

Keywords: mitomycin-C; doxifluridine; cisplatin; adjuvant chemotherapy; gastric cancer; D2 gastrectomy

Adjuvant chemotherapy for gastric cancer: a randomised phase 3 trial of mitomycin-C plus either short-term doxifluridine or long-term doxifluridine plus cisplatin after curative D2 gastrectomy (AMC0201)

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Background: This phase 3 study evaluated the efficacy of new adjuvant chemotherapy (MFP), which intensified the mitomycin-C (MMC) plus short-term doxifluridine (Mf) for gastric cancer.

Patients and methods: A total of 855 patients (424 in Mf, 431 in MFP) with pathological stage II–IV (M0) gastric cancer after D2 gastrectomy were randomly assigned to receive either Mf (MMC 20 mg m⁻², followed by oral doxifluridine 460–600 mg m⁻² per day for 3 months) or MFP (MMC 20 mg m⁻², followed by oral doxifluridine 460–600 mg m⁻² per day for 12 months with 6 monthly infusions of 60 mg m⁻² of cisplatin) chemotherapy.

Results: With a median follow-up of 6.6 years, there was no difference between the two groups in recurrence-free survival (RFS) (5-year RFS 61.1% in Mf and 57.9% in MFP; hazard ratio 1.10 (95% CI 0.89–1.35); *P* = 0.39) and overall survival (OS) (5-year OS 66.5% in Mf and 65.0% in MFP; hazard ratio 1.11 (95% CI 0.89–1.39); *P* = 0.33).

Conclusion: Intensification of Mf adjuvant chemotherapy by prolonging the duration of oral fluoropyrimidine and adding cisplatin was safe but not effective to improve the survivals in curatively resected gastric cancer patients.

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Gastric cancer (GC) is the second most common cause of cancer-related mortality worldwide, with 989 600 new cases and 738 000 deaths per year (Jemal *et al*, 2011). For locally advanced cases, the only treatment option for potential cure is complete surgical resection. Unfortunately, a significant number of patients experience recurrence even after complete resection, and the prognosis of recurrent GC is dismal. The high recurrence rate and poor prognosis have led to extensive investigation of the use of adjuvant treatments to improve survival. However, there had been a debate on the role of adjuvant treatment until early 2000, especially in western countries, primarily due to the lack of large pivotal study, although several meta-analyses consistently suggesting small but significant benefits (Hermans *et al*, 1993; Earle and Maroun, 1999).

In 2000, when the AMC0201 trial was designed, adjuvant chemotherapy was popularly used in clinical practice in Korea. Because of this widespread practice, we believed that using surgery only in the control arm would not be appropriate. Thus, we selected adjuvant chemotherapy rather than surgery alone as the control arm and aimed to focus on reinforcing its efficacy in this trial. Mitomycin-C (MMC)-based regimens were one of the most popularly used adjuvant chemotherapies at that time (Chang *et al*, 2002; Koo *et al*, 2007). A Japanese meta-analysis suggested the efficacy of MMC-based adjuvant chemotherapy, and a Spanish randomised phase 3 study showed the survival benefit of simple combination of MMC and 3 months of oral fluoropyrimidine, although sample size of this study was small (Nakajima *et al*, 1994; Cirera *et al*, 1999). On the basis of these results, we decided to use MMC plus short-term oral fluoropyrimidine (Mf) as a reference treatment. To strengthen this regimen, we mapped out the following two strategies. We prolonged the duration of oral fluoropyrimidine, and also added cisplatin, an active component of effective regimens for metastatic GC (MFP).

We then performed this prospective, randomised phase 3 study to determine whether MFP could improve recurrence-free survival (RFS) over the Mf regimen in patients with GC after curative D2 resection. Here we report the final results of the AMC0201 study with inclusion of long-term outcome.

MATERIALS AND METHODS

Study design and patients. AMC0201 was an open-label, prospective randomised phase 3 clinical trial conducted at three centres in Korea.

