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# A multicentric randomized controlled trial on the impact of lengthening the interval between neoadjuvant radiochemotherapy and surgery on complete pathological response in rectal cancer (GRECCAR-6 trial): rationale and design

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## Abstract

**Background:** Neoadjuvant radiochemotherapy (RCT) is now part of the armamentarium of cancer of the lower and middle rectum. It is recommended in current clinical practice prior to surgical excision if the lesion is classified T3/T4 or N+. Histological complete response, defined by the absence of persistent tumor cell invasion and lymph node (ypT0N0) after pathological examination of surgical specimen has been shown to be an independent prognostic factor of overall survival and disease-free survival. Surgical excision is usually performed between 6 and 8 weeks after completion of CRT and pathological complete response rate ranges around 12%. In retrospective studies, a lengthening of the interval after RCT beyond 10 weeks was found as an independent factor increasing the rate of pathological complete response (between 26% and 31%), with a longer disease-free survival and without increasing the operative morbidity. The aim of the present study is to evaluate in 264 patients the rate of pathological complete response rate of rectal cancer after RCT by lengthening the time between RCT and surgery.

**Methods/design:** The current study is a multicenter randomized trial in two parallel groups comparing 7 and 11 weeks of delay between the end of RCT and cancer surgery of rectal tumors. At the end of the RCT, surgery is planned and randomization is performed after patient's written consent for participation. The histological complete response (ypT0N0) will be determined with analysis of the complete residual tumor and double reading by two pathologists blinded of the group of inclusion. Patients will be followed in clinics for 5 years after surgery. Participation in this trial does not change patient's management in terms of treatment, investigations or visits. Secondary endpoints will include overall and disease free survival, rate of sphincter conservation and quality of mesorectal excision. The number of patients needed is 264.

**Trial registration:** ClinicalTrials.gov: NCT01648894

**Keywords:** Rectal cancer, Radiochemotherapy, Complete histological response, Procedure

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## Background

Rectal cancer represents 12 000 new cases each year in France [1]. Its management is based on a multidisciplinary management with radiotherapy, chemotherapy and surgery.

### Surgery

Surgery consisting of total excision of the mesorectum has improved over time. The studies of Heald [2,3] have showed that local recurrence rate was directly related to the preservation of the fascia recti during the dissection of the rectum. In countries where this surgical procedure was emphasized, the local recurrence rate and life expectancy after treatment of rectal cancer were improved [4,5]. In Sweden, after learning of the mesorectal excision technique, 447 patients were compared to older cohorts, the local recurrence rate at 5 years was increased from 20.5% to 8.2% and 5-year survival of 65.8% to 77.3% ( $p < 0.001$  for both comparisons).

### Radiotherapy

Before the widespread diffusion of the technique of mesorectal excision, many studies, including 4 randomized studies, had compared pre-operative radiotherapy (RT) versus surgery alone [6-9]. Results showed only a benefit in terms of local control with a reduction of 10% to 5% of the recurrence rate local for patients with optimal dissection of the mesorectum, but no survival benefit. In 1997, a randomized study in 1168 subjects showed an improved survival with neoadjuvant RT: survival at 9 years was 65% for patients with no radiotherapy compared to 74% for patients with neoadjuvant RT ( $p = 0.002$ ) [9]. Local recurrence for patients treated by surgery alone was also more frequent (27% (150/557) compared to 11% (63/553),  $p < 0.001$ ) [9]. This finding was explained by a suboptimal surgical resection without complete mesorectum excision. As a consequence, an international randomized trial study comparing RT + surgery versus surgery alone with quality control of excision mesorectum [10] was conducted in 1805 patients. The results confirmed a higher rate of local control at 2 years in patients with preoperative radiotherapy (2.4% vs. 8.3%,  $p < 0.001$ ), with no difference in terms of survival (82% vs. 81.8%,  $p = 0.84$ ). Early postoperative complications were higher in the RT + surgery group (48% vs. 41%,  $p < 0.01$ ) [11]. After a minimum of 5 years of follow-up, long-term complications; were more frequent in the preoperative radiotherapy group with (62% and 56% versus 38% and 33% of episodes of incontinence and pad wearing respectively;  $p < 0.001$ ) [12]. As postoperative morbidity and results functional obtained after preoperative RT were very significantly altered, the authors recommended such treatment only in patients at high risk of local recurrence. These recommendations were proposed by the French

experts for the treatment of rectal cancer and were approved by the *Haute Autorité de Santé* in 2007 [13].

