

The Use of Preoperative Radiotherapy in the Management of Patients with Clinically Resectable Rectal Cancer: A Practice Guideline

Alvaro Figueredo¹ MD, Lisa Zuraw² MSc, Rebecca KS Wong³, MBChB, MSc, Olusegun Agboola⁴ MD, Ved Tandan¹ MD, R Bryan Rumble BSc^{2*}, and the members of the Gastrointestinal Cancer Disease Site Group.

Address: ¹Hamilton Regional Cancer; McMaster University, Hamilton, ON, ²Department of Clinical Epidemiology & Biostatistics, McMaster University, Hamilton, ON, ³Princess Margaret Hospital; University of Toronto, Toronto, ON, ⁴Ottawa Regional Cancer Centre; University of Ottawa, Ottawa, ON.

Email: Alvaro Figueredo – alvaro.figueredo@hrcc.on.ca; Lisa Zuraw – zurawl@mcmaster.ca; Rebecca Wong - rebecca.wong@rmp.uhn.on.ca; Olusegun Agboola - Olusegun.Agboola@orcc.on.ca; Ved Tandan - tandanv@mcmaster.ca; R. Bryan Rumble – rumbleb@mcmaster.ca

* Corresponding author

Abstract

Background

This systematic review with meta-analysis was designed to evaluate the literature and to develop recommendations regarding the use of preoperative radiotherapy in the management of patients with resectable rectal cancer.

Methods

The MEDLINE, CANCERLIT, Cochrane Library databases, and abstracts published in the annual proceedings of the American Society of Clinical Oncology and the American Society for Therapeutic Radiology and Oncology were systematically searched for evidence. Relevant reports were reviewed by four members of the Gastrointestinal Cancer Disease Site Group and the references from these reports were searched for additional trials. External review by Ontario practitioners was obtained through a mailed survey. Final approval of the practice guideline report was obtained from the Practice Guidelines Coordinating Committee.

Results

Two meta-analyses of preoperative radiotherapy versus surgery alone, eighteen trials that compared preoperative radiotherapy plus surgery to surgery alone, and five trials that compared preoperative radiotherapy to alternative treatments were obtained. Randomized trials demonstrate that preoperative radiotherapy followed by surgery is significantly more effective than surgery alone in preventing local recurrence in patients with resectable rectal cancer and it may also improve survival. A single trial, using surgery with total mesorectal excision, has shown similar benefits in local recurrence.

Conclusion

For adult patients with clinically resectable rectal cancer we conclude:

Preoperative radiotherapy is an acceptable alternative to the standard practice of postoperative radiotherapy for patients with stage II and III resectable rectal cancer.

Both preoperative and postoperative radiotherapy decrease local recurrence but neither improves survival as much as postoperative radiotherapy combined with chemotherapy. If preoperative radiotherapy is used, chemotherapy should be added postoperatively, at least for patients with stage III disease.

This report does not consider the use of preoperative radiotherapy to convert locally advanced, initially unresectable rectal cancer to resectable cases, to preserve the anal sphincter, or to delay the need for colostomy. Patients who choose preoperative radiotherapy as a treatment option instead of postoperative combined radiotherapy and chemotherapy need to be aware that because pathological stage is unknown until surgery is performed, many patients, who will not benefit from treatment, will be exposed to the risk of radiation-induced morbidity and mortality.

Background

Adenocarcinoma of the rectum is a common malignancy that originates in the last 15 centimeters of the large bowel where most of its external surface is not covered by peritoneum but rather is directly surrounded by connective-adipose tissue. Resection of the rectum and surrounding tissues can cure approximately 50% of patients; the other half will eventually die of the disease. Surgical failure is related to distant micro metastases that are not apparent and/or incomplete local resection. Local recurrences increase in frequency and survival decreases as the tumour penetrates through the rectal wall and extends to regional lymph nodes [1]. These prognostic factors form the basis for the TNM staging system widely used to advise therapy (see Appendix 1).

Meticulous dissection of perirectal tissues en-bloc with total mesorectal excision (TME) decreases local relapse [2]. Preoperative and postoperative radiotherapy (RT) has also been used to reduce local recurrence, which is associated with pelvic pain and rectal obstruction, and to prevent disease dissemination from the local site, thereby improving survival. Other goals of RT are to convert inoperable tumours into resectable cases, to preserve the anal sphincter and to avert a colostomy [3]. The advantages and disadvantages of preoperative and postoperative RT have been well-described [3]. The principal advantage of postoperative RT is that it is given only to patients at high risk of recurrence according to well-investigated prognostic factors, pathological stages II and III (B2 and C). Major incentives for preoperative RT originate from a variety of perspectives. Biologically, a tumour with an undisturbed circulation and oxygenation has a better chance for full radiation effects. Moreover, a tumour reduced in size and with the surrounding tissues sterilized facilitates surgery and reduces potential tumour dissemination. A

more practical incentive is the possibility of using an equally effective five-day course of high dose fraction preoperative RT instead of a 25-30 day standard postoperative course of RT.

Another practice guideline developed by the Gastrointestinal Cancer Disease Site Group (DSG) in 1997 reviewed the effects of postoperative RT and/or chemotherapy in resected stage II/III rectal cancer [4]. This guideline recommended the combined use of postoperative radiation and chemotherapy as the preferred treatment for resected stage II and III rectal cancer. This combined treatment improved the local recurrence rate by 50% and the 5-year survival rate by 42%. The same guideline discouraged the use of RT alone as adjuvant treatment because it only decreased local recurrence rates. For the present report, we initially reviewed only the effect of RT given before definitive surgery on survival and local recurrence. Following advice received from practitioner feedback and from our resource group, we have also included a discussion of preoperative RT compared with postoperative combined RT plus chemotherapy. This report does not consider the use of preoperative radiotherapy to convert locally advanced, initially unresectable rectal cancer into resectable cases, to preserve the anal sphincter, or to delay the need for colostomy.

Methods

This practice guideline was developed by the Cancer Care Ontario Practice Guidelines Initiative (CCOPGI), using the methodology of the Practice Guidelines Development Cycle [5]. Evidence was selected and reviewed by four members of the CCOPGI's Gastrointestinal Cancer DSG and methodologists. This practice guideline report is a convenient and up-to-date source of the best available evidence on preoperative radiotherapy for clinically resectable rectal cancer, developed

through systematic reviews, evidence synthesis and input from practitioners in Ontario. The report is intended to promote evidence-based practice. The Practice Guidelines Initiative is editorially independent of Cancer Care Ontario and the Ministry of Health and Long-Term Care.

External review by Ontario practitioners was obtained through a mailed survey consisting of items that address the quality of the draft practice guideline report and recommendations, and whether the recommendations should serve as a practice guideline. Final approval of the original guideline report was obtained from the Practice Guidelines Coordinating Committee (PGCC).

The CCOPGI has a formal standardized process to ensure the currency of each guideline report. This consists of periodic review and evaluation of the scientific literature, and where appropriate, integration of this literature with the original guideline information.

Literature Search Strategy

MEDLINE (1966 to December 2001), CANCERLIT (1983 to October 2001) and the Cochrane Library (Issue 4, 2001) were searched with no language restrictions. “Rectal neoplasms” (Medical subject heading [MeSH]), “colorectal neoplasms” (MeSH) and the text word “rectal cancer” were combined with “radiotherapy” (MeSH) and the following phrases used as text words: “preoperative”, “neoadjuvant”, “radiotherapy”, “radiation”, “irradiation”. These terms were then combined with the search terms for the following study designs or publication types: practice guidelines, meta-analyses, and randomized controlled trials. The proceedings of the 1998 to 2001 annual meetings of the American Society of Clinical Oncology (ASCO) and the

1999 to 2001 annual meetings of the American Society for Therapeutic Radiology and Oncology (ASTRO) were searched for reports of new or on-going trials. Relevant articles and abstracts were selected and reviewed and the reference lists from these sources were searched for additional trials. A search of personal reprint files was also conducted.

Study Selection Criteria

Trials of preoperative RT in resectable rectal cancer are characterized by multiple methodological problems because two treatments are combined (RT and surgery) to affect a heterogeneous condition (various populations and stages of rectal carcinoma) and to achieve a variety of goals (downstaging, improving resectability, decreasing local and possibly distant recurrences and improving survival). Cummings [6] detailed many of the pitfalls that marred early trials, including deficiencies in trial design, eligibility criteria, treatment standardization and reporting of results. We used this critique to develop standard criteria for the selection of trials of preoperative RT for rectal cancer. Studies were included in this systematic review of the evidence if they met all of the following criteria:

1. Patients were randomly assigned to preoperative RT versus surgery alone or an alternative treatment.
2. The study population was well defined. Studies preferably included only rectal carcinoma, defined by tumours located within 15 centimetres of the pectinate line or anal verge on sigmoidoscopy, or rectosigmoid tumours. Patients were screened for metastases and co-morbidity by clinical and imaging procedures and were assessed as surgically resectable for cure.
3. Treatments were described clearly, including RT dose, fractionation, duration, field size and

portals of irradiation. Timing of surgery after completion of RT was clearly set. General surgical principles were described.