The criteria for eligibility were histologically proven gastric adenocarcinoma; D2 lymph-node dissection; R0 resection; an age of 18–70 years; pathological stage II–IV without distant metastasis according to the 6th edition of American Joint Committee on Cancer (AJCC) staging system; an Eastern Cooperative Oncology Group (ECOG) performance status of 0–2; and adequate renal, liver and bone marrow function. Exclusion criteria included any of the following: incomplete surgical resection, prior chemotherapy, immunotherapy, or radiotherapy, concurrent or previous malignancy within the past 5 years and positive M1 lymph nodes on postoperative pathologic examination.

The protocol was approved by the institutional review board at each participating institution and conducted in accordance with the ethical principles of the Declaration of Helsinki and also the Good Clinical Practice guidelines, as defined by the International Conference on Harmonisation. All patients provided written informed consent before entry into the study.

Study treatment. At 3 weeks after surgery, eligible pathological stage II–IV (M0) cases were randomly assigned at a 1:1 ratio to receive the Mf or MFP regimen. Randomisation was stratified according to stage and each participating centre based on a

permutation block method. Patients and investigators were not masked to study treatment.

For patients randomised to the Mf group, 20 mg m⁻² MMC was given intravenously 3–6 weeks after surgery, and then 4 weeks later, daily oral doses of 460 mg m⁻² doxifluridine were administered for 3 months. In the MFP group, 20 mg m⁻² MMC was given intravenously 3–6 weeks after surgery, and beginning 4 weeks later, 60 mg/m⁻² cisplatin was given intravenously monthly for 6 months and 460 mg m⁻² per day oral doxifluridine was administered for 12 months. One cycle lasted for 1 month. The doxifluridine dose was increased to 600 mg m⁻² per day in both groups after interim safety analysis in February 2004. Dose modification was prespecified for patients who experienced haematologic and non-haematologic toxicities (Supplementary Data).

Assessments. At baseline, patients underwent a history, physical examination, including weight, height and vital signs. Computed tomography (CT) scans of the abdomen and pelvis with contrast enhancement, chest radiography and electrocardiography were performed. In addition, laboratory tests, including complete blood count with differential counts, electrolytes, coagulation test, liver and renal function testes were obtained. Adverse events and laboratory profiles were assessed every 4 weeks during treatment. After completion of chemotherapy, clinical assessments were performed every 3 months up to 2 years from surgery, every 6 months between 2 and 5 years from surgery and every 1 year thereafter. Plain chest radiography and abdominopelvic contrast-enhanced CT scans were performed every 6 months within 5 years from surgery and every 1 year thereafter; gastroscopy was performed every 1 year. If there was any sign or symptom indicating recurrence, investigations were immediately carried out to verify the status of the patients. If findings on imaging studies were suggestive but not conclusive, rigorous serial follow-up studies were performed to detect recurrence. The date of the first recognition of findings suggestive of recurrence was defined as the date of recurrence. Adverse events were graded according to the National Cancer Institute's Common Terminology Criteria for Adverse Events, version 2.0.

Statistical analysis. The primary end point was 3-year RFS, and the secondary end points were 3-year overall survival (OS), disease recurrence and safety. All randomised patients were analysed on an intention-to-treat basis. On the basis of the result of previous trials, the 3-year RFS rate in the Mf group was estimated to be 60%. This trial was initially designed to identify a 15% improvement of 3-year RFS rate in the MFP group and a total of 475 patients were required for a two-sided α of 5% and a statistical power of 90%, considering 10% of follow-up loss.

In February 2004, when we had enrolled half of the planned number of patients (245), we performed a planned interim analysis for safety monitoring and potential sample size recalculation based on the distribution of pathological stage in study population. After the study began, several participating investigators concerned that the original dose of doxifluridine was low. As the toxicity of original dose was mild and well tolerated at interim analysis, we decided to increase the doxifluridine dose from 460 to 600 mg m⁻² per day by protocol amendment. When we analysed the distribution of pathological stage until that point, patients with early stage were included more than we originally expected. On the basis of this, the 3-year RFS of control arm was re-estimated as better than our original assumption. This estimation and rapid patient accrual enabled us to increase the sample size to detect accurately real, even if small, improvements in efficacy. Therefore, the estimated 3-year RFS rate in the Mf group was increased to 70%, while the expected additional benefit in the MFP group was lowered to 10% (80% of 3-year RFS rate). With the same type I and II error rates, we estimated that a total enrolment of 881 patients

(207 events) would be necessary for a hazard ratio (HR) of 0.6256 in the MFP group as compared with the Mf group.