### Chemoradiotherapy

Other studies have evaluated the impact of adding chemotherapy to RT [14,15]. The randomized trial of Bosset et al. included between 1993 and 2003, 1011 patients with rectal cancer T3 or T4 and analysed the impact of addition of chemotherapy based on 5-fluorouracil to RT 45Gy [14] compared with preoperative RT alone. The complete response rate (pT0) on resected specimen was 5.3% in the RT alone group and 13.7% in the chemoradiotherapy (CRT) group (odds ratio = 2.84, 95%CI: 1.75 to 4.59,  $p < 0.0001$ ) [15]. No benefit in overall survival or disease-free survival was observed in CRT group despite a significant improvement rate of local recurrence at 5 years (8.7% in the CRT compared to 17.1% after RT alone vs.,  $p = 0.002$ ). However, the results cannot be generalized because of the lack of uniform total mesorectal excision for all patients included in this study.

However, CRT has allowed the team of Pr. Habr-Gama to obtain a complete clinical regression of tumor rectal in 26.8% of cases (71/265 patients) [16]. The protocol consisted of 50 Gy radiation therapy combined with chemotherapy based on 5-fluorouracil and leucovorin. This group of patients had not been operated and simple surveillance without resection was performed. After a mean follow-up of 57 months, only 2 patients (2.8%) had local recurrence and 3 had distant metastases. The 5-year recurrence rate was 7%. The complete clinical response was associated with better overall survival at 5 years: 100% versus 88% ( $p = 0.01$ ).

In a retrospective series of the Cleveland Clinic, among 238 patients treated with neoadjuvant CRT, 58 (24.4%) had a ypT0N0 tumor on pathological exam. Postoperative morbidity was similar in the neoadjuvant CRT group and in the no-pCR group, but there was a better local control (5 years local recurrence rate: 0% vs. 10.6%,  $p < 0.001$ ) [17].

### Complete response: the importance of the period between the end of radiotherapy and surgery

The median complete response rate is about 12% [7-27%] in the main series of rectal cancer treated with RT or RCT [18-31]. A randomized study in 1999 had already found an effect in tumor reduction after RT between two intervals (2 weeks or 7 weeks) clinical response increased from 53.1% to 71.7% ( $p = 0.007$ ) and pCR or near complete (persistence of some tumor cells) increased from 10.3% to 26% ( $p = 0.0054$ ) [21]. The results of this study cannot be generalized, considering the use of a technique of mesorectal excision other than the widely distributed and administered neoadjuvant therapy (RT alone of 40 Gy). However the interest of extending the period after RT was

obvious. In 2003, Moore's team showed that among 155 rectal cancers treated with neo-adjuvant RCT (50 Gy + 5-fluorouracil), the rate of complete response increased from 9% to 23% between patients operated on before the 40<sup>th</sup> day (between 6<sup>th</sup> and 7<sup>th</sup> week) and those operated on after waiting more than 7 weeks ( $p = 0.09$ ) [27].

Following the work of Prof. Habr-Gama, other authors have attempted to identify factors associated with a pCR after surgical resection. The first study published by Tulchinsky et al. [29] in 2008 reported the retrospective findings on the time between preoperative CRT (45–50 Gy + 5-fluorouracil) and surgery (less or more than 7 weeks) in 132 patients operated on by anterior resection or abdomino-perineal resection between 2000 et 2006. The rate of pCR or near complete (persistence of microscopic foci of adenocarcinoma in the rectal wall without lymph node) was 28% in the resected specimen. The single independent factor associated with a good response was the period between the end of RT and surgery: 17% in the group operated <7 weeks against 35% in the other group operated after 7 weeks ( $p = 0.03$ ). There was no association between the duration of interval before surgery and postoperative morbidity (complications, transfusion, duration of hospitalization). Kalady et al. studied retrospectively (1997–2007) records of 306 patients operated for a rectal cancer after CRT (50 Gy and 5-fluorouracil) [30]. The dose of radiation received in each group was similar. The rate of pCR (ypT0N0) was 24% in this study. Time between the end of RCT and surgery, with a cut-off estimated at 8 weeks was the single prognostic factor for pCR in uni- and multivariate analysis (OR = 2.63-<sub>95%</sub>CI [1.13 to 6.12],  $p = 0.02$ ). A. The pathological complete response rate increased from 16.3% (14/86) in the group operated < 8 weeks against 30.8% (28/91) for other ( $p = 0.03$ ). The authors observed that the period between 8 and 10 weeks showed the greatest number of complete responses and that there was no more gain over 14 weeks. This retrospective study did not give any explanation on the reasons for variations in the time between the end of radiotherapy and surgery [32]. The group of patients with pCR had a better overall survival with a follow-up of 60 months (91% versus 80%,  $p = 0.046$ ) and less local recurrence at 5 years (0% versus 11%,  $p = 0.023$ ). A Korean study analyzed retrospectively the data from 12 centers of 306 patients with a tumor classified ypT0 following CRT. This study confirmed the favourable impact of pCR after neoadjuvant CRT on survival: the 5-year overall survival of these patients having 93.4% of tumors initially classified T3 or T4 was 92.8%. Disease-free survival at 5 years was 84.6%. Finally, Kalady et al. compared the outcomes in a retrospective series of 177 patients operated for a rectal cancer after neoadjuvant treatment between patients operated before 8 weeks or after 8 weeks following the end of CRT between 1997 and 2007. Postoperative morbidity or

