

- 4 Iniesta P, Massa MJ, Gonzalez-Quevedo R, de Juan C, Moran A, Sanchez-Pernaute A, Cerdan J, Torres A, Balibrea J and Benito ML: Loss of heterozygosity at 3p23 is correlated with poor survival in patients with colorectal carcinoma. *Cancer* 89: 1220-1227, 2000.
- 5 Arribas R, Ribas M, Risques RA, Masramon L, Tortola S, Marcuello E, Aiza G, Miro R, Capella G and Peinado MA: Prospective assessment of allelic losses at 4p14-16 in colorectal cancer: two mutational patterns and a locus associated with poorer survival. *Clin Cancer Res* 5: 3454-3459, 1999.
- 6 Gebert J, Sun M, Ridder RA, Hinz U, Lehnert T, Moller P, Schackert HK, Herfarth C and von Knebel Doeberitz M: Molecular profiling of sporadic colorectal tumors by microsatellite analysis. *Int J Oncol* 16: 169-179, 2000.
- 7 Halling KC, French AJ, McDonnell SK, Burgart LJ, Schaid DJ, Peterson BJ, Moon-Tasson L, Mahoney MR, Sargent DJ, O'Connell MJ, Witzig TE, Farr GH Jr, Goldberg RM and Thibodeau SN: Microsatellite instability and 8p allelic imbalance in stage B2 and C colorectal cancers. *J Natl Cancer Inst* 91: 1295-1303, 1999.
- 8 Choi SW, Lee KJ, Bae YA, Min KO, Kwon MS, Kim KM and Rhyu MG: Genetic classification of colorectal cancer based on chromosomal loss and microsatellite instability predicts survival. *Clin Cancer Res* 8: 2311-2322, 2002.
- 9 Diep CB, Thorstensen L, Meling GI, Skovlund E, Rognum TO and Lothe RA: Genetic tumor markers with prognostic impact in Dukes' stages B and C colorectal cancer patients. *J Clin Oncol* 21: 820-829, 2003.
- 10 Jen J, Kim H, Piantadosi S, Liu ZF, Levitt RC, Sistonen P, Kinzler KW, Vogelstein B and Hamilton SR: Allelic loss of chromosome 18q and prognosis in colorectal cancer. *N Engl J Med* 331: 213-221, 1994.
- 11 Ogunbiyi OA, Goodfellow PJ, Herfarth K, Gagliardi G, Swanson PE, Birnbaum EH, Read TE, Fleshman JW, Kodner IJ and Moley JF: Confirmation that chromosome 18q allelic loss in colon cancer is a prognostic indicator. *J Clin Oncol* 16: 427-433, 1998.
- 12 Martinez-Lopez E, Abad A, Font A, Monzo M, Ojanguren I, Pifarre A, Sanchez JJ, Martin C and Rosell R: Allelic loss on chromosome 18q as a prognostic marker in stage II colorectal cancer. *Gastroenterology* 114: 1180-1187, 1998.
- 13 Jernvall P, Makinen MJ, Karttunen TJ, Makela J and Vihko P: Loss of heterozygosity at 18q21 is indicative of recurrence and therefore poor prognosis in a subset of colorectal cancers. *Br J Cancer* 79: 903-908, 1999.
- 14 Font A, Abad A, Monzo M, Sanchez JJ, Guillot M, Manzano JL, Pinol M, Ojanguren I and Rosell R: Prognostic value of K-ras mutations and allelic imbalance on chromosome 18q in patients with resected colorectal cancer. *Dis Colon Rectum* 44: 549-557, 2001.
- 15 Watanabe T, Wu TT, Catalaño PJ, Ueki T, Satriano R, Haller DG, Benson AB III and Hamilton SR: Molecular predictors of survival after adjuvant chemotherapy for colon cancer. *N Engl J Med* 344: 1196-1206, 2001.
- 16 Massa MJ, Iniesta P, Gonzalez-Quevedo R, de Juan C, Caldes T, Sanchez-Pernaute A, Cerdan J, Torres AJ, Balibrea JL and Benito M: Differential prognosis of replication error phenotype and loss of heterozygosity in sporadic colorectal cancer. *Eur J Cancer* 35: 1676-1682, 1999.
- 17 Cohn KH, Ornstein DL, Wang F, LaPaix FD, Phipps K, Edelsberg C, Zuna R, Mott LA and Dunn JL: The significance of allelic deletions and aneuploidy in colorectal carcinoma. Results of a 5-year follow-up study. *Cancer* 79: 233-244, 1997.
- 18 Forslund A, Lonnroth C, Andersson M, Brevinge H and Lundholm K: Mutations and allelic loss of p53 in primary tumor DNA from potentially cured patients with colorectal carcinoma. *J Clin Oncol* 19: 2829-2836, 2001.
- 19 Barratt PL, Seymour MT, Stenning SP, Georgiades I, Walker C, Birbeck K and Quirke P: DNA markers predicting benefit from adjuvant fluorouracil in patients with colon cancer: a molecular study. *Lancet* 360: 1381-1391, 2002.
- 20 Carethers JM, Hawn MT, Greenson JK, Hitchcock CL and Boland CR: Prognostic significance of allelic loss at chromosome 18q21 for stage II colorectal cancer. *Gastroenterology* 114: 1188-1195, 1998.
- 21 Rooney PH, Boonsong A, McKay JA, Marsh S, Stevenson DA, Murray GI, Curran S, Haites NE, Cassidy J and McLeod H: Colorectal cancer genomics: evidence for multiple genotypes which influence survival. *Br J Cancer* 85: 1492-1498, 2001.
- 22 Yana I, Kurahashi H, Nakamori S, Kameyama M, Nakamura T, Takami M, Mori T, Takai S and Nishisho I: Frequent loss of heterozygosity at telomeric loci on 22q in sporadic colorectal cancers. *Int J Cancer* 60: 174-177, 1995.
- 23 Castells A, Gusella JF, Ramesh V and Rustgi AK: A region of deletion on chromosome 22q13 is common to human breast and colorectal cancers. *Cancer Res* 60: 2836-2839, 1995.
- 24 Zhou CZ, Peng ZH, Zhang F, Qiu GQ and He L: Loss of heterozygosity on long arm of chromosome 22 in sporadic colorectal carcinoma. *World J Gastroenterol* 8: 668-673, 2002.
- 25 Blaker H, Graf M, Rieker RJ and Otto HF: Comparison of losses of heterozygosity and replication errors in primary colorectal carcinomas and corresponding liver metastases. *J Pathol* 188: 258-262, 1999.
- 26 Sugano K, Nakashima Y, Yamaguchi K, Fukayama N, Maekawa M, Ohkura H, Kakizoe T and Sekiya T: Sensitive detection of loss of heterozygosity in the TP53 gene in pancreatic adenocarcinoma by fluorescence-based single-strand conformation polymorphism analysis using blunt-end DNA fragments. *Genes Chromosomes Cancer* 15: 157-164, 1996.
- 27 Sugano K, Tsutsumi M, Nakashima Y, Yamaguchi K, Ohkura H, Kakizoe T and Sekiya T: Diagnosis of bladder cancer by analysis of the allelic loss of the p53 gene in urine samples using blunt-end single-strand conformation polymorphism. *Int J Cancer* 74: 403-406, 1997.
- 28 Lengauer C, Kinzler KW and Vogelstein B: Genetic instability in colorectal cancers. *Nature* 386: 623-627, 1997.
- 29 Adachi Y, Yasuda K, Kakisako K, Sato K, Shiraishi N and Kitano S: Histopathologic criteria for local excision of colorectal cancer: multivariate analysis. *Ann Surg Oncol* 6: 385-388, 1999.
- 30 Nascimbeni R, Burgart LJ, Nivatvongs and Larson DR: Risk of lymph node metastasis in T1 carcinoma of the colon and rectum. *Dis Colon Rectum* 45: 200-206, 2002.
- 31 Iino H, Fukayama M, Maeda Y, Koike M, Mori T, Takahashi T, Kikuchi-Yanoshita R, Miyaki M, Mizuno S and Watanabe S: Molecular genetics for clinical management of colorectal carcinoma. 17p, 18q, and 22q loss of heterozygosity and decreased DCC expression are correlated with the metastatic potential. *Cancer* 73: 1324-1331, 1994.

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and Other Interventional Techniques

## Reduction of prolonged postoperative hospital stay after laparoscopic surgery for colorectal carcinoma

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### Abstract

**Background:** In evaluating the quality of laparoscopic surgery (LS) for colorectal carcinoma, many previous reports have used median or range values to assess the length of postoperative hospital stay and to show the complication and conversion rates separately. However, with this method, it is impossible to assess the proportion of patients who required prolonged postoperative hospital stay because of perioperative morbidities. This study investigated the proportion of patients who benefited from LS as minimally invasive surgery by assessing the percentage of patients who required prolonged postoperative hospital stay because of major perioperative morbidities.

**Methods:** A review of 202 patients who underwent LS for colorectal carcinoma at the authors' hospital between January 2002 and December 2004 was performed. Short-term outcomes were compared among the patients who underwent LS in 2002, 2003, and 2004.

**Results:** No significant differences were observed in baseline characteristics among the groups, and all the procedures in this study were completed laparoscopically. There were no significant differences in the operative times and intraoperative blood losses among the groups. Most of the patients resumed liquid intake on postoperative day 1 and solid food on day 3. However, there was a significant difference in the rate of postoperative prolonged hospital stays by year of surgery. In 2004, 97.3% of the patients (72/74) undergoing LS could be discharged to home within 8 days postoperatively. Major complications occurred at a low rate of 1.4% (1/74) in 2004. Regarding the reasons for prolonged postoperative hospital stay, inappropriate judgment of the physician in charge, based primarily on requests from patients without medical necessity, disappeared in 2004.

**Conclusions:** When LS is performed properly by specialists who have accumulated sufficient experience in

both LS and conventional open surgery for colorectal carcinoma, up to 97% of patients undergoing LS can benefit from minimally invasive surgery.

**Key words:** Colorectal carcinoma — Complication — Laparoscopic surgery — Postoperative hospital stay — Short-term outcome

In many randomized and nonrandomized studies comparing laparoscopic surgery (LS) and conventional open surgery for colorectal carcinoma, several advantages of LS have been reported, including reduction of postoperative pain, shortened duration of postoperative ileus, shortened hospital stay, and favorable effects on cytokine and hormonal responses. Consequently, LS is now termed “minimally invasive surgery” [1, 10, 15–17].

At our institution, much consideration has been given to the technical and oncologic safety of LS. Since our first LS for colorectal carcinoma in 1993, approximately 400 LS for colorectal malignancies have been performed at our institution. Most of our early experience was confined to early (Tis or T1) colorectal cancer located at the cecum, ascending colon, sigmoid colon, or rectosigmoid because of technical problems and concerns regarding port-site and peritoneal recurrences. In June 2001, we unified our surgical and postoperative management procedures and expanded our indications for LS to include advanced colorectal cancers (i.e., T2 lesions and beyond) located anywhere in the colon or rectum. As a consequence, the complication rate and mean length of hospitalization have been reduced at our institution [23, 24].