4. Compliance with treatments and follow-up were described.
5. Treatment outcomes were reported for overall survival and/or local failure. Other outcomes such as adverse effects (morbidity and mortality), downstaging (decrease in the proportion of cases with stage III disease), and resectability (total and curative) were recorded if available.

Synthesizing the Evidence

Trials of preoperative RT versus surgery alone were pooled using Review Manager 4.0.3 (Metaview© Update Software), which is available through the Cochrane Collaboration. Overall mortality, local failure, tumour resectability, tumour downstaging, and adverse effects were pooled in separate analyses for all studies, where data was available. Reported figures or estimates obtained from tables or graphs were used. For calculation of survival and local failure, all eligible patients were considered in the denominator, based on intention to treat. All deaths at the time of reporting, regardless of cause, were included in survival calculations. Patients with local failure included those with non-resected as well as those with recurrent disease. Only resected cases were considered in the calculation of downstaging.

Data were pooled using the random effects model as the more conservative estimate of effect [7]. Results were expressed as risk ratios (RR) with 95% confidence intervals (CI), where a RR less than 1.0 favours preoperative RT and a RR greater than 1.0 favours surgery alone. Odds ratios (OR) and absolute risk differences (RD) were also calculated.

Heterogeneity of results among trials was expected in view of the different treatments used and populations tested, as well as the wide time interval and geography across which these trials were conducted. For example, the RT prescription may affect the results. RT doses greater than 30 Gy₁₀ are considered necessary and pelvic fields are as effective as extended fields. Moreover, the use of three or more RT beams will lessen toxicity and short delays of surgery after RT will not demonstrate downstaging. Thus, these factors were investigated with sensitivity analyses to see whether there was an impact on results. Outcomes of predetermined groups of patients were examined initially by the graphic method described by L'Abbe et al [8] and RR calculated. For sensitivity analyses the following factors were examined:

Treatment effects:

- Biologically effective dose (BED) of RT (less than 30 Gy₁₀ versus equal to or greater than 30 Gy₁₀). BED was calculated using the formula [9] $BED=nd(1+d/\alpha/\beta)$, where n=number of fractions, d=dose per fraction, $\alpha/\beta=10$ for tumour effect and acute reactions and $\alpha/\beta=3$ for late reactions, with no time correction (not needed for late reactions) because parameters were not available and usual ranges are quite wide;
- RT fraction size (standard fractions up to 2.5 Gy/day versus high fractions of 5 Gy/day);
- Contemporary radiotherapy prescription, defined as studies employing multiple-field technique and target volume confined to the pelvis (i.e., excluding studies employing parallel pair arrangements or including para-aortics); and
- Delay of surgery after completion of RT (less than seven days versus eight or more days).

Population effects:

- Studies including a range of rectal cancer cases versus those including only advanced disease.

Sensitivity analyses were also performed for all five of the meta-analyses (overall survival, local failure, tumour resectability, downstaging, and adverse effects) considering only trials with high design quality. The quality of the 14 eligible randomized trials of preoperative RT versus surgery alone in operable rectal cancer was scored independently. Five assessors assessed each trial using the Detsky instrument [10]. This questionnaire addresses five domains of study quality: randomization process, outcomes measure, patient eligibility, treatment description, and statistical procedures. The 14 questions on the Detsky instrument can be answered “adequate”, “inadequate” or “partial” and scored 1, 0, or 0.5, respectively. The final score of each trial is a ratio of the observed points divided by the total number of questions answered. The results from the five assessors were averaged for a final score. Trials with Detsky instrument scores greater than 0.5 were considered to be of high quality.

Results

Literature Search Results

The literature search identified 23 trials. Eighteen of these trials compared preoperative RT plus surgery to surgery alone [11-28]. Five trials compared preoperative RT to alternative treatments [29-33]. Two meta-analyses of preoperative RT versus surgery alone were found in a recent search update [34,35].

Preoperative Radiotherapy versus Surgery Alone

Four of the 18 trials of preoperative RT compared with surgery alone were excluded from the review [11,12,13,14]. Preliminary results of one trial have been reported in Russian [11], but the

report and results were difficult to interpret. This trial was excluded until more mature results are available and it is clear that the trial meets the inclusion criteria. Three trials [12,13,14] had major violations to the inclusion criteria. The Memorial Hospital trial included both randomized as well as non-randomized patients in the analysis [12]. The Veterans Administration Surgical Oncology Group (VASOG) Trial I included patients who may have had apparent metastases and did not allow for an analysis excluding this group of patients [13]. The Essen trial was described in summary form and was a failed trial of preoperative plus postoperative RT treatment due to difficulties with compliance [14].

This left 14 trials of preoperative RT compared with surgery alone [15-28]. One trial [17] contributed to two comparisons (single fraction preoperative RT versus surgery alone; multiple fractions preoperative RT versus surgery alone). One trial [27] included 316 patients who were also included in another trial [28]; neither report described results for these patients separately from those who were included in only one of the two trials. Reports of three trials [16,21,22] did not provide data on compliance with treatment. Where results have been reported or updated in more than one publication, only the most recent publication is listed.

The 14 trials that were reviewed are presented in Table 1. All trials included patients with resectable rectal cancer. Two trials also included patients with rectosigmoid tumours [15,18]. Some trials excluded patients with small tumours [24,27,28] or were limited to those with locally advanced tumours [22,25,26] or those requiring abdomino-perineal resection [18]. Patients excluded were those with evidence of distant metastases, previous malignancy or previous RT.

Surgery was performed from a few hours up to 40 days following the preoperative RT course, but most surgeries were done within one to four weeks of preoperative RT. The description of surgical procedures was very general except for the distinction between palliative and curative procedures. Radiation was delivered mostly by anterior and posterior portals. Only recent trials used three or four radiation portals [26-28]. Radiation fields were mostly pelvic but some trials used extended guitar-shaped fields up to L2 [15,19,20,24]. The total radiation dose and the fractionation schedules were quite different across studies, ranging from 5 Gy in a single treatment to over 5000 cGy in five weeks.

Compliance with treatments was generally well described. Follow-up data were collected prospectively in all but one trial [16]. The follow-up schedule was described for all trials except one [15]. The rate of patients lost to follow-up was 16% in one trial [23], but most other trials seemed to have had greater than 90% compliance with follow-up. The median follow-up at the time of the trial report was five years or more in most trials.

Pooled Results of Trials Comparing Preoperative RT to Surgery Alone

Overall survival

Survival was reported for all trials except one [22]. Survival results for only resected cases were reported for one trial [27]. The overall mortality risk ratio favoured preoperative RT (RR, 0.94; 95% CI, 0.89 to 0.99; $p=0.02$), results that correspond to an absolute risk difference of 4% (95% CI, 0.7% to 7.5%; $p=0.018$). There was significant heterogeneity across trials ($X^2=20.59$; $p<0.10$). As observed in Figure 1, most heterogeneity derived from a study from Brazil [21];

removal of this study resulted in non-significant heterogeneity and the results were similar (RR, 0.95; 95% CI, 0.91 to 0.99; p=0.012).

Local failure

Local failure rate was calculated for all trials as the number of patients unable to have tumour removal as well as those with recurrent disease after resection. The overall risk ratio favoured preoperative RT (RR, 0.71; 95% CI, 0.57 to 0.89; p=0.0025) with an absolute risk reduction of 8.6% (95% CI, 3.1% to 14.2%; p=0.0024). There was significant heterogeneity when local failure rates were pooled across trials ($X^2=61.72$; p<0.001). Figure 2 displays the local failure risk ratios with the trials arranged in ascending order of the RT dose (BED) used. There was no treatment effect in the analysis of three trials of preoperative RT using doses of 7.5 to 24.0 BED (RR, 1.01; 95% CI, 0.87 to 1.17; p=0.92) while trials using doses greater than or equal to 30 BED had evidence of reduced local failures (RR, 0.64; 95% CI, 0.49 to 0.82; p=0.00047).

Tumour resectability

Total tumour resectability between the treatment arms was not significantly different when 12 trials (14 comparisons) involving 5923 patients were pooled (RR, 1.00; 95% CI 0.99 to 1.00; p=0.36). Pooling of 14 trials (16 comparisons) involving 6816 patients detected no significant difference in curative resections for preoperative radiotherapy compared with surgery alone (RR, 0.99; 95% CI, 0.98 to 1.01; p=0.59). There was no significant heterogeneity among trials in either pooled analysis.