mortality were similar between the two period group. The rate of pCR was lower in the <8 weeks group compared with the >8 weeks group (16.2% vs. 31.1%,  $p = 0.027$ ). Moreover, the 3 years local recurrence rate was significant lower in the >8 weeks (1.2% vs. 3.9%,  $p = 0.04$ ) [33].

All these publications are retrospective and the real impact of the delay between the end of CRT and surgery is still a matter of debate [34].

### Aims

The main objective of our study is to evaluate in a randomized trial the impact of a longer interval between the end of CRT and rectal cancer surgery (7 weeks versus 11 weeks) on the rate of pathological complete response. Secondary outcomes include overall and disease free survival, quality of mesorectal excision, rate of sphincter preservation.

### Methods and design

This study is a multicenter randomized open-label controlled trial in parallel groups, comparing two periods between the end of CRT and cancer surgery of rectal tumors: 7 weeks versus 11 weeks. This study has been approved by a national Institutional Review Board: the Regional Comity of Patients Protection of South-West I, N°1-12-19:30/08/12 and by the National Agency of Medicine and Medical Products (ANSM: B111580-10). This study is supported by a grant from the French Ministry of Health (PHRCN 2011, AOM 11314). The research carried out will be on accordance with Helsinki declaration.

### Participants

The institutional promoter is the AP-HP (*Assistance Publique-Hôpitaux de Paris – DRCD: Département de la Recherche Clinique*). Patients are included from several departments of surgery or oncology ( $n=26$ ) in France (see list of participating centers in the Acknowledgments section). All participating sites signed a convention with the DRCD for ethical approval before beginning of inclusion. All patients must fulfil the following criteria: T3/T4 and/or TxN+ mid or low third rectal cancer and completed RCT. The complete inclusion and exclusion criteria are given in Table 1. After oral and written explanation about the purpose of this study, the patient gives his written consent agreeing to participate to the protocol (Figure 1).

### Randomization

After completion of the pre-intervention assessments, the patients are randomly assigned to the period group (ratio 1:1) by Internet. Blocked centralized randomisation with stratification by centre will be prepared by URC-Est.

**Table 1 Inclusion and exclusion criteria**

Inclusion criteria	Exclusion criteria
<ul style="list-style-type: none"> <li>• age over 18 years, no age limit higher</li> <li>• Performance status evaluated by the Eastern Cooperative Oncology Group (ECOG) score: 0-1</li> <li>• Patients with cancer of the middle or lower rectum (lesion located within 10 cm from the dentate line or 12 cm from the anal margin) proved by pathology,</li> <li>• T3-T4N0, TxN+ on ultrasound-endoscopy and MRI, without secondary localization (M0) on the thoraco-abdominal (or chest radiography and abdominal ultrasound)</li> <li>• Patient who received a protocol between 45–50 Gy of radiotherapy and chemotherapy based on 5-fluorouracil for an average duration of 5 weeks for the management of rectal cancer,</li> <li>• Curative surgical treatment planned following radiochemotherapy with total mesorectal excision,</li> <li>• Free and informed consent signed by the patient,</li> <li>• Patient affiliated to a social security scheme or beneficiary of such plan (except AME)</li> <li>• Patient able, according to the investigator, to comply with the requirements of the study.</li> </ul>	<ul style="list-style-type: none"> <li>• Patient with metastasis,</li> <li>• T1 or T2N0 tumor classified by echo -endoscopy and MRI,</li> <li>• rectal tumor with lower pole is more than 12 cm from the anal margin or 10 cm from the dentate line,</li> <li>• Patient did not complete the full protocol of radiotherapy,</li> <li>• History of tumors (other than basal cell carcinoma and / or carcinoma in situ of the cervix)</li> <li>• A patient with impaired or incompetent</li> <li>• investigator by not allowing him a good understanding of the requirements of the study, person under guardianship, persons under guardianship, persons deprived of their liberty by judicial or administrative body, adult subject to legal protection or unable to consent.</li> </ul>

## Intervention

### ***Surgical resection with total mesorectal excision (TME)***

The anaesthesia consultation is planned before the surgery according to the habits of each department. Participation in the study does not alter the anaesthetic procedures. The patient is admitted the day before surgery in the surgical ward.