If LS is truly minimally invasive surgery, it should reflect a shortened postoperative hospital stay. With regard to assessment of the quality of LS for colorectal carcinoma, many previous reports have used median or range values to assess the length of postoperative hospital stay and to show the complication and conversion rates separately. However, with this approach, it is

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impossible to assess the proportion of patients who required prolonged postoperative hospital stay because of perioperative morbidities. Moreover, the effect of major complications on the length of postoperative hospitalization is unknown.

In the current study, short-term outcomes were compared among selected patients who underwent LS for colorectal carcinoma at our hospital in 2002, 2003, and 2004. We investigated the proportion of patients who benefited from LS as minimally invasive surgery by assessing the percentage of patients who required prolonged postoperative hospital stay because of perioperative major morbidities. Moreover, the results of our efforts to reduce postoperative hospitalization also were examined.

## Patients and methods

### Patients

Between January 2002 and December 2004, we performed 202 continuous LS for selected patients with colorectal carcinoma. Because the safety of LS for cancer patients remains to be established, our candidates for radical LS were patients with a preoperative diagnosis of T1 or T2. Additionally, our LS cases also included patients with a preoperative diagnosis of T3 who wished to undergo LS, as well as those with colon or upper rectal carcinoma for which palliative resection was considered necessary. Contraindications for LS included tumors larger than 6 cm, a history of extensive adhesions, severe obesity (body mass index  $> 32 \text{ kg/m}^2$ ), intestinal obstruction, and refusal of a patient to undergo LS.

All patients were evaluated before surgery by clinical investigation including barium enema, total colonoscopy, chest x-ray, abdominal ultrasonography, and computed tomography. For patients with rectal malignancy, a primary rectal carcinoma was defined according to its distance from the anal verge, as determined by colonoscopy. The tumors were grouped according to their location in the lower rectum (0–7 cm), the middle rectum (7.1–12 cm), and the upper rectum (12.1–17 cm). We defined conversion to open surgery as any incision larger than 7cm, except for cases in which the incision was enlarged because of a large specimen that could not be removed through a 7-cm incision.

### Laparoscopic technique

The techniques for LS have been thoroughly described previously [23–25]. For right-sided lesions, the right colon was mobilized initially, and the vascular pedicles were divided at their origin, together with the draining lymph nodes intracorporeally. For patients with a preoperative diagnosis of T2–T3 lesions, a laparoscopic no-touch isolation technique was performed. With this technique, mobilization of the right colon was performed after early proximal ligation of the tumor-feeding vessels and resection of the mesentery intracorporeally. The bowel loop was delivered under a wound protector through a small incision, and division of the marginal vessels and anastomosis was performed extracorporeally.

For transverse colon lesions, mobilization of hepatic, splenic, or both flexures was performed according to the tumor location. Proximal ligation of the right, left, or both branches of the middle colic vessels at their origins was performed intracorporeally or extracorporeally. The bowel loop was delivered, and anastomosis was performed in the same way.

Descending colon and proximal sigmoid colon lesions for which extracorporeal anastomosis was considered possible were managed by initial mobilization of the left colon. After mobilization of the splenic flexure, intracorporeal ligation of the tumor-feeding vessels (left colic artery, sigmoid arteries, inferior mesenteric vessels) at their origins was performed. The bowel loop was delivered through a small incision under a wound protector, and division of the mesentery was performed extracorporeally, followed by extracorporeal anastomosis.

For distal sigmoid colon and rectal lesions, mobilization of left colon and splenic flexure, if necessary, was followed by intracorporeal high ligation of the inferior mesenteric vessels, then by mobilization of the rectum. For higher lesions, mesorectal tissue down to 5 cm below the tumor was excised routinely. Middle and lower rectal tumors were treated by total mesorectal excision. Before rectal transection, laparoscopic rectal clamping immediately above the anticipated point of rectal transection was performed using a bowel-clamping device introduced through the 12-mm mid lower port. Rectal washout was routinely performed using 1,000 ml of 5% povidone-iodine solution. Rectal transection then was performed by the multiple firing technique using Endo GIA Universal staples (Auto Suture; U.S. Surgical Corp., Norwalk, CT, USA) introduced through the 12-mm right midabdominal port. A 4- to 5-cm incision then was made over the mid lower port site, and the bowel was exteriorized under wound protection.

After the anvil head of the circular stapler had been inserted into the end of the proximal colon, the proximal colon was internalized and the incision closed. Intracorporeal anastomosis under laparoscopic view was performed by the double-stapling technique using a circular stapler (ECS 29 or 33mm; Ethicon Endo-Surgery Inc, Cincinnati, OH, USA). For patients with lesions located within 2 cm of the dentate line, laparoscopic intersphincteric rectal resection and hand-sewn coloanal anastomosis (ISR-CAA) were performed [21]. For patients undergoing abdominoperineal resection (APR), laparoscopic procedures were followed by perineal dissection in the standard fashion and end colostomy creation using the left lower abdominal port site.

### Study parameters

The parameters analyzed included gender, age, body mass index, prior abdominal surgery, operative time, intraoperative blood loss, conversion rate, days to resumption of diet, duration of postoperative hospital stay, and both intraoperative and postoperative complications within 30 days of surgery. Pathologic staging was performed according to Dukes' stage. In the current study, major complication was defined as morbidity that required the patient to stay in the hospital 9 or more postoperative days. Prolonged postoperative hospital stay was defined as 9 or more days of postoperative hospitalization, regardless of the underlying reasons, because patients are supposed to be discharged by the postoperative day 8 when there is no major complication after LS at our institution. With regard to the operative and postoperative results, patients with colon and rectal carcinoma were evaluated separately, considering the technical difficulties of the laparoscopic procedure.

### Statistical analysis

Statistical analysis was performed using one-way analysis of variance (ANOVA) and chi-square testing as appropriate. A *p* value less than 0.05 was considered significant.

## Results

The demographics for the patients in this study are summarized in Table 1. There were no significant differences in baseline characteristics among the groups. However, the proportion of the patients with Dukes' B, C, and D stages scheduled for LS is increasing, although the difference is not yet significant (*p* = 0.093). With regard to simultaneously performed surgical techniques, one patient underwent resection of a benign submandibular gland tumor in 2002, and three patients underwent laparoscopic cholecystectomy in 2003. In 2004, three patients underwent combined surgery as follows: laparoscopic enucleation of an 8-cm hysteromyoma, partial resection of the lingua for carcinoma, and hemilateral neck lymph node dissection for

**Table 1.** Patient characteristics<sup>a</sup>

	2002 (n = 59)	2003 (n = 69)	2004 (n = 74)	p Value
No. of patients	59	69	74	
Sex ratio (male:female)	35:24	39:30	37:37	0.533
Age (years)	58.5 (30–83)	60.2 (38–88)	61.1 (34–79)	0.360
Body mass index (kg/m <sup>2</sup> )	22.9 (14.9–32.4)	23.1 (17.3–30.5)	23.1 (16.3–31.5)	0.872
Prior abdominal surgery: n (%)	15 (25.4)	16 (23.2)	16 (21.6)	0.875
Dukes' stage (n)				
A	45	52	46	
B	2	2	9	
C	8	11	18	
D	4	4	2	
A:B + C + D	45:14	52:17	46:29	0.093
Location (n)				
Cecum	3	10	5	
Ascending colon	4	13	15	
Transverse colon	6	9	7	
Descending colon	6	5	7	
Sigmoid colon	23	20	27	
Rectosigmoid/upper rectum	10	7	4	
Middle rectum	3	3	4	
Lower rectum	4	2	6	
Colon:rectum	42:12	57:12	60:14	0.238
Laparoscopic colorectal procedures (n)				
Ileocecal resection	5	7	7	
Right hemicolectomy	5	18	14	
Transverse colectomy	4	4	4	
Left hemicolectomy	0	1	1	
Descending colectomy	5	4	6	
Sigmoid colectomy	20	17	24	
Partial resection	3	6	5	
Anterior resection with DST	16	10	12	
Anterior resection with ISR-CAA	1	2	1	
Abdominoperineal resection	0	0	1	
Transverse-colooplasty pouch	0	2	2	
Covering ileostomy	4	2	5	

DST, double-stapling technique; ISR-CAA, intersphincteric rectal resection and handsewn coloanal Anastomosis

<sup>a</sup> Values are mean (range)

metachronous lymph node recurrence from lingual carcinoma. Data on these combined surgical techniques all were included in the analyses of colorectal carcinoma surgeries.

Our operative results are shown in Table 2. All the procedures in this study were completed laparoscopically. There were no significant differences in operative time or intraoperative blood loss among the groups. The postoperative courses are shown in Table 3. Most of the patients started liquid intake on postoperative day 1 and solid food on day 3. However, there was a significant difference in the rates of prolonged postoperative hospital stay by year of surgery. All the patients were discharged to home.

The postoperative complications are listed in Table 4. There were no perioperative mortalities. No significant differences in complication rates over the years were observed, although a major complication, anastomotic leakage, occurred for one patient in 2004 and was successfully treated conservatively. None of the patients in the current series required reoperation.

The reasons for prolonged postoperative hospital stays are listed in Table 5. Inappropriate judgment of the physician in charge, based primarily on requests from patients without medical necessity, disappeared in

**Table 2.** Operative results<sup>a</sup>

	2002 (n = 59)	2003 (n = 69)	2004 (n = 74)	p Value
Lap colectomy				
Operative time (min)	201 (115–345)	200 (117–348)	214 (140–495)	0.219
Intraoperative blood loss (ml)	30 (6–219)	30 (10–248)	38 (7–256)	0.157
Conversion (n)	0	0	0	
Lap-AR + APR				
Operative time (min)	244 (190–392)	263 (200–472)	283 (215–430)	0.570
Intraoperative blood loss (ml)	54 (10–265)	63 (11–250)	84 (14–477)	0.661
Conversion (n)	0	0	0	

Lap, laparoscopic; AR, anterior resection; APR, abdominoperineal resection

<sup>a</sup> Values are medians (range)

2004. By the end of the study period, two patients had experienced recurrence (hepatic metastases). At this writing, in 2005, 49 patients have undergone LS, and all have been discharged to home without major complication.