Downstaging

There was an overall significant decrease in the incidence of stage III rectal cancer among patients randomized to preoperative RT compared with surgery alone, but there was significant heterogeneity among the pooled results ($X^2=38.57$; $p<0.001$) (Figure 3). Neither radiation dose or its fractionation, nor timing of surgery affected results.

Adverse effects

Preoperative RT did not significantly increase 30-day postoperative mortality compared with surgery alone (RR, 1.33; 95% CI, 0.87 to 2.05; $p=0.19$). These results showed significant heterogeneity of trial results ($X^2=23.35$; $p<0.05$) (Figure 4). Results were not affected by radiation dose.

Postoperative morbidity was similar across trials and consisted mainly of delay of perineal wound healing and infection. The pooled results for postoperative morbidity also demonstrated significant heterogeneity ($X^2=62.74$; $p<0.001$) (Figure 5). Results were not different for patients receiving high or low dose RT or delay to surgery of <7 or >8 days.

Sensitivity analyses

Sensitivity analyses were conducted to test the robustness of the conclusions of the five pooled analyses above (overall survival, local failure, tumour resectability, downstaging, adverse effects) when excluding the three studies with quality scores of less than 0.5 [15,21,22]. Results of the pooled analyses of the best quality studies were not different from the results considering

all studies (data not shown). Details of the quality assessment results using the Detsky instrument can be found in Appendix 2.

Published meta-analyses

After completing our analysis, two literature-based meta-analyses of trials comparing preoperative RT to surgery alone for resectable rectal cancer were published [34,35]. The Camma et al [34] meta-analysis followed a methodology similar to ours. In the analysis, however, there was also an investigation of patient subgroups (Dukes' stages A, B and C, and male sex) and a regression analysis for overall survival (but not for other outcomes). The following regression variables were used: BED, stage of disease, male sex, study publication year, study size, allocation concealment and handling of withdrawals. This meta-analysis considered 14 published trials [13,15-21,23-27]: it included the VASOG-I trial [13], which we excluded, and it did not include a trial from Hungary [22] nor the single-fraction RT arm from the MRC-I trial [17]. All comparisons were made using odds ratios. Camma et al [34] detected a significant reduction in overall mortality with preoperative RT (Table 2). In a subgroup analysis, the decrease in mortality occurred in patients with Dukes' stages B and C but not in patients with stage A disease. No interaction was found between RT dose and survival (BED <30 or >30; $p=0.53$). In the regression analysis none of the factors used as variables had a significant impact on survival. Cancer-specific mortality and local recurrences were also reduced by the use of preoperative RT (Table 2) but not distant metastases. The overall rate of post-operative adverse events was higher in patients receiving preoperative RT (57.4% versus 42.4%; $p<0.001$). The 30-day postoperative mortality was not significantly different between patients receiving

preoperative RT and those having only surgery. Camma et al [34] concluded that preoperative RT reduced overall and cancer-specific mortality rates, and particularly local recurrence rates, while not affecting distant metastases. Postoperative mortality was not affected by the use of preoperative RT in spite of a higher rate of adverse events in patients.

The Colorectal Cancer Collaborative Group (CCCG) [35] identified trials of adjuvant preoperative and postoperative RT started before January 1st, 1987. The CCCG search yielded 19 trials of preoperative RT which included five trials not included in our review: two trials used of preoperative plus postoperative RT [14,30], one trial included patients with metastatic disease [13], another combined RT with 5-FU administration [32] and a Japanese trial published in 1989. Analysis was done by the log-rank method for overall and disease-specific mortality and for all and isolated recurrences. The overall yearly death rate was 5.6% (standard error ± 3.3). All recurrences and isolated recurrences were significantly decreased by preoperative RT at 5 years (45.9% vs. 52.9% and 12.5% vs. 22.2%, respectively) and at 10 years (55.1% vs. 60.8% and 16.7% vs. 25.8%, respectively). There was significant heterogeneity between the results of the 12 trials analyzed ($p=0.002$), which was explained by the greater efficacy of RT at higher biologically effective dose (>30 Gy). The reduction in local recurrence was proportionally similar for the various stages of the disease, and not affected by either sex or age. A multivariate analysis was not done [35].

The results of the published meta-analyses [34,35] and the one conducted for this systematic review are shown in Table 2.

Preoperative Radiotherapy versus Alternative Treatments

Preoperative RT versus postoperative adjuvant RT in high-risk cases

In a multi-institutional randomized trial in Sweden, patients with operable rectal cancer were randomized to preoperative RT or selective postoperative RT if the pathological stage was II or III [29,36]. Overall survival was the same for both treatment groups. When only patients with radical resection were considered, local recurrence was less likely for those receiving preoperative RT (11% vs. 22%; $p=0.02$) (Table 3). Postoperative complications, both early [29] and late [36], were significantly more frequent after higher-dose postoperative RT. The investigators emphasized that a short course of high fraction preoperative RT is preferable to a standard course of postoperative RT. Preoperative RT was better in reducing local recurrence rates and was associated with lower morbidity.

Two other randomized trials have investigated the benefit of preoperative RT in patients who also received postoperative RT if the pathological stage was II or III [30,31] (Table 3). In a study by the Radiation Therapy Oncology Group (RTOG), investigators randomized patients to a single dose of 500 cGy preoperatively and gave 45 Gy postoperatively to all high-risk patients [30]. After more than five years of follow-up, survival and local failure were similar in patients with or without preoperative RT. The preoperative RT dose in this trial was very small and has been shown to be ineffective [16,17]. German investigators performed a similar but smaller trial using a higher preoperative RT dose [31]. Patients were randomized to immediate surgery or to receive 16.5 Gy in 5 fractions preoperatively. After surgery, patients at high risk of local

recurrence (T4 stage, R1-2 or intraoperative tumour perforation) also received 41.4 Gy if they had preoperative RT and 59.8 Gy if they did not have preoperative RT. In a multivariate analysis of local recurrence, the only significant variable was staging (International Union Against Cancer [UICC]) ($p=0.0003$) while preoperative RT and T4 stage had non-significant effects ($p=0.08$ and 0.07 , respectively). In a similar analysis of survival, three variables were significant: age ($p=0.0003$), UICC stage ($p=0.001$) and residual disease status ($p=0.01$). Preoperative RT had a non-significant effect ($p=0.078$). These trials [30,31] indicate that selective postoperative RT annuls any potential positive effect of preoperative RT in low dose.

Preoperative RT alone versus preoperative RT plus chemotherapy

An early randomized trial by the European Organization for Research and Treatment of Cancer (EORTC) [32] compared preoperative RT with or without chemotherapy. Fluorouracil by bolus injection was given for four days during the first week of the radiation course. The trial was marred by many difficulties, with 27% of the cases being ineligible or not evaluable. The combined treatment did not reveal any advantage over RT alone (Table 3). Of interest was a marginally significant decrease in liver metastases for patients receiving preoperative combined treatment ($p=0.06$).

Preoperative RT with surgery at different intervals

One trial by French investigators tested whether the delay of surgery after the completion of preoperative RT is important [33] (Table 3). Operable patients with rectal tumours accessible to

digital rectal examination (stage T2-3, NX, M0) received preoperative RT (39 Gy in 13 fractions over 17 days through 3 fields) and were randomized to surgery after a short (two-week) or a long (six- to eight-week) interval. The only significant difference in outcomes, favouring the long over the short delay, was the higher proportion of patients with a clinical tumour response (partial plus complete) (53.1% versus 71.7%; $p=0.007$) and pathological downstaging (10.3% versus 26%; $p=0.005$). The three-year local failure and survival rates were not significantly different.

A recently published trial, coordinated by the Dutch Colorectal Cancer Group, included patients with rectal cancer not fixed or amenable to local excision. This study used standardized surgery with total mesorectal excision and randomized patients to surgery alone or surgery preceded by RT [37]. Nine hundred and twenty-four patients were randomized to surgery alone and 937 to preoperative RT consisting of 25 Gy in 5 fractions. The analysis of results is based on 908 patients allocated to surgery alone and 897 patients allocated to preoperative RT. After a median follow-up of only two years, the recurrence rate was significantly lower in patients receiving preoperative RT (2.4% vs. 8.2%; $p<0.001$) but overall survival was the same for both treatment groups. In a univariate subgroup analysis, the risk of local recurrence was significant for tumours with an inferior margin ≤ 5 cm ($p<0.05$) from the anal verge, an inferior margin 5.1 to 10cm ($p<0.001$) from the anal verge, TNM stage II tumours ($p<0.01$), and TNM stage III tumours ($p<0.001$). In a multivariate subgroup analysis, tests for interaction between tumour location, TNM stage and treatment were not significant, suggesting that treatment effect was similar for all the subgroups analyzed.

These early results of the trial would indicate that even with optimal surgery (total mesorectal excision) the rate of local recurrence can be reduced by preoperative RT. As the absolute difference in recurrence rates is 5.8%, 17 patients would require treatment to prevent or delay one local recurrence. It would be beneficial to know more about the toxicity, especially long-term anal dysfunction. Results of this trial were not added into the meta-analysis because it only reported two-year median follow-up data.