The type of resection (coloanal anastomosis, abdomino-perineal resection, delayed anastomosis, drainage, laparoscopic approach...) is not influenced by the participation to this protocol. As the rectal tumor is located in the mid- or low rectum a TME is required.

### **Pathological exam**

The pathological exam required has been already published in the National French Guidelines [35]. A standardized routine pathology examination was performed using the protocol of Quirke et al. [36]. After fixation in 10% formalin and inking to assess the circumferential margin, the whole tumor was cut transversely. According to the macroscopical features, different techniques for sampling of tumor tissues were applied in order to avoid any under staging of the specimen:

- If there was only ulceration and fibrotic changes without any visible tumoral lesion or if the tumor residual measured less than 3 cm in diameter, the lesion was examined entirely;
- If the residual tumor measured more than 3 cm in diameter, one selected block per cm of tumor was

processed. If the first selected blocks were free of tumor, complementary blocks were taken and the macroscopically residual lesion was examined in toto. A diagnosis of pCR will only be made after examination of the whole macroscopically residual lesion [37]. The tumor response is evaluated by inclusion of all residual tumor and the response to CRT is graded with the scales of Rodol and Dvorak [22,38]. A double reading of slides will be made for each patient by two independent pathologists blinded to the randomization group of the patient to confirm the ypT stage as the two regression scores. In case of disagreement between the two pathologists, they should jointly give a mutual result.

## Outcomes and assessments

### ***Primary outcome***

Rate of histological complete response after double reading by two different pathologist.

### ***Secondary outcomes***

Morbidity, sphincter preservation rate, overall and disease-free survival.

### ***Surgical data***

During surgery, the operating data are provided on the e-CRF (digital rectal examination under general anaesthesia, type of surgery (anterior resection or abdomino-perineal resection), operative time, intraoperative bleeding, macroscopic