**Table 3.** Postoperative results

	2002 (n = 59) n (%)	2003 (n = 69) n (%)	2004 (n = 74) n (%)	p Value
<b>Lap colectomy</b>				
Liquid intake range (days)				
1 POD	38 (90.4)	54 (94.7)	59 (98.3)	
2 POD	2 (4.8)	3 (5.3)	0 (0)	
3 ≤ POD	2 (4.8)	0 (0)	1 (1.7)	
Solid food (days)				
2 POD	0 (0)	0 (0)	0 (0)	
3 POD	31 (73.8)	51 (89.5)	56 (93.3)	
4 ≤ POD	11 (26.2)	6 (10.5)	4 (6.7)	
Hospital stay (days)				
7 POD	5 (11.9)	19 (33.3)	28 (46.7)	
8 POD	17 (40.5)	28 (49.1)	31 (51.7)	
9 ≤ POD	20 (47.6)	10 (17.5)	1 (1.7)	
Range	7–20	7–15	7–21	
<b>Lap-AR + APR</b>				
Liquid intake range (days)				
1 POD	16 (94.1)	10 (83.3)	13 (92.9)	
2 POD	1 (5.9)	1 (8.3)	0 (0)	
3 ≤ POD	0 (0)	1 (8.3)	1 (7.1)	
Solid food (days)				
2 POD	3 (17.6)	1 (8.3)	3 (21.4)	
3 POD	6 (35.3)	9 (75.0)	9 (64.3)	
4 ≤ POD	8 (47.1)	2 (16.7)	2 (14.3)	
Hospital stay (days)				
7 POD	2 (11.8)	4 (33.3)	5 (35.7)	
8 POD	3 (17.6)	6 (50.0)	8 (57.1)	
9 ≤ POD	12 (70.6)	2 (16.7)	1 (7.1)	
Range	7–12	7–17	7–23	
<b>Total (Lap-colectomy + AR + APR)</b>				
Hospital stay (days)				
7–8 POD	27 (45.8)	57 (82.6)	72 (97.3)	< 0.0001
9 ≤ POD	32 (54.2)	12 (17.4)	2 (2.7)	

Lap, laparoscopic; POD, postoperative days; AR, anterior resection; APR, abdominoperineal resection

## Discussion

To date, the quality of LS for colorectal carcinoma has been assessed by the median or range of the postoperative hospital stay, the complication rate, and the conversion rate. However, with only these values, it is impossible to assess accurately the degree of the effect that each complication has on the length of postoperative hospitalization for patients overall. This means that the rate of patients undergoing LS who have benefited from minimally invasive surgery has not been properly evaluated. If the greatest advantage of LS is minimal invasiveness, LS must ultimately be linked to shortened postoperative hospitalization. However, no reports have focused on the rate of reduction in the length of postoperative hospital stay after LS. In our hospital, patients are supposed to be discharged after LS until postoperative day 8. As experience with LS cases accumulated, surgical and postoperative management procedures became unified. The timing for the start of solid food intake became earlier in 2004 than in 2002, and this may have contributed to the significantly shortened period of postoperative hospitalization in 2004. Furthermore, major complications that required prolonged postoperative hospital stay were reduced. As a result, in 2004, 97.3% (72/74) of our patients undergoing LS could be discharged to home within 8 days postoperatively. Major complications occurred at a low rate of 1.4% (1/74) in 2004. Needless to say, this low rate contributed greatly to the current results.

The current report deals with the length of postoperative hospital stay. Recent reports from randomized controlled trials (RCTs) and single institutions investigating a number of cases in western countries indicate that the median or mean length of hospital stay after LS for colorectal carcinoma ranges from 5 to 9 days [1, 2, 6, 7, 9, 14, 16]. The appearance of this range may be attributable to social factors such as differences in medical fees, medical insurances, and medical systems among countries rather than differences in the quality of surgery. According to former studies from Japan, Japanese patients tend to stay in the hospital longer than patients in western countries [20]. The reasons for this tendency include the following facts. From the perspective of patients in Japan, public health insurance covers 70% of the total medical cost for every patient, with the patients paying only 30% of the cost. Socially disadvantaged people do not have to bear their medical expense no matter how many days they are hospitalized. Furthermore, for a patient undergoing surgery, the cost of surgery accounts for the greater part of the total medical cost. Hence, if the duration of hospital stay is lengthened by 1 day, the patient pays only several tens of dollars in additional cost. Furthermore, many Japanese patients have private health insurance, which pays the patient a specified amount of money per day of hospitalization. Under some types of insurance contract, the longer the patient stays in hospital, the more the insurance dividend will be, thereby yielding greater “earn-

**Table 4.** Morbidities and mortality<sup>a</sup>

	2002 (n = 59)	2003 (n = 69)	2004 (n = 74)	p Value
Lap colectomy				
Mortality	0	0	0	
Morbidity				
Wound sepsis	3 (1)	2	6	
Bowel obstruction	2 (1)	3 (1)	1	
Urinary tract infection	2 (2)	1	0	
Pneumonia	0	1 (1)	0	
Pneumothorax	0	1 (1)	0	
Pulmonary embolism	0	1 (1)	0	
Enterocolitis	0	1	0	
Total	7 (4)	10 (4)	7 (0)	
Reoperation	0	0	0	
Readmissions	1	3	0	
Lap-AR + APR				
Mortality	0	0	0	
Morbidity				
Wound sepsis	0	2	1	
Bowel obstruction	0	1 (1)	0	
Anastomotic leakage	0	0	1 (1)	
Abscess	0	1 (1)	0	
Pneumonia	0	0	1	
Neurogenic bladder	1 (1)	0	0	
Total	1 (1)	4 (2)	3 (1)	
Reoperation	0	0	0	
Readmissions	0	0	0	
Total complication: n (%)	8 (13.5)	14 (20.3)	10 (13.5)	0.4595
Major complication: n (%)	5 (8.5)	6 (8.7)	1 (1.4)	0.111

Lap, laparoscopic; AR, anterior resection; APR, abdominoperineal resection

No. of major complications in parentheses

**Table 5.** Reasons for prolonged postoperative hospital stay

	2002 (n = 32)	2003 (n = 12)	2004 (n = 2)
Major complication	5	6	1
Others			
Treatment for comorbid disease	3	1	1
Ileostomy management	4	0	0
Inappropriate judgment of the physician in charge	20	5	0

ings.” Under these circumstances, patients do not need to be discharged from the hospital quickly.

In contrast, at our institution, if there is no major complication after LS, the patient is supposed to be discharged by postoperative day 8, but no patients wished to leave hospital earlier than postoperative day 7. Furthermore, the results of the current study in terms of postoperative stay after LS for colorectal carcinoma are some of the shortest reported in Japan. Obviously, this situation in Japan is wasting medical funds. It goes without saying that the situation must be improved. From the results of the current study, we consider that the appropriate duration of postoperative hospital stay after LS is 5 to 7 days. Early discharge within 5 days might be possible for some patients. However, it is necessary to confirm the safety of early discharge, especially for patients with rectal carcinoma who have

undergone anterior resection, because for some patients, fatal complications accompanied by anastomotic leakage might occur approximately 7 days after surgery.

With regard to the oncologic outcome of LS for colorectal carcinoma, recently reported RCTs have demonstrated that LS is comparable with open surgery or superior to it [6, 12, 14]. The results of some other RCTs to be published in the near future also are attracting attention. However, in Japan, RCTs for gastrointestinal malignancies have not been widely accepted in the past because of concerns about consequences if one form of treatment is shown to be inferior to the other. For this reason, a prospective multicenter trial with patients undergoing laparoscopic colectomy for advanced carcinoma has not been performed. However, fertile ground for RCTs comparing surgical techniques has finally begun to develop among both patients and surgeons in Japan. Consequently, a multicenter RCT comparing LS and open surgery outcomes for advanced colorectal cancer was begun in 2004 [11]. The distinctive features of this trial are that all the participating institutions have sufficient experience not only in open surgery, but also in LS, and that D3 lymphadenectomy is being required as a rule for all patients because this has been regarded as the standard surgical procedure for advanced colorectal carcinoma in both LS and open surgery. The results of this Japanese study will be published later than those of similar studies in western countries, but this study is receiving attention as an RCT performed by surgeons with sufficient accumulated experience in LS and specializing in colorectal carcinoma surgery using LS.

The issue in the expansion of the indications for LS for colorectal carcinoma is whether it can be performed for patients with middle or lower rectal carcinoma. The technical difficulty of surgery is high in such cases. If the rate of conversion to open surgery increases, the short-term outcomes of LS will be shifted to the outcomes of open surgery, thus making it difficult to detect differences between the two surgeries [9, 22]. In addition, if the complication rate increases, it could lead to prolonged postoperative hospitalization, thereby canceling the advantages of LS. The most important issue is whether LS can yield a treatment outcome comparable with that of open surgery in cases of advanced rectal cancer. The most difficult complication to manage is anastomotic leakage. In cases of middle and lower rectal carcinoma, the occurrence of anastomotic leakage requires not only prolonged hospitalization, but also a temporary or permanent stoma in some patients, thereby resulting in an unavoidable deterioration in quality of life. Moreover, anastomotic leakage may cause fatal peritonitis, or may promote intrapelvic recurrence in some cases [4]. However, with regard to the technical issue, as shown by the results of this study and recently published papers, when surgeons with sufficient experience in LS for rectal carcinoma are in charge of the procedure, it can be performed successfully as minimally invasive surgery in cases of middle and lower rectal carcinoma [2, 3, 5, 8, 13, 18, 19, 24].

In the previous report from our institution, short-term outcomes were compared between patients with

colon carcinoma and those with rectal carcinoma, all of whom underwent LS. The complication rates and postoperative courses between the two approaches were similar [23]. Needless to say, in cases of middle and lower rectal carcinoma as well, further investigations in multicenter RCTs are needed regarding short- and long-term outcomes.

One distinctive feature of LS for rectal carcinoma at our institution is that only one patient underwent APR. The background factor behind this is that whether we select open surgery or LS, we usually perform ISR-CAA in T1–T2 cases of lower rectal carcinoma, and also in many T3 cases if the patient expresses a request. Recently, favorable oncologic and functional outcomes of ISR-CAA have been reported, and the number of patients undergoing ISR-CAA using LS is expected to increase in the coming years [18, 21]. Only one patient who underwent laparoscopic APR in 2004 was preoperatively judged to be a candidate for ISR-CAA by LS. However, that patient's choice was APR.

The mean operative time in the current study was slightly longer than that reported in previous studies. This may be partially because of gradual increases in the proportion of patients with relatively advanced stages of disease who underwent LS. Other reasons might be that trainee doctors perform part or all of the surgical procedure under the guidance of staff doctors in many cases, and that we have been unable to establish a laparoscopic team. However, it is evident that the quality of our operations has not been lowered, as demonstrated by the results of this study.

In conclusion, the surgical outcome for LS at our institution demonstrated that when LS is performed properly by specialists who have accumulated sufficient experience in both LS and open surgery for colorectal carcinoma, up to 97% of patients undergoing LS can benefit from minimally invasive surgery. To expand the use of minimally invasive surgery for advanced colorectal carcinoma, it goes without saying that while making efforts to acquire high-level technical skills, it is also necessary to confirm the oncologic safety of LS.