External Review

Practitioner feedback was obtained through a mailed survey. The survey consisted of items evaluating the methods, results, and interpretive summary used to inform the draft recommendations and whether the draft recommendations should be approved as a practice guideline. Written comments were invited. Follow-up reminders were sent at two weeks (post card) and four weeks (complete package mailed again). The Gastrointestinal Cancer DSG reviewed the results of the survey.

1. Number surveyed: 155 practitioners in Ontario (30 medical oncologists, 21 radiation oncologists, 100 surgeons, and four gastroenterologists).
2. Return rate: 63%
3. Written comments attached: 54%
4. Agreement with the summary of the evidence: 87%
5. Agreement with the recommendation: 85%
6. Approval of the recommendation as a practice guideline: 68%

Summary of Main Findings

Written comments provided by practitioners varied. Several respondents expressed concern with the statement in the draft recommendations that “Postoperative radiotherapy for patients with stage II/III rectal cancer is as effective in prolonging survival as preoperative radiotherapy for all rectal cancer patients regardless of stage of disease”. These respondents noted that preoperative versus postoperative radiotherapy was not the topic of this guideline and randomized trials addressing this issue are still ongoing.

Other respondents also questioned the recommendation that “postoperative radiotherapy combined with chemotherapy should remain the standard treatment”. They argued that if the results of the on-going trials demonstrate that preoperative and postoperative RT are equally effective, then one treatment should not be recommended over the other as the standard treatment. Several respondents raised the question of the role of total mesorectal excision.

Modifications or Actions

Practitioner feedback indicated a need to clarify the role of preoperative RT in the context of the companion guideline [4] recommending postoperative RT plus chemotherapy for stage II and III rectal cancer. In this context, the magnitude of benefits and drawbacks of preoperative and postoperative RT with and without chemotherapy are further discussed in the Gastrointestinal Cancer DSG Consensus section.

Gastrointestinal Cancer DSG Consensus

The discussion of the Gastrointestinal Cancer DSG focused on results from recent trials of preoperative RT in Europe that demonstrated significant improvements in local failure and survival rates. These results, achieved with a short course of radiation (five fractions) and with less toxicity than standard longer courses of radiation, have prompted the widespread use of this treatment modality in Europe and more recently in North America. Some treatment centres in Ontario have started phase II studies of preoperative RT, in some cases with concurrent chemotherapy.

There are, however, some concerns about the widespread use of preoperative RT. Some potential risks of the treatment seem preventable. The use of radiation given to smaller volumes and multiple fields, instead of the past practice of two fields, has been shown to decrease both early postoperative morbidity and mortality [24,27,28]. The exclusion of patients with poor performance status and those with ischemic changes in the electrocardiogram (ECG) [38] reduced both mortality and morbidity in the first two months. More difficult to predict is the long-term ano-rectal dysfunction, which restricts the social life of one third of survivors in some series following both preoperative and postoperative adjuvant RT [39]. Another concern is that some of the preoperatively irradiated patients would not have required this treatment based on the postoperative staging of the disease. Furthermore, the prognostic value of the postoperative staging of irradiated patients remains uncertain, particularly if there is downstaging of the disease. The postoperative pathological staging is very important to determine the need for adjuvant chemotherapy, which improves survival and reduces local recurrence [4].

Is preoperative RT an acceptable option to be offered to patients for adjuvant treatment in resectable rectal cancer? The previous recommendations from this group for patients with resected stages II and III rectal cancer was postoperative radiotherapy plus chemotherapy. This combined treatment has been demonstrated to significantly reduce local failure by 50% (95% CI, 8% to 73%) and improve patient survival by 42% (95% CI; 8% to 63%) in patients with stage II and III rectal cancer when compared to postoperative RT alone [4]. In similar patients, postoperative RT alone compared to observation after surgery decreased local recurrences by 27% (95% CI, 4% to 45%) but did not improve survival. Postoperative RT alone is, therefore, discouraged [4]. Preoperative RT alone, when compared to surgery, has been shown to decrease local failure by approximately 50% and to improve survival by approximately 15% (Table 2). The improvement in local recurrence occurs even after optimal surgery with TME. In a single trial [29,36], preoperative short-course RT has detected less local recurrence (11% versus 22%; $p=0.02$) and less morbidity than conventional postoperative RT alone. From these results it can be inferred that preoperative RT is a better treatment choice than postoperative RT with less local failures and less morbidity. A comparison of preoperative RT followed by postoperative chemotherapy versus combined postoperative RT plus chemotherapy is presently being investigated in clinical trials but mature results are not yet available for review.

While we confirm our recommendation for combined postoperative RT plus chemotherapy for resected patients with stage II and III rectal cancer, based on the evidence from the Swedish and Dutch trials [27,37], preoperative RT (followed by chemotherapy for at least patients with stage III) is an alternative provided the patient is made aware of the potential benefits and drawbacks. Benefits are the decrease in local failure and in treatment morbidity. Local failure is an

important outcome in rectal cancer as recurrences are associated with significant disability. Drawbacks are the need to use preoperative RT in most patients compared to RT administered according to postoperative staging and the possibility that patients not requiring radiation may develop treatment associated complications.

Physicians should encourage patients to participate in clinical trials of the primary treatment of rectal cancer. These trials should require the best possible surgery, the confirmation of the accuracy of clinical staging versus pathological staging, and the use of measures of quality of life. Patients must also be clearly advised of the differences between treatment approaches.

Discussion

Results of three meta-analyses (Table 2) indicate that preoperative RT compared to surgery alone significantly decreases the risk of local failure and overall mortality. The absolute reduction in local failure is 8.6% (95% CI, 3.1% to 14.2%) while the absolute reduction in overall mortality at five years is 3.5% (95% CI, 1.1% to 6.0%). Early results of the Dutch trial [37] confirm the decrease in local recurrence with preoperative RT even after optimal surgery with total mesorectal excision. The improved results of recent trials can be explained by better patient selection and radiation prescription. Swedish investigators, comparing the results of the Stockholm-I [24] and Stockholm-II [28] trials of preoperative RT, showed that the overall survival of patients is significantly affected by the 60-day postoperative mortality rate [38]. This early fatality rate is due to an excess of infectious, cardiovascular and thromboembolic causes. The cause for this excess in mortality is attributed to the delivery of similar doses of radiation by larger versus smaller

volumes, by two rather than multiple radiation portals and to patient characteristics such as evidence of ischemia by ECG and poor performance status. An increase in morbidity was also observed and consisted of venous thromboembolism, femoral neck and pelvic fractures and intestinal obstruction. A subgroup of patients who participated in the Swedish trial [27] completed a questionnaire about ano-rectal dysfunction. Bowel disturbances led to social restrictions in 30% of patients who received preoperative RT compared with 10% of patients who received surgery alone ($p < 0.01$). The abnormalities included more frequent bowel movements, urgency and incontinence. No single factor could be identified to explain the complications, but the authors postulated the radiation effect on the anal sphincter itself or on its nerve supply [39]. Similar anorectal dysfunction has been reported after postoperative RT combined with chemotherapy [40].

Preoperative RT in high fractions has been compared with standard low-fraction postoperative adjuvant RT [29]. The overall survival was the same for both treatment groups but the local recurrence rate was lower and the morbidity less for preoperative RT. While preoperative RT was given to all cases, postoperative RT was given only to high-risk cases (stages B2 and C), a group equal to half the number of cases treated with the preoperative approach. The increased morbidity of the postoperative RT must be related in part to the higher radiation dose.

The use of preoperative RT in small doses did not decrease the indications for postoperative adjuvant RT [30,31]. Patients who received four bolus injections of 5-FU during the preoperative RT in one old trial increased postoperative mortality and decreased overall survival [32]. The cause for this toxicity is not known and has not been observed in subsequent trials of combined therapy. The delay of surgery after preoperative RT for more than two weeks

decreased the rate of stage III disease but had no impact on resectability, local recurrence or survival [33].

Practice Guidelines Coordinating Committee Approval Process

The guideline was circulated to 11 members of the Practice Guideline Coordinating Committee. Seven members returned ballots; five approved the guideline report as written, and two members approved the guideline conditional on the Gastrointestinal Cancer DSG addressing suggestions for revision. The suggestions referred to the recommendations and the meta-analyses. Changes were made to the guideline based on these suggestions, and a revised version was resubmitted to the PGCC for further consideration. The practice guideline was approved with one dissenting vote.

Conclusions

This practice guideline reflects the integration of the draft recommendations with feedback obtained from the external review process. It has been approved by the Gastrointestinal DSG and the Practice Guidelines Coordinating Committee.