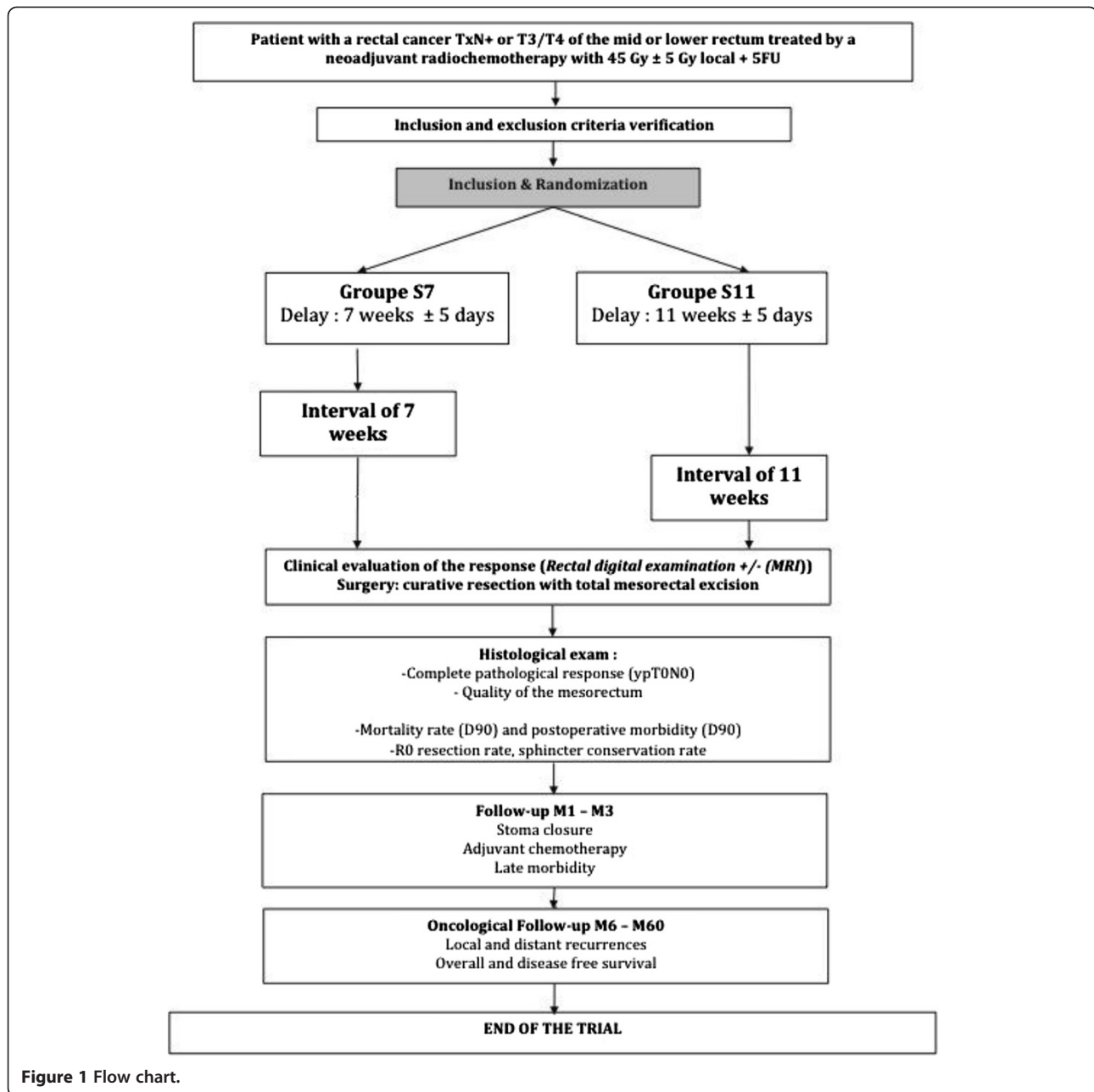


Figure 1 Flow chart.

appearance of the mesorectum, distance from the distal limit of resection.

#### Morbidity and mortality

The postoperative complications are noted by the surgeon in the e-CRF during hospitalization (about 10–15 days) and during the first 3 months. Postoperative death is defined as death occurring within 30 postoperative days or during the first hospitalization. Postoperative complications are defined by the occurrence of medical or surgical complications within 90 postoperative days

or during the first hospitalization. Morbidity will be evaluated with the new classification of surgical complications by Dindo et al. which includes 5 grades [39,40].

#### Pathological exam

Usual data are recorded: distal and circumferential margins, number of resected and invaded nodes, tumoral differentiation, presence of vascular embols (veinous or lymphatic, intra or extra-mural), perineural engainment, quality of mesorectal excision. The resected specimen will

be staged according to American Joint Committee on Cancer (AJCC) criteria (7<sup>th</sup> version).

#### **Rate of sphincter preservation**

Comparison between the planned intervention at the time of the randomisation and the resection performed after peroperative digital exam will be performed.

#### **Oncological follow-up**

Patients will be followed in clinics for 5 years according to the habits of each department. Usual follow-up is composed of clinical exam, CEA analysis, CT-scan or Chest Radiography with abdominal ultra-sound every 3–4 months during the first three years and every 6 months for the last two years.

#### **Samples size and statistical considerations**

With a sample of 264 patients, GRECCAR6 trial has 80% of power to detect at least a two fold increase in the complete response rate in the 11 weeks group compared to the 7 weeks group. This hypothesis assumes that the complete response rate in the 7 weeks group is similar to the usual rate of complete response after 6–8 weeks (12% (Kalady et al. Bosset et al. Hiotis et al.)), with an expected complete response rate in the 11 weeks group of 26% and 10% of drop-outs, using a two-sided test at the 0.05 significance level.

Intention to treat analysis of the primary endpoint will be performed once all randomized patients have 6 months of follow-up. Analysis of survival will be performed after at least 5 years of follow-up. A futility analysis [41] is planned after randomization after 132 patients, using a Bayesian statistical interference, with no impact on Type 1 error [42].

#### **Discussion**

While several retrospective studies has emphasised the role of longer interval on efficiency of radiochemotherapy on histological tumoral response, this multicentric randomized trial is the first to evaluate this factor. The major drawback of previous studies is that no explanations are given on the reasons for variations in the time between the end of radiotherapy and surgery [29,32,43]. Indeed, the time interval was decided by the surgeon or maybe in case of favourable clinical response, mainly authors advocate other factors such as patient morbidity or logistical scheduling issues [33].

This trial could then identify a simple and cheap factor influencing the rate of pCR. While is it still not obvious that pCR is a good prognostic marker of the overall survival [44] based on previous randomized control trial on CRT, it is probable that it could be a prognosis marker of rectal conservation. One publication showed that

patients operated after 8 weeks may have significantly less local recurrences.