## References

1. Abraham NS, Young JM, Solomon MJ (2004) Meta-analysis of short-term outcomes after laparoscopic resection for colorectal cancer. *Br J Surg* 91: 1111–1124
2. Anthuber M, Fuerst A, Elser F, Berger R, Jauch KW (2003) Outcome of laparoscopic surgery for rectal cancer in 101 patients. *Dis Colon Rectum* 46: 1047–1053
3. Bärlechner E, Benhidjeb T, Anders S, Schicke B (2005) Laparoscopic resection for rectal cancer: outcomes in 194 patients and review of the literature. *Surg Endosc* 19: 757–766
4. Branagan G, Finnis D (2005) Prognosis after anastomotic leakage in colorectal surgery. *Dis Colon Rectum* 48: 1021–1026
5. Bretagnol F, Lelong B, Laurent C, Moutardier V, Rullier A, Monges G, Delperio JR, Rullier E (2005) The oncological safety of laparoscopic total mesorectal excision with sphincter preservation for rectal carcinoma. *Surg Endosc* 19: 892–896
6. Clinical Outcomes of Surgical Therapy Study Group (2004) A comparison of laparoscopically assisted and open colectomy for colon cancer. *N Engl J Med* 350: 2050–2059
7. COLOR Study Group (2005) Impact of hospital case volume on short-term outcome after laparoscopic operation for colonic cancer. *Surg Endosc* 19: 687–692
8. Delgado S, Momblán D, Salvador L, Bravo R, Castells A, Ibarzabal A, Piqué JM, Lacy AM (2004) Laparoscopic-assisted approach in rectal cancer patients: lessons learned from >200 patients. *Surg Endosc* 18: 1457–1462
9. Guillou PJ, Quirke P, Thorpe H, Walker J, Jayne DG, Smith AMH, Heath RM, Brown JM, MRC CLASICC Trial Group (2005) Short-term end points of conventional versus laparoscopic-assisted surgery in patients with colorectal cancer (MRC CLASICC trial): multicentre, randomized, controlled trial. *Lancet* 365: 1718–1726
10. Hasegawa H, Kabeshima Y, Watanabe M, Yamamoto S, Kitajima M (2003) Randomized controlled trial of laparoscopic versus open colectomy for advanced colorectal cancer. *Surg Endosc* 17: 636–640
11. Kitano S, Inomata M, Sato A, Yoshimura K, Moriya Y, Colorectal Cancer Study Group (CCSG) of Japan Clinical Oncology Group (2005) Randomized controlled trial to evaluate laparoscopic surgery for colorectal cancer: Japan Clinical Oncology Group study JCOG 0404. *Jpn J Clin Oncol* 35: 475–477
12. Lacy AM, García-Valdecasas JC, Delgado S, Castells A, Taurá P, Piqué JM, Visa J (2002) Laparoscopic-assisted colectomy versus open colectomy for treatment of non-metastatic colon cancer: a randomized trial. *Lancet* 359: 2224–2229
13. Leroy J, Jamali F, Forbes L, Smith M, Rubino F, Mutter D, Marescaux J (2004) Laparoscopic total mesorectal excision (TME) for rectal cancer surgery: long-term outcomes. *Surg Endosc* 18: 281–289
14. Leung KL, Kwok SPY, Lam SCW, Lee JFY, Yiu RYC, Ng SSM, Lai PBS, Lau WY (2004) Laparoscopic resection of rectosigmoid carcinoma: prospective randomised trial. *Lancet* 363: 1187–1192
15. Leung KL, Lai PBS, Ho RLK, Meng WCS, Yiu RYC, Lee JFY, Lau WY (2000) Systemic cytokine response after laparoscopic-assisted resection of rectosigmoid carcinoma: a prospective randomized trial. *Ann Surg* 231: 506–511
16. Milsom JW, Böhm B, Hammerhofer KA, Fazio V, Steiger E, Elson P (1998) A prospective, randomized trial comparing laparoscopic versus conventional techniques in colorectal cancer surgery: a preliminary report. *Am Coll Surg* 187: 46–57
17. Nishiguchi K, Okuda J, Toyoda M, Tanaka K, Tanigawa N (2001) Comparative evaluation of surgical stress of laparoscopic and open surgeries for colorectal carcinoma. *Dis Colon Rectum* 44: 223–230
18. Rullier E, Sa Cunha A, Couderc P, Rullier A, Gontier R, Saric J (2003) Laparoscopic intersphincteric resection with coloplasty and coloanal anastomosis for mid and low rectal cancer. *Br J Surg* 90: 445–451
19. Tsang WWC, Chung CC, Kwok SY, Li MKW (2005) Minimally invasive surgery for rectal cancer. *Surg Clin North Am* 85: 61–73
20. Uchiyama K, Takifuji K, Tani M, Onishi H, Yamaue H (2002) Effectiveness of the clinical pathway to decrease length of stay and cost for laparoscopic surgery. *Surg Endosc* 16: 1594–1597
21. Watanabe M, Teramoto T, Hasegawa H, Kitajima M (2000) Laparoscopic ultralow anterior resection combined with per anum intersphincteric rectal dissection for lower rectal cancer. *Dis Colon Rectum* 43: S94–S97
22. Weeks JC, Nelson H, Gelber S, Sargent D, Schroeder G, Clinical Outcomes of Surgical Therapy (COST) Study Group (2002) Short-term quality-of-life outcomes following laparoscopic-assisted colectomy vs open colectomy for colon cancer: a randomized trial. *JAMA* 287: 321–328
23. Yamamoto S, Fujita S, Akasu T, Moriya Y (2004) A comparison of the complication rates between laparoscopic colectomy and laparoscopic low anterior resection. *Surg Endosc* 18: 1447–1451
24. Yamamoto S, Fujita S, Akasu T, Moriya Y (2005) Safety of laparoscopic intracorporeal rectal transection with double-stapling technique anastomosis. *Surg Laparosc Endosc Percutan Tech* 15: 70–74
25. Yamamoto S, Watanabe M, Hasegawa H, Kitajima M (2002) Prospective evaluation of laparoscopic surgery for rectosigmoidal and rectal carcinoma. *Dis Colon Rectum* 45: 1648–1654



# The Risk of Multiple Primary Malignancies with Colorectal Carcinoma

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**PURPOSE:** With advances in diagnostic techniques and treatment modalities, the number of patients identified with colorectal carcinoma who develop multiple primary malignancy during long-term follow-up has been increasing. We investigated multiple primary malignancies occurring in a large number of colorectal carcinoma patients who had undergone surgery in the 1980s at our institution. **METHODS:** A total of 1,304 Japanese patients with colorectal carcinoma treated between January 1980 and December 1989 were prospectively followed to investigate the situations in which multiple primary malignancies occurred. To determine whether the incidence of multiple primary malignancies in this series was higher than expected, we calculated the expected numbers of carcinoma occurrences and evaluated these findings by exact binomial test. **RESULTS:** The median follow-up period was 95 months. The incidence of multiple primary malignancy was 18.7 percent (143/765) among males and 14.7 percent (79/539) among females. The most common site of multiple primary malignancy among males was the stomach, followed by the lung, prostate, larynx, liver, esophagus, and urinary bladder. The most common site among females was the uterus, followed by the stomach, breast, and liver. The sites that showed a higher incidence of multiple primary malignancy than the expected value were: the prostate, larynx, urinary bladder, oral cavity/pharynx and thyroid among males, and the uterus and oral cavity/pharynx among females. **CONCLUSIONS:** Fifteen to 20 percent of Japanese colorectal carcinoma patients experienced multiple primary malignancies. Postoperative long-term screening methods should be established considering the actual occurrence numbers and risk rate of multiple primary malignancies in addition to metachronous colorectal carcinoma. [Key words: Colorectal carcinoma; Multiple

primary malignancy; Follow-up; Expected numbers of carcinoma occurrences]

The incidence rates of each organ carcinoma vary with the times, racial or ethnical groups, and countries. In terms of age-adjusted incidence rates in Japan, a recent report showed that the most common carcinoma among males is gastric carcinoma, followed by colorectal carcinoma and lung carcinoma; and the most common carcinoma among females is colorectal carcinoma, followed by breast carcinoma and gastric carcinoma.<sup>1</sup> Among those carcinomas, the incidence of colorectal carcinoma is rising among both males and females. With advances in diagnostic techniques and treatment modalities, the outcomes of colorectal carcinoma treatment have improved, whereas the number of patients who develop multiple primary malignancy during long-term follow-up has simultaneously increased. However, with regard to the incidence of concurrent colorectal carcinoma and multiple primary malignancy in Japanese patients, many previous reports have merely indicated the number of concurrences, and few reports have described the incidence of the concurrences in relation to patient age and follow-up period.<sup>2-5</sup>

Taking patient age and follow-up period into consideration, we investigated the situations in which multiple primary malignancies occurred during long-term follow-up of a large number of colorectal carcinoma patients who had received treatment in the 1980s at our institution. This paper reports the findings of the investigation.

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## PATIENTS AND METHODS

A total of 1,304 Japanese patients underwent colorectal carcinoma surgery at our institution between January 1980 and December 1989, and patient information and follow-up data were prospectively collected and added to the department database. In terms of follow-up, we routinely conducted periodic check-ups for the recurrence of colorectal carcinoma until the fifth postoperative year. No routine examinations were performed for multiple primary malignancies. Multiple primary malignancies were confirmed only when patients with multiple primary malignancies were diagnosed or treated at our institution, or documentation from other hospitals was obtained. The follow-up period was defined as the interval between the date of surgery for colorectal carcinoma and the date at which information regarding the occurrence or absence of multiple primary malignancies was confirmed. We defined metachronous and synchronous carcinomas according to the criteria used by Warren and Gates<sup>6</sup>; synchronous carcinoma was defined as tumors detected after an interval of less than one year, and metachronous carcinoma was defined as tumors detected after an interval of one year or longer. Fifteen patients with familial adenomatous polyposis were excluded, but six patients with hereditary nonpolyposis colorectal carcinoma (HNPCC) were included in this study.

### Statistical Analysis

To determine whether the incidence of multiple primary malignancies in this series was higher than the average incidence in Japan, we calculated the expected numbers of carcinoma occurrences by gender and tumor site for each of the following three periods: 1) from the date of birth to the date of surgery, 2) from the date of surgery to the final date of confirmation of survival, and 3) from the date of birth to the final date of confirmation; then we compared those expected numbers with the observed numbers.