Target Population

These recommendations apply to adult patients with clinically resectable rectal cancer. This report does not consider the use of preoperative radiotherapy to convert locally advanced, initially unresectable rectal cancer to resectable cases, to preserve the anal sphincter, or to delay the need for colostomy.

Recommendations

- Preoperative radiotherapy is an acceptable alternative to the standard practice of postoperative radiotherapy for patients with stage II and III resectable rectal cancer.
- Both preoperative and postoperative radiotherapy decrease local recurrence but neither improves survival as much as postoperative radiotherapy combined with chemotherapy. Therefore, if preoperative radiotherapy is used, chemotherapy should be added postoperatively, at least for patients with stage III disease.

Qualifying Statement

Patients who choose preoperative radiotherapy as a treatment option instead of postoperative combined radiotherapy and chemotherapy need to be made aware that because pathological stage is unknown until surgery is performed, many patients, who will not benefit from treatment, will be exposed to the risk of radiation-induced morbidity and mortality.

List of Abbreviations Used

TNM, Tumour, Node, Metastases (the staging system of the UICC); RT, radiotherapy; DSG, Disease Site Group; CCOPGI, Cancer Care Ontario Practice Guidelines Initiative; MeSH, medical subject heading; ASCO, American Society of Clinical Oncology; ASTRO, American Society for Therapeutic Radiation and Oncology; RR, Relative Risk ratio; CI, Confidence Interval; OR, Odds Ratio; RD, Risk Difference; Gy, Gray; BED, biologically effective dose; VASOG, Veterans Administration Surgical Oncology Group; cGy, cyclic Gray; MRC, Medical Research Council; CCCG, Colorectal Cancer Collaborative Group; RTOG, Radiation Therapy

Oncology Group; UICC, International Union Against Cancer; EORTC, European Organization for Research and Treatment of Cancer; ECG, electrocardiogram; TME, Total Mesorectal Excision; 5-FU, 5-Fluorouracil; PGCC, Practice Guidelines Coordinating Committee.

Competing Interests

None declared.

Authors' Contributions

AF and LZ participated in the statistical analysis and did the majority of the drafting of the manuscript with input from KSW, OA, VT, and other Gastrointestinal Cancer DSG members. KSW, OA, and VT participated in study selection and study quality assessments, and provided feedback on all drafts.

Acknowledgements

Additional members of Cancer Care Ontario's Practice Guidelines Initiative's Gastrointestinal Cancer Disease Site Group include: Dr. J. Maroun MD (Chair), Dr. B. Cummings MD (Vice-Chair), Dr. O. Agboola MD, Mr. M. Citron (Community Representative), Dr. F.G. DeNardi MD, Dr. C. Earle MD, Dr. A. Figueredo MD, Dr. S. Fine MD, Dr. B. Fisher MD, Dr. C. Germond MD, Dr. D. Jonker MD, Dr. K. Khoo MD, Dr. W. Kocha MD, Ms. M. Lethbridge (Community Representative), Dr. W. Lofters MD, Dr. R. Malthaner MD, Dr. M. Moore MD, Dr. V. Tandan MD, and Dr. R. Wong MD. Please see the Cancer Care Ontario Practice Guidelines Initiative (CCOGPI) web site (<http://www.ccopebc.ca/>) for a complete list of current Gastrointestinal Cancer Disease Site Group members.

References

1. Cohen AM, Minsky BD, Schilsky RL: **Cancer of the rectum.** In: *Cancer: Principles and Practice of Oncology*. 5th ed. (Edited by DeVita VT, Hellman S, Rosenberg SA) Philadelphia-New York: Lippincott-Raven 1997. 1197-1233
2. MacFarlane JK, Ryall RDH, Heald RJ: **Mesorectal excision for rectal cancer.** *Lancet* 1993, **341**: 457-460
3. Mohiuddin M: **Radiation: The preoperative adjuvant strategy.** In: *Cancer of the Colon, Rectum, and Anus* (Edited by Cohen AM, Winawer SJ) New York, McGraw-Hill 1995, 657-668
4. Figueredo A, Germond C, Taylor B, Maroun J, Agboola O, Wong R, Zwaal C, Micucci S, for the Gastrointestinal Cancer Disease Site Group. **Post-operative adjuvant radiotherapy or chemotherapy for resected stage II or III rectal cancer.** *Curr Oncol* 2000, **7**: 37-51
5. Browman GP, Levine MN, Mohide EA, Hayward RSA, Pritchard KI, Gafni A, Laupacis A: **The practice guidelines development cycle: a conceptual tool for practice guidelines development and implementation.** *J Clin Oncol* 1995, **13**: 502-512
6. Cummings BJ: **A critical review of adjuvant pre-operative radiation therapy for adenocarcinoma of the rectum.** *Br J Surg* 1986, **73**: 332-338
7. DerSimonian R, Laird N: **Meta-analysis in clinical trials.** *Controlled Clin Trials* 1986, **7**: 177-188
8. L'Abbe KA, Detsky AS, O'Rourke K: **Meta-analysis in clinical research.** *Ann Int Med* 1987, **107**: 224-233
9. Fowler JF: **The linear-quadratic formula and progress in fractionated radiotherapy.** *Br J Radiol* 1989, **62**: 679-694

10. Detsky AS, Naylor CD, O'Rourke K, McGeer AJ, L'Abbe KA: **Incorporating variations in the quality of individual randomized trials into meta-analysis.** *J Clin Epidemiol* 1992, **45**: 255-265
11. Knysh VI, Alieve BM, Barsukov YA: **Combined treatment of rectal cancer using two variants of preoperative concentrated radiotherapy** [Russian]. *Med Radiol Mosk* 1983, **28**: 12-17
12. Stearns MW Jr, Deddish MR, Quan SHQ, Leaming RH: **Preoperative roentgen therapy for cancer of the rectum and rectosigmoid.** *Surg Gynecol Obstet* 1974, **138**: 584-586
13. Higgins GA Jr, Conn JH, Jordan PH Jr, Humphrey EW, Roswit B, Keehn RJ: **Preoperative radiotherapy for colorectal cancer.** *Ann Surg* 1975, **181**: 624-630
14. Niebel W, Schulz U, Ried M, Erhard J, Beersiek F, Blöcher G, Nier H, Halama H, Scherer E, Zeller G, Eigler FW, Tacke J: **Five-year results of a prospective randomized study: experience with combined radiotherapy and surgery of primary rectal carcinoma.** *Recent Results Cancer Res* 1988, **110**: 111-113
15. Kligerman MM, Urdaneta N, Knowlton A, Vidone R, Hartman PV, Vera R: **Preoperative irradiation of rectosigmoid carcinoma including its regional lymph nodes.** *Am J Roengenol* 1972, **114**: 498-503
16. Rider WD, Palmer JA, Mahoney LJ, Robertson CT: **Preoperative irradiation in operable cancer of the rectum: report of the Toronto trial.** *Can J Surg* 1977, **20**: 335-338
17. Second Report of an MRC Working Party. **The evaluation of low dose pre-operative x-ray therapy in the management of operable rectal cancer; results of a randomly controlled trial.** *Br J Surg* 1984, **71**: 21-25

18. Higgins GA, Humphrey EW, Dwight RW, Roswit B, Lee LE Jr, Keehn RJ: **Preoperative radiation and surgery for cancer of the rectum.** Veterans Administration Surgical Oncology Group Trial II. *Cancer* 1986, **58**: 352-359
19. Dahl O, Horn A, Morild I, Halvorsen JF, Odland G, Reinertsen S, Reisaeter A, Kavli H, Thunold J: **Low-dose preoperative radiation postpones recurrences in operable rectal cancer. Results of a randomized multicenter trial in western Norway.** *Cancer* 1990, **66**: 2286-2294
20. Gérard A, Buyse M, Nordlinger B, Loygue J, Pene F, Kempf P, Bosset J-F, Gignoux M, Arnaud J-P, Desai C, Duez N: **Preoperative radiotherapy as adjuvant treatment in rectal cancer. Final results of a randomized study of the European Organization for Research and Treatment of Cancer (EORTC).** *Ann Surg* 1988, **208**: 606-614
21. Reis-Neto JA, Quilici FA, Reis JA Jr: **A comparison of nonoperative vs. preoperative radiotherapy in rectal carcinoma. A 10-year randomized trial.** *Dis Colon Rectum* 1989, **32**: 702-710
22. Illényi L, Grexa E, Gecser G, Kött I: **Local recurrence of rectal cancer following preoperative irradiation.** *Acta Chir Hung* 1994, **34**: 333-347
23. Goldberg PA, Nicholls RJ, Porter NH, Love S, Grimsey JE: **Long-term results of a randomized trial of short-course low-dose adjuvant pre-operative radiotherapy for rectal cancer: reduction in local treatment failure.** *Eur J Cancer* 1994, **30A**: 1602-1606
24. Stockholm Rectal Cancer Study Group. **Preoperative short-term radiation therapy in operable rectal carcinoma. A prospective randomized trial.** *Cancer* 1990, **66**: 49-55