Indeed, if waiting 4 more weeks increase the rate of pCR, more patients could avoid a morbid surgical procedure such as an abdomino-perineal resection. This new management is still limited due to the actual difficulties to identify patients with pCR before surgery. Advances in radiological imaging may facilitate this important point.

Finally, this long delay would allow the use of chemotherapy during this 3 months waiting period. This could reduce the risk of synchronous metastasis that may occur during the waiting period.

#### **Competing interests**

The authors declare that they have no competing interests.

#### **Authors' contributions**

JHL: conception of the study, writing of the manuscript. AR: statistical analysis and help to draft the manuscript. MS: pathological exam and help to draft the manuscript. YP: writing of the manuscript. TS: coordination of the study. ET: conception and coordination of the study. All authors read and approved the manuscript.

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#### References

1. Bouvier AM, Remontet L, Jouglu E: Incidence of gastrointestinal cancers in France. *Gastroenterol Clin Biol* 2004, **28**:877-881.
2. Heald RJ, Ryall RD: Recurrence and survival after total mesorectal excision for rectal cancer. *Lancet* 1986, **1**(8496):1479-1482.
3. MacFarlane JK, Ryall RD, Heald RJ: Mesorectal excision for rectal cancer. *Lancet* 1993, **341**(8843):457-460.
4. Martling A, Cedermark B, Johansson H, Rutqvist LE, Holm T: The surgeon as a prognostic factor after the introduction of total mesorectal excision in the treatment of rectal cancer. *Br J Surg* 2002, **89**(8):1008-1013.
5. Martling A, Holm T, Rutqvist LE, Johansson H, Moran BJ, Heald RJ, Cedermark B: Impact of a surgical training programme on rectal cancer outcomes in Stockholm. *Br J Surg* 2005, **92**(2):225-229.
6. Gerard A, Berrod JL, Pene F, Loygue J, Laugier A, Bruckner R, Camelot G, Arnaud JP, Metzger U, Buyse M, et al: Preoperative radiotherapy and radical surgery as combined treatment in rectal cancer. *Recent Results Cancer Res* 1988, **110**:130-133.
7. Stockholm Rectal Cancer Study Group: Preoperative short-term radiation therapy in operable rectal carcinoma. A prospective randomized trial. *Cancer* 1990, **66**(1):49-55.
8. Randomised trial of surgery alone versus radiotherapy followed by surgery for potentially operable locally advanced rectal cancer. Medical research council rectal cancer working party. *Lancet* 1996, **348**(9042):1605-1610.
9. Improved survival with preoperative radiotherapy in resectable rectal cancer. Swedish rectal cancer trial. *N Engl J Med* 1997, **336**(14):980-987.
10. Kapiteijn E, Marijnen CA, Nagtegaal ID, Putter H, Steup WH, Wiggers T, Rutten HJ, Pahlman L, Glimelius B, van Krieken JH, et al: Preoperative radiotherapy combined with total mesorectal excision for resectable rectal cancer. *N Engl J Med* 2001, **345**(9):638-646.
11. Marijnen CA, Kapiteijn E, van de Velde CJ, Martijn H, Steup WH, Wiggers T, Kranenborg EK, Leer JW: Acute side effects and complications after short-term preoperative radiotherapy combined with total mesorectal excision in primary rectal cancer: report of a multicenter randomized trial. *J Clin Oncol* 2002, **20**(3):817-825.
12. Peeters KC, van de Velde CJ, Leer JW, Martijn H, Junggeburst JM, Kranenborg EK, Steup WH, Wiggers T, Rutten HJ, Marijnen CA: Late side effects of short-course preoperative radiotherapy combined with total mesorectal excision for rectal cancer: increased bowel dysfunction in irradiated patients—a Dutch colorectal cancer group study. *J Clin Oncol* 2005, **23**(25):6199-6206.
13. Portier G: [Recommendations for clinical practice. Therapeutic choices for rectal cancer. How should neoadjuvant therapies be chosen?]. *Gastroenterol Clin Biol* 2007, **31** Spec No 1:23-33.
14. Bosset JF, Collette L, Calais G, Mineur L, Maingon P, Radosevic-Jelic L, Daban A, Bardet E, Beny A, Ollier JC: Chemotherapy with preoperative radiotherapy in rectal cancer. *N Engl J Med* 2006, **355**(11):1114-1123.
15. Bosset JF, Calais G, Mineur L, Maingon P, Radosevic-Jelic L, Daban A, Bardet E, Beny A, Briffaux A, Collette L: Enhanced tumorocidal effect of chemotherapy with preoperative radiotherapy for rectal cancer: preliminary results—EORTC 22921. *J Clin Oncol* 2005, **23**(24):5620-5627.
16. Habr-Gama A, Perez RO, Nadalin W, Sabbaga J, Ribeiro U Jr, Silva e Sousa AH Jr, Campos FG, Kiss DR, Gama-Rodrigues J Jr: Operative versus nonoperative treatment for stage 0 distal rectal cancer following chemoradiation therapy: long-term results. *Ann Surg* 2004, **240**(4):711-717.
17. de Campos-Lobato LF, Stocchi L, da Luz MA, Geisler D, Dietz DW, Lavery IC, Fazio VW, Kalady MF: Pathologic complete response after neoadjuvant treatment for rectal cancer decreases distant recurrence and could eradicate local recurrence. *Ann Surg Oncol* 2011, **18**(6):1590-1598.
18. Medich D, McGinty J, Parda D, Karlovits S, Davis C, Caushaj P, Lembersky B: Preoperative chemoradiotherapy and radical surgery for locally