The expected numbers of carcinoma incidences were computed by summing the cumulative risk of developing carcinoma for each patient during the period; those numbers were calculated based on the age-specific and gender-specific carcinoma incidence rates in Japan.<sup>1,7</sup> For example, the cumulative risk of stomach cancer from the date of surgery (1985) to

the final date of confirmation of survival (1995) for a female patient aged 60 years at surgery was obtained by the sum of the incidence rates of stomach cancer for females aged 60 years in 1985, that for females aged 61 years in 1986, . . . , and that for females aged 70 years in 1995. In the case of a period of less than one year, the probability was obtained by multiplying the incidence rate by the number of days per 365.25. The methods of estimating cancer incidence in Japan and their limitations have been explained in previous reports, and corrections were applied to minimize any possible bias.<sup>7-9</sup> The cancer incidence rates after 2000 and before 1974 were assumed to be equal to those of 1999 and 1975, respectively, because data before 1974 and after 2000 have not been published. The two-tail *P* value was calculated exactly based on binomial distribution (exact binomial test).

Clinicopathologic parameters, such as gender, age, location of tumor, Dukes stage, and presence or absence of adjuvant treatment were compared by using Student's *t*-test or the chi-squared test where appropriate. *P* < 0.05 was considered significant.

## RESULTS

### Patient Characteristics

The follow-up periods for all patients ranged from 1 to 269 (median, 95) months. The mortality rate for male patients was 51.9 percent (397/765), and that for female patients was 41.7 percent (225/539). The patient demographics are summarized in Table 1. The incidence of multiple primary malignancy was 18.7 percent (143/765) among males and 14.7 percent (79/539) among females, showing no difference between the two groups (*P* = 0.0614). A comparison between patients with only colorectal carcinoma (O) and patients with multiple primary malignancies (M) demonstrated that the mean age at the onset of colorectal carcinoma was significantly higher in the M group among both males and females (*P* < 0.0001, *P* = 0.0008, respectively). With regard to the locations of colorectal carcinoma, the proportion of M was significantly higher among male colon carcinoma patients (*P* = 0.0002), but there was no difference among females (*P* = 0.6277). Patients with a more advanced Dukes stage had a significantly lower proportion of M in both males and females (*P* < 0.0001, *P* = 0.0049, respectively). With regard to adjuvant treatment, 37 patients underwent adjuvant radiotherapy and no patients developed subsequent

**Table 1.**  
Characteristics of the Patients

Variable	Male (n = 765)		Female (n = 539)	
	Only Colorectal Carcinoma (n = 622)	Multiple Primary Malignancies (n = 143)	Only Colorectal Carcinoma (n = 460)	Multiple Primary Malignancies (n = 79)
Mean age at surgery for colorectal carcinoma (yr)	58.9 <sup>a</sup>	65 <sup>a</sup>	58.4 <sup>b</sup>	63.2 <sup>b</sup>
Synchronous		67.5		67.5
Metachronous—colorectal carcinoma preceding				
Age at colorectal carcinoma (yr)		61.2		59.4
Age at multiple primary malignancies (yr)		69.6		67.1
Metachronous—multiple primary malignancies preceding				
Age at multiple primary malignancies (yr)		58.1		53.5
Age at colorectal carcinoma (yr)		67.4		64.6
Location <sup>g</sup>				
Colon	272 <sup>c</sup>	87 <sup>c</sup>	229 <sup>d</sup>	37 <sup>d</sup>
Rectum	348 <sup>c</sup>	55 <sup>c</sup>	229 <sup>d</sup>	42 <sup>d</sup>
Dukes stage				
A	105 <sup>e</sup>	41 <sup>e</sup>	86 <sup>f</sup>	17 <sup>f</sup>
B	177 <sup>e</sup>	49 <sup>e</sup>	110 <sup>f</sup>	31 <sup>f</sup>
C	194 <sup>e</sup>	33 <sup>e</sup>	142 <sup>f</sup>	22 <sup>f</sup>
D	146 <sup>e</sup>	20 <sup>e</sup>	122 <sup>f</sup>	9 <sup>f</sup>

<sup>a</sup> $P < 0.0001$ ;

<sup>b</sup> $P = 0.0008$ ;

<sup>c</sup> $P = 0.0002$ ;

<sup>d</sup> $P = 0.6276$ ;

<sup>e</sup> $P < 0.0001$ ;

<sup>f</sup> $P = 0.0049$ .

<sup>g</sup>Five patients with synchronous or metachronous carcinoma of the colon and rectum were excluded from the analysis.

**Table 2.**  
Observed and Expected Number of Multiple Primary Malignancies in Males (n = 143)

Site	Total No. of Malignancies			Multiple Primary Malignancies Preceding and Synchronous			Colorectal Carcinoma Preceding		
	Obs	Exp	P Value	Obs	Exp	P Value	Obs	Exp	P Value
Stomach	59	54.7	0.5277	37	33.4	0.8596	22	20.9	0.7395
Lung	25	22.9	0.5957	13	10.2	0.0063	12	12.3	1
Prostate	12	5.5	0.0144	1	1.7	<0.001	11	3.6	0.0013
Larynx	11	2.1	<0.001	8	1.2	0.0066	3	1	0.0735
Liver	10	14	0.3441	3	6.9	0.2435	7	7	1
Esophagus	10	6.1	0.1468	5	3.2	0.0156	5	2.9	0.2198
Urinary bladder	10	4.9	0.0361	7	2.3	0.0283	3	2.6	0.7475
Oral cavity/pharynx	7	2.9	0.0274	4	1.6	0.0217	3	1.3	0.1412
Malignant lymphoma	5	3.7	0.4248	3	2.1	0.4786	2	1.5	0.6642
Kidney	4	2.8	0.372	2	1.3	0.0428	2	1.5	0.6643
Skin	4	1.7	0.093	4	0.9	0.5772	0	0.8	1
Pancreas	3	5.8	0.3958	1	2.9	0.7657	2	2.8	1
Thyroid	3	0.6	0.025	3	0.3	0.2849	0	0.3	1
Other	6			1			5		
Total	169			92			77		

Obs = observed; exp = expected.

**Table 3.**  
Observed and Expected Number of Multiple Primary Malignancies in Females (n = 79)

Site	Total No. of Malignancies			Multiple Primary Malignancies Preceding and Synchronous			Colorectal Carcinoma Preceding		
	Obs	Exp	P Value	Obs	Exp	P Value	Obs	Exp	P Value
Uterus	26	8.1	<0.001	19	6.3	0.3079	7	1.8	0.0026
Stomach	18	17.1	0.8053	7	10.9	0.4458	11	6.1	0.061
Breast	14	9.4	0.1342	12	6.2	0.5413	2	3.1	0.7761
Liver	4	3.1	0.5618	0	1.4	0.0589	4	1.6	0.0845
Biliary tract	3	3.0	0.7744	1	1.3	0.1521	2	1.6	0.6755
Oral cavity/pharynx	3	0.7	0.0398	1	0.4	0.0617	2	0.3	0.0456
Malignant lymphoma	3	1.4	0.161	1	0.7	0.1713	2	0.6	0.1287
Skin	3	0.9	0.0628	1	0.4	0.0619	2	0.5	0.0842
Thyroid	3	1.4	0.1638	2	0.8	0.0434	1	0.6	0.4602
Lung	2	4.6	0.3431	1	2.1	0.7298	1	2.4	0.7397
Other	11			9			2		
Total	90			54			36		

Obs = observed; exp = expected.

malignancies. On the other hand, subsequent malignancies developed in 36 of 516 patients (7 percent) who received adjuvant chemotherapy and in 85 of 788 patients (10.8 percent) who did not receive adjuvant chemotherapy ( $P = 0.0263$ ).

### Multiple Primary Malignancies

The most common site of multiple primary malignancy among males was the stomach, followed by the lung, prostate, larynx, liver, esophagus, and urinary bladder (Table 2). In detail, the most common site of multiple primary malignancy in male colon carcinoma patients was the stomach (45 percent, 40/88) followed by the lung (14 percent, 12/88), whereas the incidences of stomach (35 percent, 19/55) and lung (23.4 percent, 13/55) carcinoma differed in male rectal carcinoma patients. The most common site among females was the uterus, followed by the stomach, breast, and liver (Table 3).

The sites that showed a higher incidence of multiple primary malignancy than the expected value were the prostate, larynx, urinary bladder, oral cavity/pharynx, and thyroid among males, and the uterus and oral cavity/pharynx among females. In particular, sites showing a significantly higher rate of malignancy than the expected value after colorectal carcinoma surgery were the prostate among males, and the uterus and oral cavity/pharynx among females.

With regard to uterine carcinoma, 14 cases had cervical carcinoma (12 cases of squamous-cell carcinoma and 2 cases of adenocarcinoma), 10 cases had corpus carcinoma (9 cases of adenocarcinoma and 1

case of adenosquamous carcinoma), and the details were unknown in 2 cases. The mean age at the onset of carcinoma was 55.3 (range, 33–76) years in cervical carcinoma cases, and 59.4 (range, 35–82) years in corpus carcinoma cases.

### DISCUSSION

In Japan, the incidence of colorectal carcinoma has shown a tendency to increase in recent years, and as the treatment outcomes for each organ carcinoma have improved, it is not unusual to see patients with multiple malignancies involving the colorectum and other organs.<sup>1,2</sup> In the current series of Japanese colorectal carcinoma patients, multiple primary malignancy occurred in 18.7 percent of males, among whom the most common site was the stomach, and in 14.7 percent of females, among whom the most common site was the uterus. The organs in which multiple primary malignancies occurred in colorectal carcinoma patients at a higher incidence than the expected values were the prostate, larynx, urinary bladder, oral cavity/pharynx, and thyroid among males, and the uterus and oral cavity/pharynx among females, which were not necessarily correlated with the numbers of carcinoma occurrences. In previous studies, the reported incidence of other organ carcinomas among colorectal carcinoma patients in Japan ranged from 3 to 8.7 percent, but all of those studies used shorter follow-up periods than our study.<sup>2-5</sup> Our long-term follow-up demonstrated that 15 to 20 percent of Japanese colorectal carcinoma patients experience multiple primary malignancies.

In the results indicated above, there are some noteworthy observations. First, for gastric carcinoma, its incidence in Japan ranks high among both males and females, and also in our study, the occurrence of gastric carcinoma ranked high among both males and females,<sup>1,10,11</sup> but the number was almost the same as the expected value. Hence, it is conceivable that the high occurrence of gastric carcinoma in Japanese colorectal carcinoma patients is merely a reflection of the high incidence of gastric carcinoma in Japan. However, uterine carcinoma was the most common carcinoma among female patients, showing a significantly higher rate than the expected value. With regard to the occurrence of uterine carcinoma among Japanese females, a comparison between data from 1975 and data from 1998 shows that the age-standardized incidence of invasive cervical carcinoma decreased by approximately one-half from 13.4 to 7.2 per 100,000 females; conversely, the corpus carcinoma incidence increased from 1.4 to 4.5 per 100,000 females,<sup>12</sup> and in 1998, the ratio of invasive cervical carcinoma to corpus carcinoma was 1.6:1. In this study, these two carcinomas occurred to ratio of 14:10, and the inclusion of HNPCC cases did not result in a particularly high proportion of corpus carcinoma occurrence. The incidence of uterine carcinoma in Japanese females with colorectal carcinoma needs further investigation in the light of the increasing tendency of patients with corpus carcinoma.