25. Medical Research Council Rectal Cancer Working Party. **Randomised trial of surgery alone versus radiotherapy followed by surgery for potentially operable locally advanced rectal cancer.** *Lancet* 1996, **348**: 1605-1610
26. Marsh PJ, James RD, Schofield PF: **Adjuvant preoperative radiotherapy for locally advanced rectal carcinoma. Results of a prospective, randomized trial.** *Dis Colon Rectum* 1994, **37**: 1205-1214
27. Swedish Rectal Cancer Trial: **Improved survival with preoperative radiotherapy in resectable rectal cancer.** *New Engl J Med* 1997, **336**: 980-987
28. Stockholm Colorectal Cancer Study Group: **Randomized study on preoperative radiotherapy in rectal carcinoma.** *Ann Surg Oncol* 1996, **3**: 423-430
29. Pahlman L, Glimelius B: **Pre- or postoperative radiotherapy in rectal and rectosigmoid carcinoma: report from a randomized multicenter trial.** *Ann Surg* 1990, **211** : 187-195
30. Sause WT, Pajak TF, Noyes RD, Dobelbower R, Fischbach J, Doggett S, Mohiuddin M: **Evaluation of preoperative radiation therapy in operable colorectal cancer.** *Ann Surg* 1994, **220**: 668-675
31. Herrmann T, Petersen S, Hellmich G, Baumann M, Ludwig K: **Delayed toxicity of brief preoperative irradiation and risk-adjusted postoperative radiotherapy of operative rectal carcinoma. Results of a randomized prospective study [German].** *Strahlentherapie und Onkologie* 1999, **175**: 430-436

32. Boulis-Wassif S, Gerard A, Loygue J, Camelot D, Buyse M, Duez N: **Final results of a randomized trial on the treatment of rectal cancer with preoperative radiotherapy alone and in combination with 5-fluorouracil, followed by radical surgery.** Trial of the European Organization on Research and Treatment of Cancer Gastrointestinal Tract Cancer Cooperative Group. *Cancer* 1984, **53**: 1811-1818
33. Francois Y, Nemoz CJ, Baulieux J, Vignal J, Grandjean JP, Partensky C, Souquet JC, Adeleine, Gerard J-P: **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**: 2396-2402
34. Cammà C, Giunta M, Fiorica F, Pagliaro L, Craxi A, Cottone M: **Preoperative radiotherapy for resectable rectal cancer. A meta-analysis.** *JAMA* 2000, **284**: 1008-1015
35. Colorectal Cancer Collaborative Group: Adjuvant radiotherapy for rectal cancer: a systematic overview of 8507 patients from 22 randomized trials. *Lancet* 2001, **358**: 1291-1304
36. Frykholm GJ, Glimelius B, Pahlman L: **Preoperative or postoperative irradiation in adenocarcinoma of the rectum: final treatment results of a randomized trial and an evaluation of late secondary effects.** *Dis Colon Rectum* 1993, **36**: 564-572
37. Kapiteijn E, Marijnen CAM, Nagtegaal ID, Putter H, Steup WH, Wiggers T, van Krieken JHJM, Hermans J, Leer JWH, van de Velde CJH: **Preoperative radiotherapy combined with total mesorectal excision for resectable rectal cancer.** *New Engl J Med* 2001, **345**: 638-646
38. Holm T, Singnomklao T, Rutqvist LE, Cedermark B: **Adjuvant preoperative radiotherapy in patients with rectal carcinoma. Adverse effects during long term follow-up of two randomized trials.** *Cancer* 1996, **78**: 968-976

39. Dahlberg M, Glimelius B, Graf W, Pahlman L: **Preoperative irradiation affects functional results after surgery for rectal cancer. Results from a randomized study.** *Dis Colon Rectum* 1998, **41**: 543-551
40. Kollmorgen CF, Meagher AP, Wolff BG, Pemberton JH, Martenson JA, Ilstrup DM: **The long-term effect of adjuvant postoperative chemoradiotherapy for rectal carcinoma on bowel function.** *Ann Surg* 1994, **20**: 676-682

Table 1 - Randomized trials of preoperative radiotherapy (RT) versus surgery alone in resectable rectal cancer.

Trial (Reference)	No. of Patients Randomized RT / no RT (Analyzed)	RT Prescription*	Biologically Effective Dose (Gy ₁₀)	Compliance with RT [†]			Surgical Delay after RT (Days)	Median Follow-up (Months)	Study Quality Score [‡]
				0	<1	>1			
Yale [1§]	15 (15) / 16 (16)	45 in 25 2 fields, G	53.1	-	7%	-	2-42	>60	0.29
Toronto [1¶]	60 (60) / 65 (65)	5 in 1 2 fields, P	7.5	-	-	-	0	72	0.55
MRC-I [17]	549 (549) / 275 (275)	5 in 1 <u>or</u> 20 in 10 2 fields, G	7.5 <u>or</u> 24.0	2%	6%	-	<7	>60	0.66
VASOG-II [1§]	180 (180) / 181 (181)	31.5 in 18 2 fields, G	37.0	7%	11%	4%	~40	60	0.70
Norway [19]	159 (155) / 150 (145)	31.5 in 18 2 fields, G	37.0	1%	-	-	<21	54	0.80
EORTC [20]	236 (231) / 230 (228)	34.5 in 15 2 fields, G	42.4	1%	-	-	1-69	>72	0.67
Brazil [21]	34 (34) / 34 (34)	40 in 20 2 fields, P	48.0	-	-	-	7	120	0.48
Hungary [22]	171 (171) / 165 (165)	40 in 20 <u>or</u> 50 in 25 2 fields, P	48.0 60.0	-	-	-	42	>60	0.44
ICRF-UK [23]	228 (228) / 239 (239)	15 in 3 2 fields, P	22.5	-	10%	-	<2	60	0.64
Stockholm-I [24]	424 (424) / 425 (425)	25 in 5 2 fields, G	37.5	-	12%	-	<7	107	0.74
MRC-II [25]	139 (139) / 140 (140)	40 in 20 2 fields, P	48.0	1%	5%	4%	<7	>60	0.83
NW-UK [26]	143 (143) / 141 (141)	20 in 4 3 field, P	30.0	-	1%	3%	<7	96	0.57
Sweden [27]	583 (573) / 585 (574)	25 in 5 3-4 fields, P	37.5	3%	1% [§]	-	<7	>60	0.82
Stockholm-II [28]	272 (272) / 285 (285)	25 in 5 4 fields, P	37.5	-----	5%	-----	<7	50	0.88

Note: EORTC indicates European Organization for Research and Treatment of Cancer; ICRF-UK, Imperial Cancer Research Fund United Kingdom; MRC, Medical Research Council; NW-UK, Northwest Region Rectal Cancer Group United Kingdom; VASOG, Veterans Administration Surgical Oncology Group.

* Total dose in Gy, number of fractions, number of treatment fields, target volume (G, guitar-shaped; P, pelvic).

† 0 indicates per cent of patients receiving no treatment; <1, percent receiving less than planned treatment; >1, percent receiving more than planned treatment.

‡ Based on independent assessment by five reviewers using the Detsky instrument¹⁰.

§ 5% of patients also received RT over >7 days, and 8% of patients received radiation through a two portal beam.

Table 2 - Results of three meta-analyses of preoperative radiotherapy versus surgery alone in resectable rectal cancer.

Outcome	Pooled Results		
	Odds Ratio, 95% Confidence Interval and p value		
	Camma et al [34]	CRCCG [35]*	Present Review
Overall Mortality	0.84 (0.72, 0.98) p=0.03	0.95 (0.86, 1.06) p=0.34	0.85 (0.74, 0.97) p=0.019
Local Failure	0.49 (0.38, 0.62) p<0.001	0.58 (0.42, 0.80) p=0.00008	0.62 (0.45, 0.85) p=0.0029
30-day Postoperative Mortality	1.38 (0.86, 2.32) p=0.22	- -	1.42 (0.90, 2.23) p=0.13

* ORs calculated from data shown in Figure 1 [35].

Table 3 - Randomized trials of preoperative radiotherapy compared to alternative treatments in rectal cancer.