- advanced distal rectal adenocarcinoma: pathologic findings and clinical implications. *Dis Colon Rectum* 2001, **44**(8):1123-1128.
19. Kim NK, Baik SH, Seong JS, Kim H, Roh JK, Lee KY, Sohn SK, Cho CH: Oncologic outcomes after neoadjuvant chemoradiation followed by curative resection with tumor-specific mesorectal excision for fixed locally advanced rectal cancer: impact of postirradiated pathologic downstaging on local recurrence and survival. *Ann Surg* 2006, **244**(6):1024-1030.
  20. Hiotis SP, Weber SM, Cohen AM, Minsky BD, Paty PB, Guillem JG, Wagman R, Saltz LB, Wong WD: Assessing the predictive value of clinical complete response to neoadjuvant therapy for rectal cancer: an analysis of 488 patients. *J Am Coll Surg* 2002, **194**(2):131-135. discussion 135-136.
  21. Francois Y, Nemoz CJ, Baulieux J, Vignal J, Grandjean JP, Partensky C, Souquet JC, Adeleine P, Gerard JP: Influence of the interval between preoperative radiation therapy and surgery on downstaging and on the rate of sphincter-sparing surgery for rectal cancer: the Lyon R90-01 randomized trial. *J Clin Oncol* 1999, **17**(8):2396.
  22. Rodel C, Martus P, Papadopoulos T, Fuzesi L, Klimpfinger M, Fietkau R, Liersch T, Hohenberger W, Raab R, Sauer R, et al: Prognostic significance of tumor regression after preoperative chemoradiotherapy for rectal cancer. *J Clin Oncol* 2005, **23**(34):8688-8696.
  23. Gerard JP, Conroy T, Bonnetain F, Bouche O, Closon-Dejardin MT, Untereiner M, Leduc B, Francois E, Maurel J, et al: Preoperative radiotherapy with or without concurrent fluorouracil and leucovorin in T3-4 rectal cancers: results of FFD 9203. *J Clin Oncol* 2006, **24**(28):4620-4625.
  24. Stipa F, Chessin DB, Shia J, Paty PB, Weiser M, Temple LK, Minsky BD, Wong WD, Guillem JG: A pathologic complete response of rectal cancer to preoperative combined-modality therapy results in improved oncological outcome compared with those who achieve no downstaging on the basis of preoperative endorectal ultrasonography. *Ann Surg Oncol* 2006, **13**(8):1047-1053.
  25. Garcia-Aguilar J, Hernandez de Anda E, Sirivongs P, Lee SH, Madoff RD, Rothenberger DA: A pathologic complete response to preoperative chemoradiation is associated with lower local recurrence and improved survival in rectal cancer patients treated by mesorectal excision. *Dis Colon Rectum* 2003, **46**(3):298-304.
  26. Ruo L, Tickoo S, Klimstra DS, Minsky BD, Saltz L, Mazumdar M, Paty PB, Wong WD, Larson SM, Cohen AM, et al: Long-term prognostic significance of extent of rectal cancer response to preoperative radiation and chemotherapy. *Ann Surg* 2002, **236**(1):75-81.
  27. Moore HG, Gittleman AE, Minsky BD, Wong D, Paty PB, Weiser M, Temple L, Saltz L, Shia J, Guillem JG: Rate of pathologic complete response with increased interval between preoperative combined modality therapy and rectal cancer resection. *Dis Colon Rectum* 2004, **47**(3):279-286.
  28. Stein DE, Mahmoud NN, Anne PR, Rose DG, Isenberg GA, Goldstein SD, Mitchell E, Fry RD: Longer time interval between completion of neoadjuvant chemoradiation and surgical resection does not improve downstaging of rectal carcinoma. *Dis Colon Rectum* 2003, **46**(4):448-453.
  29. Tulchinsky H, Shmueli E, Figer A, Klausner JM, Rabau M: An interval >7 weeks between neoadjuvant therapy and surgery improves pathologic complete response and disease-free survival in patients with locally advanced rectal cancer. *Ann Surg Oncol* 2008, **15**(10):2661-2667.
  30. Kalady MF, de Campos-Lobato LF, Stocchi L, Geisler DP, Dietz D, Lavery IC, Fazio VW: Predictive factors of pathologic complete response after neoadjuvant chemoradiation for rectal cancer. *Ann Surg* 2009, **250**(4):582-589.
  31. Lim SB, Choi HS, Jeong SY, Kim DY, Jung KH, Hong YS, Chang HJ, Park JG: Optimal surgery time after preoperative chemoradiotherapy for locally advanced rectal cancers. *Ann Surg* 2008, **248**(2):243-251.
  32. Yeo SG, Kim DY, Kim TH, Chang HJ, Oh JH, Park W, Choi DH, Nam H, Kim JS, Cho MJ, et al: Pathologic complete response of primary tumor following preoperative chemoradiotherapy for locally advanced rectal cancer: long-term outcomes and prognostic significance of pathologic nodal status (KROG 09-01). *Ann Surg* 2010, **252**(6):998-1004.
  33. de Campos-Lobato LF, Geisler DP, da Luz MA, Stocchi L, Dietz D, Kalady MF: Neoadjuvant therapy for rectal cancer: the impact of longer interval between chemoradiation and surgery. *J Gastrointest Surg* 2011, **15**(3):444-450.
  34. Bujko K: Timing of surgery following preoperative therapy in rectal cancer: there is no need for a prospective randomized trial. *Dis Colon Rectum* 2012, **55**(3):31-32.
  35. Rullier A, Laurent C: [Recommendations for clinical practice. Therapeutic choices for rectal cancer. What quality criteria are important for surgical