An interesting point in this study is that, in male patients, the incidence of malignancies such as larynx, urinary bladder, and oral cavity/pharynx carcinomas, was significantly higher than the expected value. One of the background factors contributing to such a higher rate may be cigarette smoking. In Japan, adult males have a smoking rate of approximately 50 percent, compared with 20 to 30 percent in Western countries.<sup>13</sup> Recently it has been reported that smoking also is associated with colorectal polyp and colorectal carcinoma.<sup>14-16</sup> To determine whether colorectal carcinoma is a cigarette-associated malignancy and whether cigarette-associated malignancies are likely to occur frequently in colorectal carcinoma patients, it is necessary to conduct further study analyzing a large number of patients followed for a long period.

Regarding the occurrence of other organ carcinomas in colorectal carcinoma patients, the role of genetic factors that contribute to diseases, such as HNPCC, also should be investigated. The reported frequency of HNPCC accounts for up to 5 percent in

Western countries, whereas in Japan the frequency ranges from 0.15 to 0.2 percent, which is a greatly low incidence.<sup>17-20</sup> This low rate will possibly rise in future long-term investigations, because the surveillance of patients with HNPCC has just begun in Japan.<sup>19</sup> The incidence of multiple primary malignancy has been reported to be high in patients with HNPCC.<sup>18,21</sup> In this study, we found ten cases of corpus carcinoma; however, as described above, the inclusion of HNPCC cases did not result in a particularly high proportion of corpus carcinoma. Similarly, we also found carcinoma of the renal pelvis in only one case among males, and carcinoma of the small intestine and ureter in only one case among females. It has been pointed out that gastric carcinoma also may occur frequently in patients with HNPCC, but as indicated above, it could be speculated that the high occurrence of gastric carcinoma in Japanese colorectal carcinoma patients is merely the result of the high gastric carcinoma incidence in Japan.<sup>3,19</sup>

In this study, patients with a more advanced Dukes stage had a significantly lower proportion of multiple primary malignancies in both males and females (Table 1). This can be explained by patients with a more advanced Dukes stage having a lower survival rate, resulting in shorter term follow-up and a lower proportion of multiple primary malignancies. However, patients with multiple primary malignancies demonstrated that the mean age at the onset of colorectal carcinoma was significantly high in both males and females compared with that in patients with single colorectal carcinoma. Although patients with single colorectal carcinoma are candidates for subsequent multiple primary malignancies with colorectal carcinoma, the reason for this result needs further investigation. Regarding the proportion of multiple primary malignancies, the proportion rate was significantly higher among male colon carcinoma patients than male rectal carcinoma patients. One possible reason for this is that HNPCC patients were included in this study. Obviously, the rates of colon carcinoma and multiple primary malignancies are elevated in HNPCC patients; however, there was no difference among females, although HNPCC patients also were included. The reason for the difference in incidence between male colon and rectal carcinoma patients needs further investigation.

Obviously, this study has some limitations. With regard to the effects of cigarette smoking, the patients in our series were not interviewed in detail

about their smoking history and we, therefore, could not accurately investigate the contribution of smoking to carcinogenesis in patients with colorectal carcinoma or in patients who developed multiple primary malignancy. In addition, no investigation into carcinoma occurrence in family members was performed during the long-term follow-up period, and hence we also could not obtain data regarding situations in which HNPCC and other hereditary colorectal carcinomas occurred. Moreover, the influence of adjuvant treatment on subsequent malignancies needs further investigation. In this study, no patient developed subsequent malignancies after radiotherapy. One possible reason for this is that patients who received radiotherapy were in a relatively advanced stage, and that only 9 of 37 patients who received radiotherapy survived for more than five years from the surgery for colorectal carcinoma. These points need to be improved in further studies.

### CONCLUSIONS

This study was able to obtain some interesting findings on the incidence of other organ carcinomas in Japanese colorectal carcinoma patients. By conducting long-term follow-ups, we found that 15 to 20 percent of these patients in Japan develop multiple primary malignancies. This frequency is projected to rise in future studies with long-term follow-up. For colorectal carcinoma patients, postoperative long-term screening methods should be established, considering the actual occurrence numbers and the risk rate of multiple primary malignancies in addition to metachronous colorectal carcinoma.

### REFERENCES

- Ajiki W, Tsukuma H, Oshima A. Research Group for Population-based Cancer Registration in Japan. Cancer incidence and incidence rates in Japan in 1999: estimates based on data from 11 population-based cancer registries. *Jpn J Clin Oncol* 2004;34:352-6.
- Ueno M, Muto T, Oya M, Ota H, Azekura K, Yamaguchi T. Multiple primary cancer: an experience at the cancer institute hospital with special reference to colorectal cancer. *Int J Clin Oncol* 2003;8:162-7.
- Tomoda H, Taketomi A, Kohnoe S, Seo Y, Saito T. Second primary multiple primary cancers in Japanese hereditary nonpolyposis colorectal cancer. *Oncol Rep* 1998;5:143-5.
- Maruyama H, Hasuike Y, Furukawa J, *et al.* Multiple colorectal carcinomas and colorectal carcinoma associated with multiple primary malignancies. *Surg Today* 1992;22:99-104.
- Kobayashi Y, Arimoto H, Watanabe S. Occurrence of multiple primary cancer at the National Cancer Center Hospital, 1962-1989. *Jpn J Clin Oncol* 1991;21:233-51.
- Warren S, Gates O. Multiple primary malignant tumors: a survey of the literature and a statistical study. *Am J Cancer* 1932;16:1358-414.
- Research Group for Population-based Cancer Registration in Japan. Cancer incidence in Japan. In: Tajima K, Kuroishi T, Oshima A, eds. *Gann monograph on cancer research: cancer mortality and morbidity statistics*. Tokyo: Japan Scientific Societies Press, 2004:95-130.
- Anonymous. Cancer incidence in Japan, 1985-89: re-estimation based on data from eight population-based cancer registries. The Research Group for Population-based Cancer Registration in Japan. *Jpn J Clin Oncol* 1998;28:54-67.
- Anonymous. Cancer incidence and incidence rates in Japan in 1988: estimates based on data from ten population-based cancer registries. The Research Group for Population-based Cancer Registration in Japan. *Jpn J Clin Oncol* 1994;24:299-304.
- Tomoda H, Taketomi A, Baba H, Kohnoe S, Seo Y, Saito T. Multiple primary colorectal and gastric carcinoma in Japan. *Oncol Rep* 1998;5:147-9.
- Ikeda Y, Mori M, Kajiyama K, Haraguchi Y, Sugimachi K. Multiple primary gastric and colorectal cancer in Japan. *Int Surg* 1995;80:37-40.
- Ioka A, Tsukuma H, Ajiki W, Oshima A. Trends in uterine cancer incidence in Japan 1975-98. *Jpn J Clin Oncol* 2003;33:645-6.
- Forey B, Hamling J, Lee P, Wald N. *International smoking statistics*. London: Oxford, 2002.
- Colangelo LA, Gapstur SM, Gann PH, Dyer AR. Cigarette smoking and colorectal carcinoma mortality in a cohort with long-term follow-up. *Cancer* 2004;100:288-93.
- Slattery ML, Samowitz W, Ma K, *et al.* CYP1A1, cigarette smoking, and colon and rectal cancer. *Am J Epidemiol* 2004;160:842-52.
- Otani T, Iwasaki M, Yamamoto S, *et al.* Alcohol consumption, smoking, and subsequent risk of colorectal cancer in middle-aged and elderly Japanese men and females: Japan Public Health Center-based prospective study. *Cancer Epidemiol Biomarkers Prev* 2003;12:1492-500.
- Dunlop MG, Farrington SM, Carothers AD, *et al.* Cancer risk associated with germline DNA mismatch repair gene mutations. *Hum Mol Genet* 1997;6:105-10.

18. Aarnio M, Mecklin JP, Aaltonen LA, Nystrom-Lahti M, Jarvinen HL. Life-time risk of different cancers in hereditary non-polyposis colorectal cancer (HNPCC) syndrome. *Int J Cancer* 1995;64:430-3.
19. Kawakami K, Yasutomoi M, Baba S. Analysis of the HNPCC registries data reported at the 43rd Japanese Society for Cancer of the Colon and Rectum (JSCCR) meeting. In: Baba S, ed. *New strategies for treatment of hereditary colorectal cancer*. Tokyo: Churchill Livingstone, 1996:229-33.
20. Utsunomiya J, Miyaki M. Studies of hereditary non-polyposis colorectal cancer in Japan. *Int J Clin Oncol* 1998;3:53-74.
21. Watson P, Vasen HF, Mecklin JP, Jarvinen H, Lynch HT. The risk of endometrial cancer in hereditary non-polyposis colorectal cancer. *Am J Med* 1994;96:516-20.

## Second Hepatectomy for Recurrent Colorectal Liver Metastasis: Analysis of Preoperative Prognostic Factors

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**Background:** Second hepatectomy is a potentially curative treatment for patients with hepatic recurrence of colorectal cancer. However, there is still no consensus about the patient selection criteria for second hepatectomy under these circumstances, and the factors affecting prognosis after second hepatectomy remain uncertain.

**Methods:** Clinicopathologic data for 111 consecutive patients with colorectal liver metastasis who underwent second hepatectomy at a single institution between 1985 and 2004, and for whom complete clinicopathologic reports were available, were subjected to univariate and multivariate analyses.

**Results:** The morbidity and mortality rates were 14% and 0%, respectively, and the overall 5-year survival rate was 41%. Multivariate analysis revealed that synchronous resection for the first liver metastasis (hazard ratio, 1.8), more than three tumors at the second hepatectomy (1.9), and histopathological involvement of the hepatic vein and/or portal vein by the first liver metastasis (1.7) were independently associated with poor survival. We used these three risk factors to devise a preoperative model for predicting survival. The 5-year survival rates of patients without any risk factors, and with one, two, or three risk factors, were 62%, 38%, 19%, and 0%, respectively.

**Conclusions:** Second hepatectomy is beneficial for patients without any risk factors. Before second hepatectomy, chemotherapy should be considered for patients with any of these risk factors, especially with two or three factors, in the adjuvant or neoadjuvant setting to prolong survival. These results need to be confirmed and validated in another data set or future prospective trial according to the scoring scheme we outline.