Trial (Reference)	Treatment			No. Patients	Local Failure (%)	Survival Rate at 5 years (%)
	Preoperative RT	Surgery Delay	Postoperative RT (Delay)			
Pahlman & Glimelius [29]	5.1 Gy x 5	1 week	-	236	22%	43%
	-	-	60 Gy in 54* (4-6 weeks)	235	33%	37%
					p=0.012	p=0.43
RTOG [30]	0.5 Gy x 1	1 day	45-51 Gy*	175	32%	43%
	-	-	(2-6 weeks)	178	32%	32%
					p=NS	p=NS
Herrmann et al [31]	3.3 Gy x 5	1-2 days	41.5 Gy in 48*	48	25%	49%
	-	-	59.8 Gy in 56* (6-14 weeks)	46	39%	37%
					p=0.142	p=NS
EORTC [32]	2.3 x 15	2 weeks	-	121	15%	59%
	2.3 x 15	2 weeks	-	126	15%	46%
	+ 5-FU				p=NS	p=0.06
Francois et al [33]	3.0 Gy x 13	2 weeks	-	99	9%	78% [†]
	3.0 Gy x 13	6-8 weeks	-	102	9%	73% [†]
					p=NS	p=NS

Note: NS indicates not statistically significant; RTOG, Radiation Therapy Oncology Group.

* Radiation given only to high-risk cases (stages B2 and C).

[†] 3-year survival rate.

Figure 1 - Meta-analysis examining preoperative radiotherapy for patients with resectable rectal cancer: overall mortality.

[insert Figure 1 here]

Results were reported for only the eligible or evaluable patients for Norway [19] trial.
Results were reported for only the eligible patients undergoing surgery for the Sweden [27] trial.

Figure 2 - Meta-analysis examining preoperative radiotherapy for patients with resectable rectal cancer: local failure.

[insert Figure 2 here]

Results were reported for only the eligible or evaluable patients for the Norway [19] trial.

Figure 3 - Meta-analysis examining preoperative radiotherapy for patients with resectable rectal cancer: incidence of stage III tumours.

[insert Figure 3 here]

Figure 4 - Meta-analysis examining preoperative radiotherapy for patients with resectable rectal cancer: 30-day postoperative mortality.

[insert Figure 4 here]

Figure 5 - Meta-analysis examining preoperative radiotherapy for patients with resectable rectal cancer: postoperative morbidity.

[insert Figure 5 here]

