There were no reported data on positive distal margins for any of these groups.

CONCLUSION

Introduction of new surgical techniques to further surgical innovation and improve patient outcomes should be judiciously undertaken to ensure patient care, most notably that oncologic outcomes are not compromised. The majority of the high-level available evidence has found no differences between the two surgical approaches relative to TME completeness, lymph node harvest, positive CRM, or proximal resection margin. A longer distal resection margin has been found with robotic compared to laparoscopic approaches in meta-analyses, but not in RCTs. However, there is no evidence that a longer distal margin translates to better oncological outcomes.

Based on the current literature, either approach, laparoscopic or robotic, is safe and effective from a pathology standpoint. Since the two techniques are comparable, other outcomes and factors need to be considered when recommending one versus the other to our patients. The non-pathology outcomes are discussed in a separate review and should be strongly considered\textsuperscript{[29]}.

Scrutinizing one’s own rectal cancer resection outcomes is even more important than reviewing the literature. The Commission on Cancer’s National Accreditation Program for Rectal Cancer was established to ensure the highest quality metrics based on the highest level of evidence available are followed\textsuperscript{[30]}. The NAPRC requires data collection and monitoring, which should help the provision of optimal care. A national program of this caliber allows for further tracking of current care processes to better evaluate.
the care model to continue improving patient care. Regardless of what surgical technique is chosen by the surgeon, a multidisciplinary team approach must be applied to optimize oncologic outcomes.[31,32].

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