**Key Words:** Second hepatectomy—Colorectal cancer—Liver metastasis—Prognostic factor—Neoadjuvant chemotherapy.

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Hepatectomy is the best and most potentially curative treatment for patients with colorectal liver metastases, yielding a 5-year survival rate of 38% to 51%.<sup>1–5</sup> After a first hepatectomy, 60% to 70% of

patients will develop recurrent disease, and one-third of these recurrences are limited to the liver.<sup>6</sup> The safety of hepatectomy has been increasing as a result of improvements in surgical techniques and perioperative management, and second liver resection has also been performed for patients with recurrent colorectal liver metastases. During the past decade, the reported outcomes of second hepatectomy have ranged from 21% to 49% in terms of 5-year survival after surgery. However, after second hepatectomy, some patients develop early recurrence in the liver,

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lung, and other organs, and in most of them, the disease is unresectable. Patients who experience recurrence within 6 months after second hepatectomy are already considered to have systemic disease before they undergo surgery. For such patients with systemic colorectal metastases, second hepatectomy is not beneficial and can even be harmful. However, the factors predicting early recurrence and poor outcome have not been established.

The purpose of this study was to find criteria that could be used to identify patients with recurrent liver metastases from colorectal cancer before surgery who would have a poor prognosis after second hepatectomy.

## MATERIALS AND METHODS

Between October 1985 and November 2004, data for 111 consecutive patients with recurrent liver metastases from colorectal cancer who underwent first and second hepatectomies at the National Cancer Center Hospital, Tokyo, were collected and reviewed. Patients who did not undergo initial liver resection at our hospital were excluded from the study because we were unable to obtain enough clinicopathologic data for them.

We investigated 27 clinicopathologic variables pertaining to patient characteristics, clinical data, and histopathologic findings, such as sex, age, primary cancer location, lymph node status, timing of first hepatectomy, number of hepatic metastases, tumor diameter, tumor distribution, preoperative serum carcinoembryonic antigen level, extent of liver resection, surgical margin, venous invasion by liver metastases, and bile duct invasion. The extent of liver resection was defined according to the nomenclature; wedge, segmental, and bisegmental resections were classified as minor resection, and hemihepatectomy or more extended resections were classified as major resection. Patient outcomes were determined on the basis of clinical data obtained from the files as of August 2005. The median follow-up period for the 111 patients after second liver resection was 43 months (range, 1–207 months).

The prognostic significance of clinicopathologic factors in relation to survival was investigated by univariate and multivariate analyses. Data were censored in the analysis of overall survival if a patient was alive, and in the analysis of disease-free survival if a patient was alive without recurrent colorectal cancer. Survival rates were calculated by the Kaplan-Meier method and compared statistically by the log-

rank test. Univariate comparisons of survival were performed by the log-rank test and multivariate analysis by the Cox regression model with the forward stepwise method (likelihood ratio). All variables were dichotomized for analysis. All statistical analyses were performed by SPSS for Windows, version 6.0 (SPSS-Japan Inc., Tokyo, Japan). All *P* values were two-sided, and differences at *P* < .05 were considered to be statistically significant.

## RESULTS

### Patient Characteristics and Follow-up

The 111 patients who underwent second hepatic resection with curative intent included 74 men and 37 women with a mean age of 59 years. The median interval between the first and second hepatic resections was 16 months (range, 4–96 months). At the last follow-up, 37 patients (34%) were alive with no evidence of recurrence, 12 (11%) were alive with disease, 61 (55%) had died of disease, and 1 patient was lost to follow-up. There were 23 actual 5-year survivors. The median follow-up time from primary resection was 69 months (range, 11–249 months), and the median follow-up from the second liver resection was 43 months (range, 1–207 months).

### Clinical Features and Pathology

#### Primary Tumor

The site of the primary cancer was the colon in 75 patients (68%) and the rectum in 36 (32%). Histologically, there were 50 well-differentiated, 58 moderately differentiated, and 1 poorly differentiated adenocarcinoma, and 2 mucinous carcinomas. Metastatic lesions in the liver were found synchronously with the primary tumor in 58 patients (stage IV). Of the 53 patients with metachronous liver metastases, 13 patients had no lymph node metastasis (stage I or II), and 40 patients had lymph node metastasis (stage III).

#### First Liver Resection

Of the 53 patients with metachronous liver metastases, the median interval between the primary resection and the first hepatectomy was 16 months (range, 4–60 months), and 25 patients (47%) underwent the second hepatectomy within 12 months. Unilobar involvement was observed in 68 patients and bilobar involvement in 43. At the first hepatectomy, 43 patients had a solitary hepatic lesion, 67 had

2 to 12 (median, 3) metastatic nodules, and one patient had more than 50 lesions. The median size of the largest hepatic lesion was 3.3 cm (range, 1.2–10 cm). Minor resection was performed in 93 patients and major resection in 18. The median blood loss was 698 mL (range, 50–3215 mL). Blood transfusion was performed in 17 patients. The surgical margin was negative in 89 patients and positive in 22. Invasions of the portal vein or hepatic vein by the liver metastases were found in 22 patients, and bile duct involvement was found in 39.

#### Second Liver Resection

The median interval between the first hepatectomy and detection of recurrence was 13 months (range, 2–92 months). Sixty-two patients had a solitary metastasis, and 49 had multiple metastases. The recurrent metastases ranged from 1.2 to 10 cm (median, 3.3 cm) in greatest dimension. Before second hepatectomy, pulmonary resection for lung metastasis was conducted in eight patients, and three patients underwent second hepatectomy and pulmonary resection simultaneously. After first hepatectomy, performed mostly in the 1980s, nine patients received adjuvant hepatic arterial infusion chemotherapy with 5-fluorouracil (5-FU), mitomycin C, and oral capecitabine regimen,<sup>7</sup> and six patients received oral anticancer drugs for adjuvant therapy (capecitabine in five patients, uracil-tegafur in one). Two patients who underwent colectomy plus simultaneous hepatectomy received adjuvant intravenous 5-FU plus leucovorin or mitomycin C, and one who had initially unresectable liver metastases was provided irinotecan for downstaging.

The second hepatectomy procedures included minor resection in 99 patients, hemihepatectomy in 6 patients, extended hemihepatectomy in 5, extended hemihepatectomy with bile duct reconstruction in 1, and central bisectionectomy in 1. Ninety-three patients had negative margins and 18 had positive margins. The median blood loss during the second liver resection was 913 mL (range, 95–4803 mL), and 22 patients received blood transfusions. No patient died during the perioperative course. Complications occurred in 16 patients (14%), including bile leakage in 8, abscess formation in 4, pleural effusion in 3, cholangitis in 1, and wound infection in 1. Invasions of the portal vein or hepatic vein were found in 25 patients, and bile duct involvement was found in 40.

#### Survival and Recurrence After Second Hepatectomy

Of the 111 patients who underwent second hepatectomy with curative intent, 61 had died by August 31, 2005. Overall 1-, 3-, and 5-year survival rates were

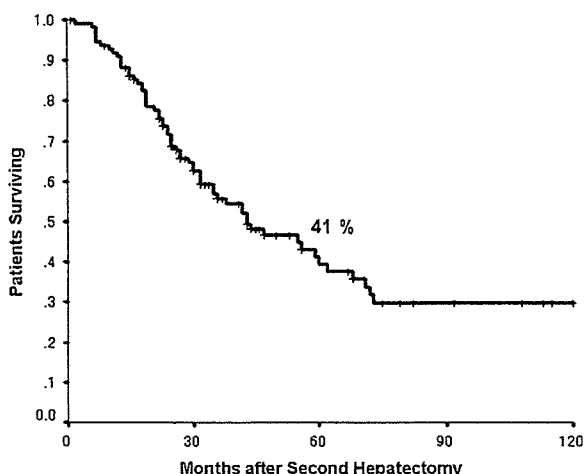


FIG. 1. Overall survival after second hepatectomy in patients with colorectal liver metastases.

91%, 74%, and 41%, respectively, from the time of second liver resection, with a median survival of 43 months (Fig. 1). There were 23 actual 5-year survivors. Recurrence after second hepatectomy occurred in 74 patients. Of these, 39 patients developed liver metastases (27 confined to the liver; 12 involving the liver plus other sites), and 37 developed lung metastases. Of them, 21 patients underwent surgery, including 13 third hepatectomies and 8 pulmonary resections. Twenty-seven patients experienced recurrence within 6 months after the second hepatectomy, and their median survival time was significantly worse than that of the others (15 vs. 60 months,  $P = .0001$ ). Forty-four of the patients who experienced recurrence after second hepatectomy received chemotherapy for treatment. Hepatic arterial infusion chemotherapy was performed in seven patients who had isolated hepatic recurrence. 5-FU was given by infusion to 14 patients; seven received additional mitomycin C. Oral chemotherapy drugs were administered to 7 patients (uracil-tegafur plus leucovorin in 2, S-1 [tegafur/5-chloro-2,4-dihydroxypyridine/potassium oxonate] in 3, capecitabine in 1, and capecitabine in 1), and intravenous 5-FU plus leucovorin was provided to 11. Twenty patients received irinotecan, eight received oxaliplatin, and one received bevacizumab. Irinotecan has been commonly used since 1999 in Japan, and most of the patients who experienced recurrence after 1999 benefited from irinotecan. The group that developed recurrence until 1998 ( $n = 19$ ) had significantly worse survival than those after 1999 ( $n = 25$ ) (median survival time, 23 months vs. 55 months,  $P = .004$ ).

### Univariate Analysis of Survival

#### Factors Related to Primary Lesion

The results of univariate analysis of survival are listed in Table 1. The presence of metastatic lymph nodes was a significant predictor of worse outcome after second liver resection ( $P = .03$ ). The location of the primary colorectal cancer and the histology of the primary lesion did not influence survival.

#### Factors Related to First Hepatectomy

Patients with metachronous liver metastases had a significantly better median survival time after second hepatectomy (60 vs. 32 months,  $P = .009$ ). The patients who received blood transfusions during the first hepatectomy had significantly worse survival ( $P = .049$ ). The number of tumors, size of the largest tumor, bilobar involvement, extent of hepatectomy, and serum carcinoembryonic antigen level were not prognostic factors. In terms of microscopic findings, invasions of the portal vein or hepatic vein, and a positive surgical margin tended to be associated with poor survival, but the difference was marginal ( $P = .07$  and  $.07$ , respectively). Bile duct invasion was not a prognostic factor.

#### Factors Related to Second Hepatectomy

The 5-year survival was significantly better for patients with a disease-free interval of more than 6 months between the first hepatectomy and recurrence, as compared with patients with a disease-free interval of less than 6 months (49% vs. 22%,  $P = .02$ ). Patients who had less than four nodules had significantly better survival than those with four or more (45% vs. 18%,  $P = .001$ ). The size of the largest lesion at the second operation influenced survival, but not to a significant degree ( $P = .09$ ). Similar to the first hepatectomy, patients who received blood transfusions showed significantly worse survival ( $P = .03$ ), and bilobar involvement, extent of hepatectomy, and serum carcinoembryonic antigen level were not prognostic factors. Patients who had undergone resection of extrahepatic disease before second hepatectomy did not show worse survival. With respect to the microscopic features of the recurrent metastatic disease, a surgical margin, invasions of the portal vein or hepatic vein, and bile duct invasion had no statistically significant influence on survival.