- excision of rectal cancer?]. *Gastroenterol Clin Biol* 2007, **31 Spec No 1**:34–51. 31S91-35.
36. Quirke P, Durdey P, Dixon MF, Williams NS: **Local recurrence of rectal adenocarcinoma due to inadequate surgical resection. Histopathological study of lateral tumour spread and surgical excision.** *Lancet* 1986, **2(8514)**:996–999.
  37. Tranchart H, Lefevre JH, Svrcek M, Flejou JF, Tiret E, Parc Y: **What is the incidence of metastatic lymph node involvement after significant pathologic response of primary tumor following neoadjuvant treatment for locally advanced rectal cancer?** *Ann Surg Oncol* 2012, **20(5)**:1551–1559.
  38. Dworak O, Keilholz L, Hoffmann A: **Pathological features of rectal cancer after preoperative radiochemotherapy.** *Int J Colorectal Dis* 1997, **12(1)**:19–23.
  39. Clavien PA, Barkun J, de Oliveira ML, Vauthey JN, Dindo D, Schulick RD, de Santibanes E, Pekolj J, Slankamenac K, Bassi C, *et al*: **The Clavien-Dindo classification of surgical complications: five-year experience.** *Ann Surg* 2009, **250(2)**:187–196.
  40. Dindo D, Demartines N, Clavien PA: **Classification of surgical complications: a new proposal with evaluation in a cohort of 6336 patients and results of a survey.** *Ann Surg* 2004, **240(2)**:205–213.
  41. Armitage P: **Interim analysis in clinical trials.** *Stat Med* 1991, **10(6)**:925–935. discussion 936–927.
  42. Kramar A, Mathoulin-Pelissier S: *Methodes biostatistiques appliquees à la recherche clinique en cancérologie.* Paris: John Libbey Eurotext; 2011.
  43. Evans J, Tait D, Swift I, Pennert K, Tekkis P, Wotherspoon A, Chau I, Cunningham D, Brown G: **Timing of surgery following preoperative therapy in rectal cancer: the need for a prospective randomized trial?** *Dis Colon Rectum* 2011, **54(10)**:1251–1259.
  44. Methy N, Bedenne L, Conroy T, Bouche O, Chapet O, Ducreux M, Gerard JP, Bonnetain F: **Surrogate end points for overall survival and local control in neoadjuvant rectal cancer trials: statistical evaluation based on the FFCD 9203 trial.** *Ann Oncol* 2010, **21(3)**:518–524.

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