

# Local excision of scars after a complete clinical response to neoadjuvant CRT in rectal cancer—organ-preservation without function-preservation?

# Adrian Mattacheo, Rodrigo Oliva Perez

Angelita & Joaquim Gama Institute, Rua Manuel da Nóbrega 1564, São Paulo, Brazil

*Correspondence to*: Rodrigo Oliva Perez, MD, PhD. Angelita & Joaquim Gama Institute, Rua Manuel da Nóbrega 1564, São Paulo, 04001-005, Brazil. Email: rodrigo.operez@gmail.com.

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Management of rectal cancer has changed significantly over the last few decades (1). First, appropriateness of surgical technique with standardization of total mesorectal excision led to a substantial decrease in local recurrence rates (2). Second, the role of neoadjuvant chemoradiotherapy (nCRT) in further decreasing local failures after TME is currently well established (3). Finally, with optimal local disease control, the concept of organ-preservation has become an attractive alternative to patients with significant primary tumor regression after nCRT. In this setting, selecting patients with complete clinical response (cCR) could be managed by no immediate surgery or transanal local excision providing not only excellent oncological outcomes but also acceptable functional results (4-7).

Ghiselli *et al.* have investigated the functional outcomes of patients that have developed a cCR and were managed by routine transanal endoscopic microsurgery (TEM) (8). The data suggests minimal postoperative complication rates and excellent functional outcomes leading the authors to conclude that such approach could be considered the preferred organ-preserving treatment strategy for patients with cCR after nCRT for patients with rectal cancer. In fact, perhaps the most appropriate tool to assess functional outcomes among these patients would have been the low anterior resection syndrome score recently validated in multiple languages instead of the fecal incontinence scores and questionnaires used in the study (9).

Nevertheless, there are a few additional considerations that should be considered prior to full implementation of this approach into clinical practice.

First of all, assessment of patients after 30 days from completion of nCRT may have significantly affected the outcomes of the study. Not only this relatively short interval may have underestimated the proportion of patients with cCR but also significantly underestimated functional outcomes at "baseline". Acute effects of radiation after 30 days may have worsened functional outcomes at baseline leading to less striking differences with postoperative assessment results at 1 year (10).

Second, even though a significant number of patients were entered into the study, the absence of sample size calculation due to its retrospective design may also represent a significant source of bias. Ultimately, it is likely that this number of patients led insufficient power of the study to demonstrate any clinically relevant differences.

Third, the inclusion of patients with considerably highlocated tumors (70% were beyond 5 cm from the anal verge) is relevant here. This is because organ-preservation is more critical for most distal rectal tumors. Preservation of the rectum to these patients may provide a more significant benefit than for patients with more proximal lesions, where anterior resections may provide acceptable functional results. Local excision in this latter group of patients is unlikely to require any amount of sphincteric resection and therefore result in minimal functional consequences (11). It would have been perhaps more meaningful to provide a comparison of functional outcomes of these two organpreserving strategies among patients that otherwise would

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have required ultra-low anterior resections or abdominal perineal excisions. In this case, local excision would probably require at least partial resection of sphincter complex leading to potentially worse functional outcomes.

Finally, when comparing non-operative management to transanal local excision as organ-preserving strategies for patients with rectal cancer following nCRT, one has to consider the consequences of each alternative to regular follow-up. In both cases, thorough follow-up of the preserved rectum is crucial to provide early detection of local recurrences and appropriate salvage resection in this situation (12). Here non-operative management after a cCR may provide minimal distortion of the anatomy of the rectum seen by endoscopic or radiological assessment. In contrast, local excision and significant postoperative scarring may represent a significant challenge for endoscopic and radiological assessment during post-operative follow-up (13). These difficulties may result in clinically relevant consequences oncological outcomes after a local recurrence (14).

Altogether, even though the present study may provide data suggesting the advantages of local excision for the management of cCR over observation alone, a few significant limitations of the study may ultimately have underestimated the proportion of patients that worsened their anorectal function after a local excision. Considering that nearly one out of five patients became incontinent after TEM, local excision of a fibrotic scar (with not a single cancer cell in the resected specimen) may ultimately be already significantly deleterious in the absence of no oncological benefit to close surveillance alone with no immediate resection.

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# References

- Kosinski L, Habr-Gama A, Ludwig K, et al. Shifting concepts in rectal cancer management: a review of contemporary primary rectal cancer treatment strategies. CA Cancer J Clin 2012;62:173-202.
- Heald RJ, Husband EM, Ryall RD. The mesorectum in rectal cancer surgery--the clue to pelvic recurrence? Br J Surg 1982;69:613-6.
- Sauer R, Becker H, Hohenberger W, et al. Preoperative versus postoperative chemoradiotherapy for rectal cancer. N Engl J Med 2004;351:1731-40.
- 4. Habr-Gama A, Perez RO, Nadalin W, et al. Operative versus nonoperative treatment for stage 0 distal rectal cancer following chemoradiation therapy: long-term results. Ann Surg 2004;240:711-7; discussion 7-8.
- Garcia-Aguilar J, Renfro LA, Chow OS, et al. Organ preservation for clinical T2N0 distal rectal cancer using neoadjuvant chemoradiotherapy and local excision (ACOSOG Z6041): results of an open-label, singlearm, multi-institutional, phase 2 trial. Lancet Oncol 2015;16:1537-46.
- Rullier E, Rouanet P, Tuech JJ, et al. Organ preservation for rectal cancer (GRECCAR 2): a prospective, randomised, open-label, multicentre, phase 3 trial. Lancet 2017;390:469-79.
- Dossa F, Chesney TR, Acuna SA, et al. A watch-andwait approach for locally advanced rectal cancer after a clinical complete response following neoadjuvant chemoradiation: a systematic review and meta-analysis. Lancet Gastroenterol Hepatol 2017;2:501-13.
- 8. Ghiselli R, Ortenzi M, Cardinali L, et al. Functional

outcomes after TEM in patients with complete clinical response after neoadjuvant chemoradiotherapy. Surg Endosc 2017;31:2997-3003.

- Juul T, Ahlberg M, Biondo S, et al. Low anterior resection syndrome and quality of life: an international multicenter study. Dis Colon Rectum 2014;57:585-91.
- Hayne D, Vaizey CJ, Boulos PB. Anorectal injury following pelvic radiotherapy. Br J Surg 2001;88:1037-48.
- Habr-Gama A, Lynn PB, Jorge JM, et al. Impact of Organ-Preserving Strategies on Anorectal Function in Patients with Distal Rectal Cancer Following Neoadjuvant Chemoradiation. Dis Colon Rectum 2016;59:264-9.
- 12. Habr-Gama A, Gama-Rodrigues J, Sao Juliao GP, et

### doi: 10.21037/ales.2018.01.02

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al. Local recurrence after complete clinical response and watch and wait in rectal cancer after neoadjuvant chemoradiation: impact of salvage therapy on local disease control. Int J Radiat Oncol Biol Phys 2014;88:822-8.

- Sao Juliao GP, Ortega CD, Vailati BB, et al. Magnetic resonance imaging following neoadjuvant chemoradiation and transanal endoscopic microsurgery for rectal cancer. Colorectal Dis 2017;19:O196-O203.
- Perez RO, Habr-Gama A, Sao Juliao GP, et al. Transanal Endoscopic Microsurgery (TEM) Following Neoadjuvant Chemoradiation for Rectal Cancer: Outcomes of Salvage Resection for Local Recurrence. Ann Surg Oncol 2016;23:1143-8.