### Multivariate Analysis of Survival

Multivariate analysis identified three independent risk factors: synchronous first hepatectomy, four or

TABLE 1. Univariate analysis related to survival

Prognostic factor	No. of Patients	5-Year survival rate of second hepatectomy (%)	<i>P</i> value
<b>Demographics</b>			
<b>Age</b>			
< 60 y	56	43	.95
≥ 60 y	55	38	
<b>Sex</b>			
Male	74	38	.64
Female	27	46	
<b>Primary lesion</b>			
<b>Location</b>			
Colon	75	36	.17
Rectum	36	52	
<b>Lymph nodes</b>			
Negative	21	74	.03
Positive	90	33	
<b>First hepatectomy</b>			
<b>Number of lesions</b>			
< 4	90	43	.33
≥ 4	21	33	
<b>Size</b>			
< 5 cm	87	42	.45
≥ 5 cm	24	37	
<b>Timing with primary</b>			
Metachronous	53	51	.009
Synchronous	58	30	
<b>Distribution</b>			
Unilobar	68	44	.25
Bilobar	43	36	
<b>Resection</b>			
Minor	93	38	.92
Major	18	53	
<b>CEA before first hepatectomy</b>			
< 50 ng/dL	89	42	.31
≥ 50 ng/dL	22	37	
<b>Blood loss</b>			
< 1000 mL	91	35	.27
≥ 1000 mL	20	57	
<b>Blood transfusion</b>			
No	94	41	.049
Yes	17	35	
<b>Surgical margin</b>			
No	88	45	.07
Yes	23	29	
<b>Vessel invasion</b>			
No	89	46	.07
Yes	22	23	
<b>Bile duct invasion</b>			
No	72	36	.1
Yes	39	50	
<b>Second hepatectomy</b>			
<b>Interval between first hepatectomy and recurrence</b>			
< 6 mo	79	49	.02
≥ 6 mo	36	22	
<b>Extrahepatic disease before second hepatectomy</b>			
No	100	39	.31
Yes	11	54	
<b>Number of lesions</b>			
< 4	93	45	.001
≥ 4	18	18	
<b>Size</b>			
< 5 cm	100	44	.09
≥ 5 cm	11	16	

TABLE 1. Continued

Prognostic factor	No. of Patients	5-Year survival rate of second hepatectomy (%)	P value
Distribution			
Unilobar	68	45	.11
Bilobar	43	34	
Resection			
Minor	96	41	.67
Major	15	44	
CEA before second hepatectomy			
< 50 ng/dL	89	42	.31
> 50 ng/dL	22	37	
Blood loss			
< 1000 mL	73	40	.77
> 1000 mL	38	42	
Blood transfusion			
No	89	43	.03
Yes	22	29	
Surgical margin			
No	90	45	.16
Yes	21	22	
Vessel invasion			
No	75	36	.17
Yes	36	52	
Bile duct invasion			
No	71	43	.95
Yes	40	35	

TABLE 2. Multivariate analysis with Cox proportional hazard model

Prognostic factor	Relative risk	95% Confidence interval	P value
Synchronous timing of first hepatectomy	1.85	1.10-3.11	.02
Presence of vessel invasion at first hepatectomy	1.79	1.00-3.19	.049
Number of lesions at second hepatectomy $\geq 4$	1.94	1.10-3.41	.022

more lesions at second hepatectomy, and invasion of the portal vein or hepatic vein at first hepatectomy (Table 2). Any of the variables related to microscopic findings—information that could only be obtained after the second hepatectomy—were not statistically significant prognostic factors.

All three risk factors identified in the multivariate analysis were based on information obtained before the second hepatectomy. Therefore, we tried to group patients according to risk factors. Thirty-four patients had no risk factors, 53 had one factor, 22 had two factors, and 2 had three factors. Survival expectancies at 5 years for patients with no risk factors, one or two risk factors, and three risk factors were 62%, 31%, and 0%, respectively, and these differences were statistically significant ( $P = .001$ ) (Fig. 2). Two patients with three risk factors developed recurrence

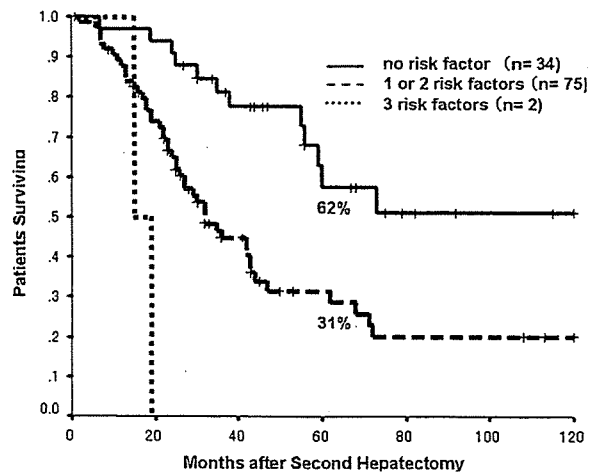


FIG. 2. Overall survival according to groups divided by number of risk factors identified in current study.

within 6 months and died at 15 months and 19 months, respectively, after the second hepatectomy.

DISCUSSION

The only potentially curative treatment for patients with liver-isolated colorectal metastases is surgical resection. However, most cannot be considered candidates for surgery for reasons such as very large tumor, unfavorable tumor location, multinodularity, or inadequate hepatic reserve. Nevertheless, because of its impact on survival, surgical resection is the treatment of choice when it is feasible. Improvements in surgical techniques and perioperative management have made surgical resection more feasible, with a mortality rate of less than 3%.<sup>2-4,8</sup> For patients with recurrent colorectal liver metastases, second hepatic resection has been performed more frequently during the past decade.<sup>6,9-22</sup> The results of previous series involving more than 20 patients are listed in Table 3. Five-year survival rates of 38% to 51% have been reported for first hepatic resections,<sup>1-5,23</sup> and similar 5-year survival rates, ranging from 21% to 49%, can be achieved after second hepatic resection for well-selected patients. In fact, morbidity and mortality after second hepatic resection are almost comparable to those after initial hepatic resection. The median survival and 5-year survival rate in the present series were 43 months and 41%, respectively. Because conventional chemotherapy alone cannot achieve such favorable results, a second hepatectomy has become the treatment of choice for recurrent liver metastases from colorectal cancer.

TABLE 3. Reports of second hepatectomy

Author	Year	No. Patients	Mortality (%)	Morbidity (%)	5-Year survival	MST (mo)
Fong et al. <sup>9</sup>	1994	25	0	28	NR	30
Nordlinger et al. <sup>10</sup>	1994	116	.9	25	33 <sup>a</sup>	NR
Que and Nagorney <sup>11</sup>	1994	21	5	NR	43 <sup>b</sup>	41
Fernandez-Trigo et al. <sup>12</sup>	1995	170	NR	19	32	34
Riesener et al. <sup>13</sup>	1996	25	0	20	24 <sup>a</sup>	NR
Adam et al. <sup>6</sup>	1997	64	0	20	26	46
Tuttle et al. <sup>14</sup>	1997	23	0	22	32	40
Yamamoto et al. <sup>15</sup>	1999	75	0	11	31	31
Muratore et al. <sup>16</sup>	2001	29	3.4	10	35 <sup>a</sup>	NR
Suzuki et al. <sup>17</sup>	2001	26	0	33	32	31
Petrowsky et al. <sup>18</sup>	2002	126	1.6	28	34	37
Takahashi et al. <sup>19</sup>	2003	22	0	18	49 <sup>a</sup>	23
Tanaka et al. <sup>20</sup>	2004	26	0	30	48 <sup>c</sup>	NR
Sugawara et al. <sup>21</sup>	2005	27	0	22	49	41
Pessaux et al. <sup>22</sup>	2006	42	0	14	21	25

MST, median survival time; NR, not reported.

<sup>a</sup> Three-year survival.

<sup>b</sup> Four-year survival.

<sup>c</sup> Five-year disease-free survival.

Recently, systemic chemotherapy has led to markedly improvements in median overall survival and progression-free survival.<sup>24-26</sup> These benefits are most pronounced with regimens containing irinotecan or oxaliplatin in combination with 5-FU plus leucovorin; median overall survival durations consistently approach 20 months, and some are as high as 24 months. Some patients experience early repeat recurrence within 6 months after second hepatectomy, and most of the disease is in an unresectable state when repeat recurrence is detected. In our series, 27 patients developed repeat recurrence within 6 months. Two patients underwent additional surgery, and the others received systemic chemotherapy, with a median survival time of 15 months. Considering these results, because their median survival was worse than that of patients who received systemic chemotherapy, the patients must have already had systemic disease at the time of surgery. Interestingly, we found that the patients who developed recurrence after 1999, when we started chemotherapy with irinotecan, had much better survival than those before 1998. This result seems mainly attributable to progress in chemotherapy, although diagnostic modality and perioperative management have been improved during these periods, and such improvements may have influenced survival. Now that we are in an era of effective chemotherapy, the indications for second liver resection need to be reconsidered. To improve the results of second hepatectomy, it is necessary to identify patients whose disease is likely to develop early repeat recurrence and who therefore should receive systemic chemotherapy.

Several studies have tried to identify factors predictive of a favorable outcome after repeat hepatectomy. To date, three reports have identified independent prognostic factors by multivariate analysis. Adam et al.<sup>6</sup> showed that the disease-free interval between initial and second liver resections and a second liver resection with curative intent were independently associated with survival. In our study, univariate analysis showed that a disease-free interval of more than 6 months between the first hepatectomy and recurrence was a significant prognostic factor, although it did not reach statistical significance by multivariate analysis. Our multivariate analysis showed that synchronous first hepatectomy was an independently predictive factor. Petrowsky et al.<sup>18</sup> showed that the presence of multiple lesions at repeat hepatectomy and a maximum tumor size exceeding 5 cm were independent prognostic factors after repeat hepatectomy. The third report, by Yamamoto et al.<sup>15</sup> from our hospital in 1999, involved data from 90 repeat hepatectomies (second = 75; third = 12; fourth = 3). Multivariate analysis revealed two independent prognostic factors after the second hepatectomy: four or more tumors, and the presence of extrahepatic disease. The present study detected three independent prognostic factors: synchronous first hepatectomy, four or more lesions evident at the second hepatectomy, and invasions of the portal vein or hepatic vein at the first hepatectomy. Thus, only the number of lesions was a common predictor of outcome, whereas the other factors differed from those highlighted in the first study. We speculate that the reason for this difference was patient selection. In