Comparison: 01 Overall Mortality

Outcome: 01 Overall Mortality

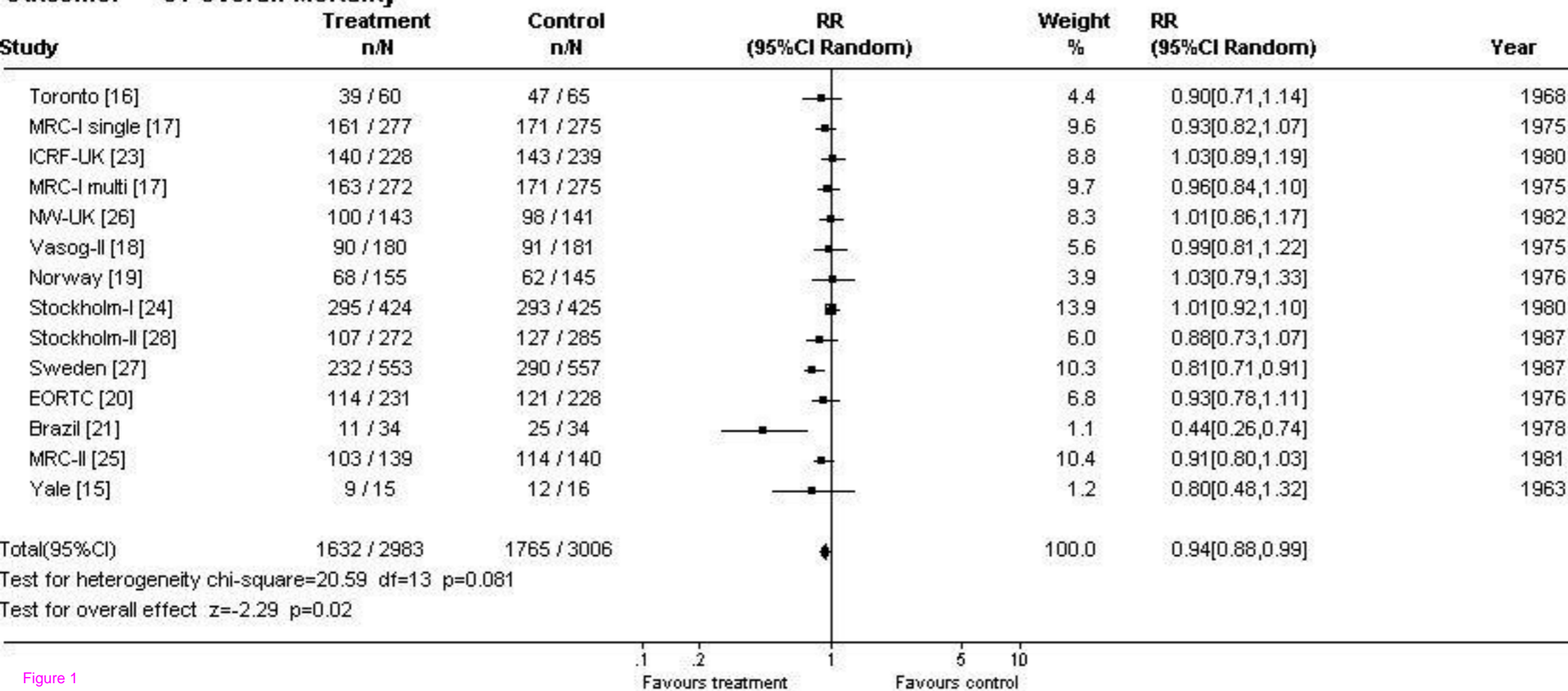


Figure 1

Comparison: 02 Local Failure

Outcome: 01 Local Failure

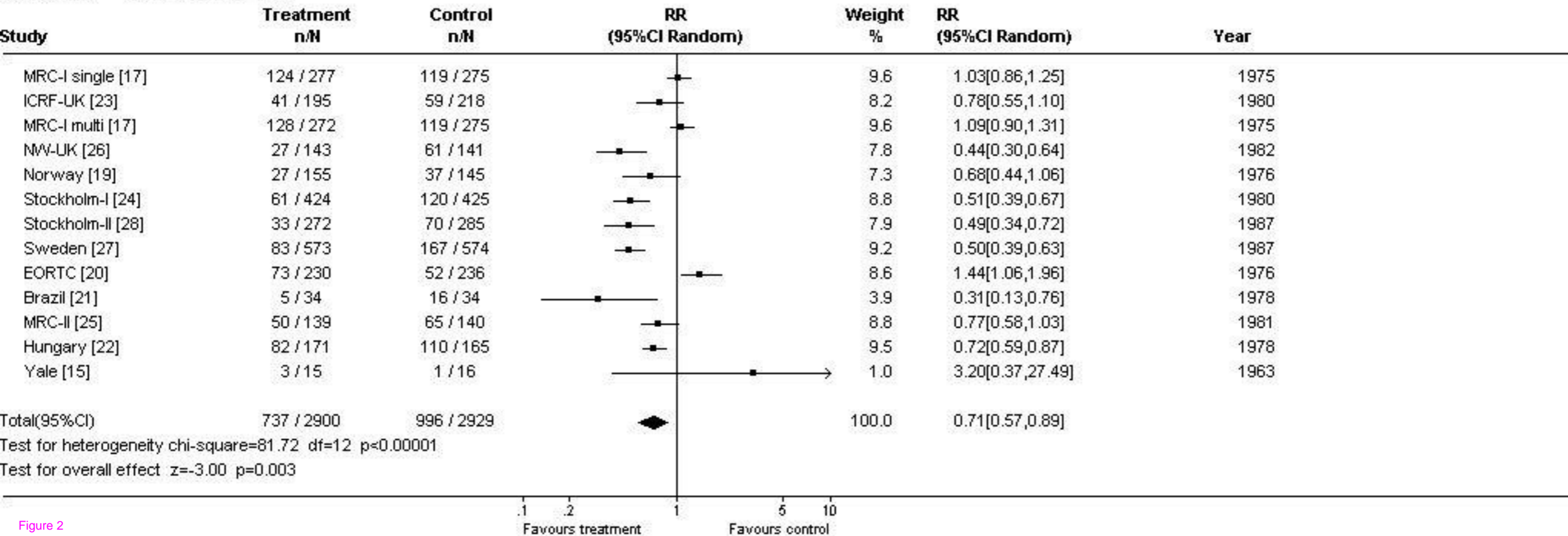


Figure 2

Comparison: 03 Incidence of Stage III Tumours

Outcome: 01 Incidence of Stage III Tumours

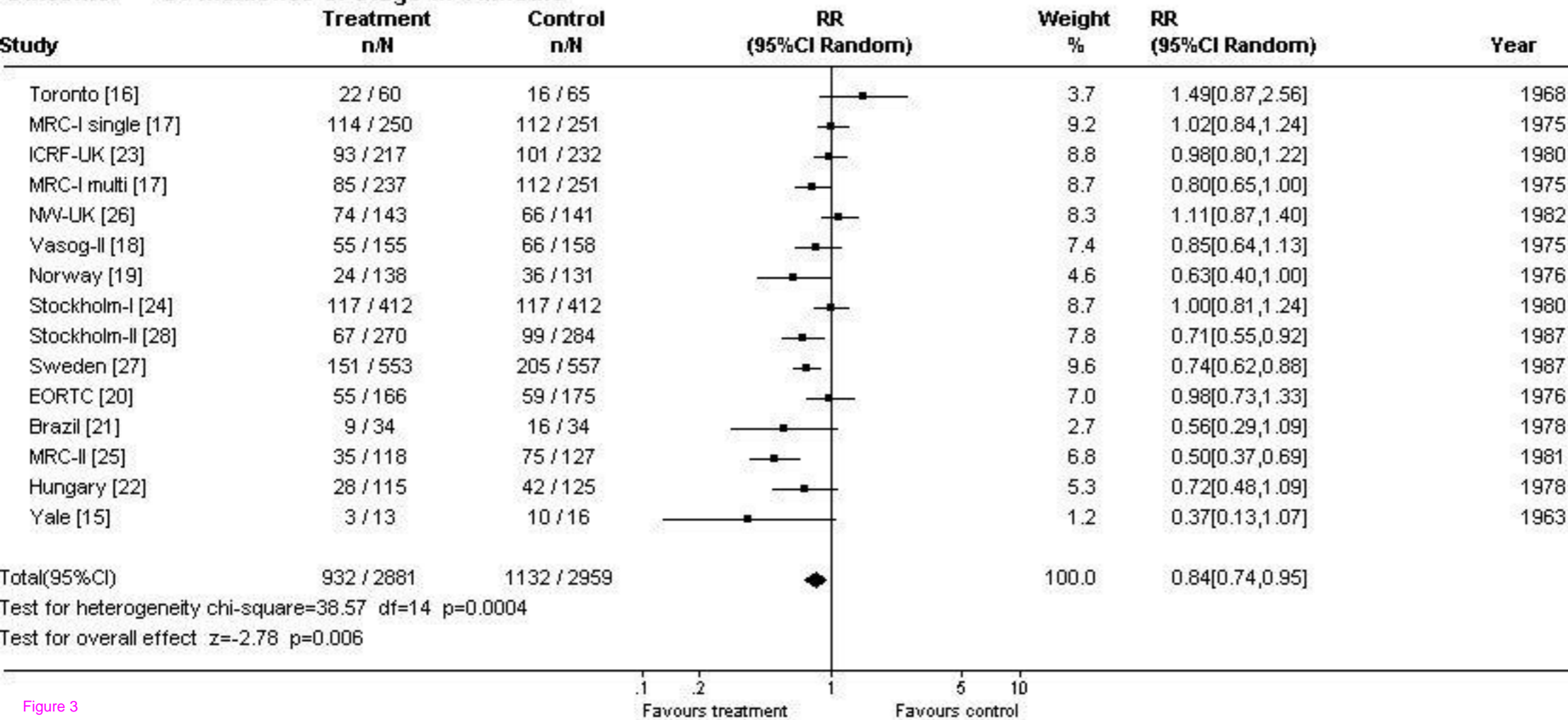


Figure 3

Comparison: 04 30-day Postoperative Mortality

Outcome: 01 30-day Postoperative Mortality

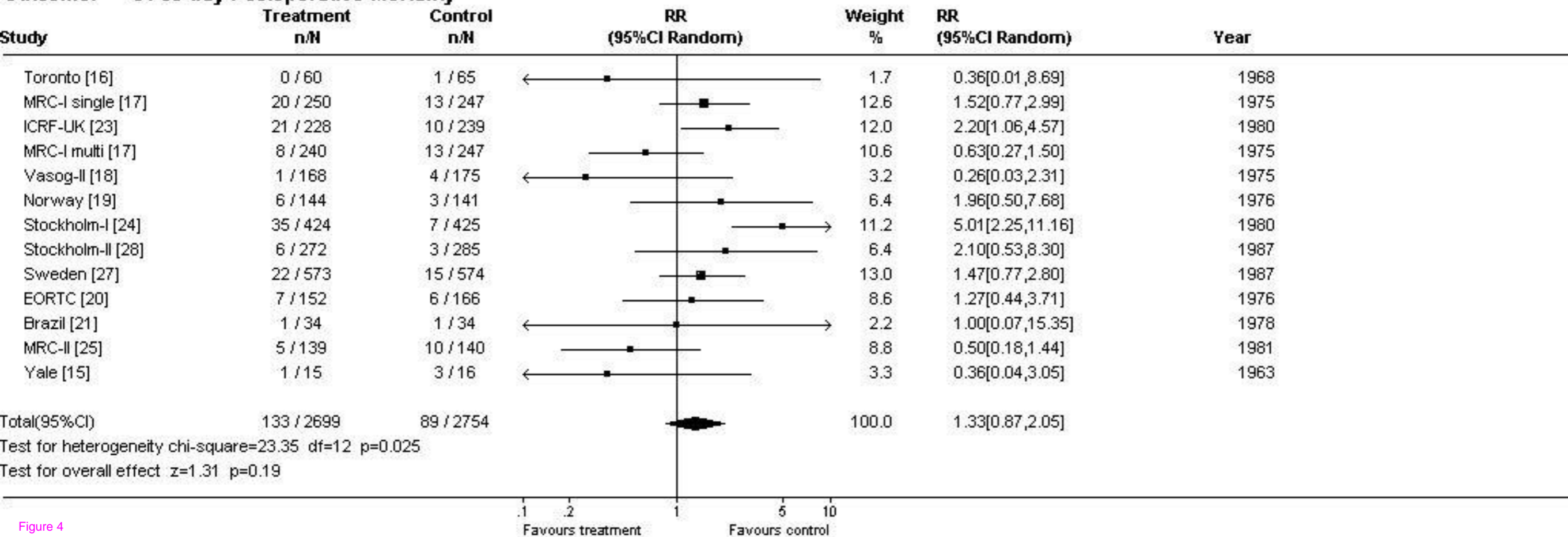


Figure 4

Comparison: 05 Postoperative Morbidity

Outcome: 01 Postoperative Morbidity

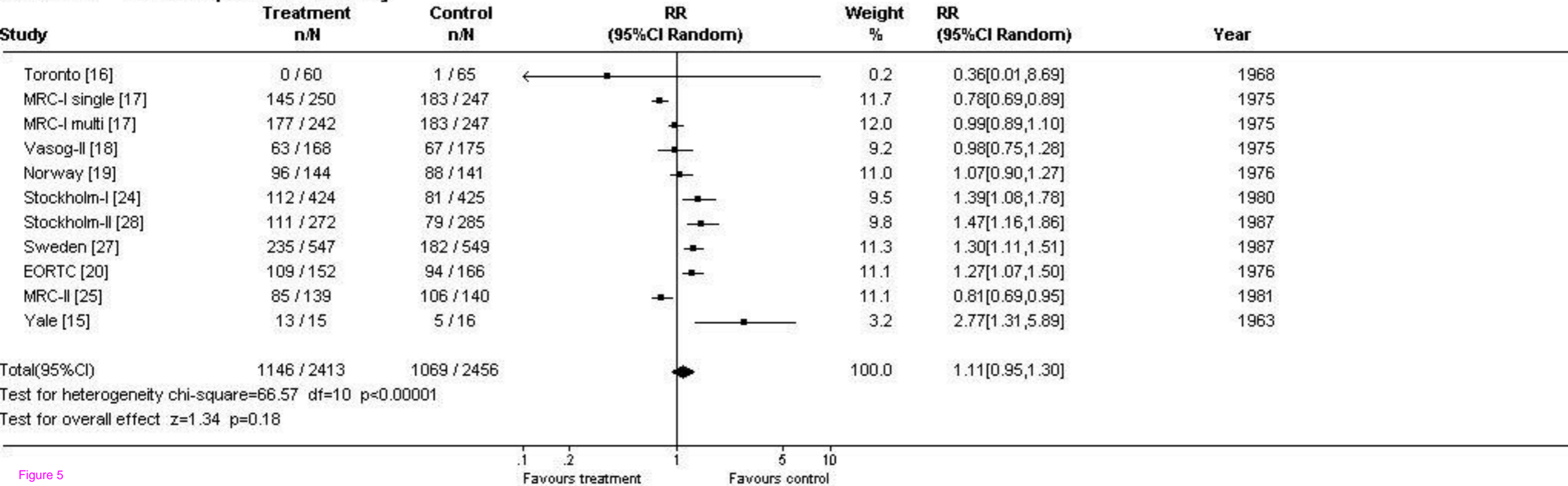


Figure 5

Java Exception error '80004005'

com.biomedcentral.BMCEException: com.ms.wfc.data.AdoException: ORA-01722: invalid number : 1895843828146123 : 1

/msp_sup.asp, line 9