Recurrence-free survival was defined as the time from randomisation to documented disease recurrence or death, and OS as the time from randomisation to death from any cause. Patients were censored if they were recurrence-free or alive at the last follow-up. Kaplan–Meier curves were used to estimate survival, and compared using the log-rank test. The Cox regression model was used to estimate HRs based on the comparison of the efficacy between the Mf and MFP arms in both primary analysis and subgroup analyses. All tests were two-sided, and a *P*-value < 0.05 was considered to indicate statistical significance. Confidence intervals (CI) are at the 95% level. Descriptive analyses were used to represent adverse events. The relative dose intensity (RDI) was calculated as the percentage of actually administered dose to planned dose per unit time. Statistical analyses were performed by a qualified biostatistician (BKP) using SAS version 9.1 (SAS Institute Inc., Cary, NC, USA) and SPSS version 14.0 (SPSS Inc., Chicago, IL, USA).

After 229 events occurred with a median follow-up period of 3.5 years, the database was initially locked on 20 March 2008 and planned primary analysis was performed. These results were previously presented at ASCO annual meeting, 2008 (Chang *et al*, 2008). Afterwards, to verify the long-term results of this study, 3-year extension analysis was performed. The data cutoff for this extension analysis was 11 April 2011. Median follow-up at that time was 6.6 years (maximum 109.4 months).

RESULTS

Patient characteristics. Between February 2002 and August 2006, 855 patients were randomly allocated to study arms (424 in MF and 431 in MFP; Figure 1). Baseline characteristics turned out to be well balanced between the two groups (Table 1). Median time interval between surgery and chemotherapy was 23 days in the Mf group and 24 days in the MFP group. Postoperative stages were II in 51%, IIIA in 31%, IIIB in 9% and IV in 9% of patients. All patients received D2 surgery with R0 resection.

Treatment delivery. Planned treatment was completed in 93% of the Mf group and 72% of the MFP group. Throughout the treatment period, 145 patients (34%) underwent dose reduction for doxifluridine in the Mf group compared with 322 (75%) and 19 (4%) for doxifluridine and cisplatin in the MFP group. The reasons for dose reduction were haematologic toxicities (97% in the Mf group; 97% for doxifluridine and 42% for cisplatin in the MFP group), or non-haematologic toxicities (3% in the Mf group; 3% for doxifluridine and 58% for cisplatin in the MFP group). Dose delay was applied in 64 patients (15%) in the Mf group and 189 patients (44%) in the MFP group, mainly because of haematologic toxicities (83% in the Mf group and 86% in the MFP group) of all adverse events. Median RDI per cycle for doxifluridine was 96.7–100.0% in the Mf group and 79.0–100.0% in the MFP group. The median RDI per cycle for cisplatin was 100.0% throughout all cycles.

Safety. Adverse events that occurred in 10% or more of patients are summarised in Table 2. The population evaluated for safety comprised 847 patients (422 in the Mf group and 425 in the MFP group), with the exclusion of the patients who did not receive allocated treatment after randomisation or were lost to follow-up after MMC administration. Both regimens were well tolerated, and the adverse events of all grades were more frequent in patients who received MFP than in those who received Mf. Grade 3 or 4 neutropenia was more common in the MFP group (35%) than in the Mf group (11%). Two patients in the Mf group and four patients in the MFP group developed febrile neutropenia. Grade 3 or 4 thrombocytopenia was rare and similar in both arms (3% in both groups). Grade 3 or 4 non-haematologic adverse events were uncommon in both groups; however, patients in the MFP group had slightly higher rates of fatigue, anorexia, nausea, vomiting and diarrhoea. There was no treatment-related death. Between patients enrolled before and after the introduction of protocol amendment in each Mf and MFP groups, there were no significant differences in the frequency of grade 3 or 4 haematologic or non-haematologic toxicities, except that grade 3 or 4 neutropenia in the MFP group was significantly increased with higher dose of doxifluridine (22% vs 14%; *P* = 0.003). Actually administered dose of doxifluridine during six cycles of study treatments was increased after the protocol amendment than before (mean 517.89 vs 409.24 mg m⁻²

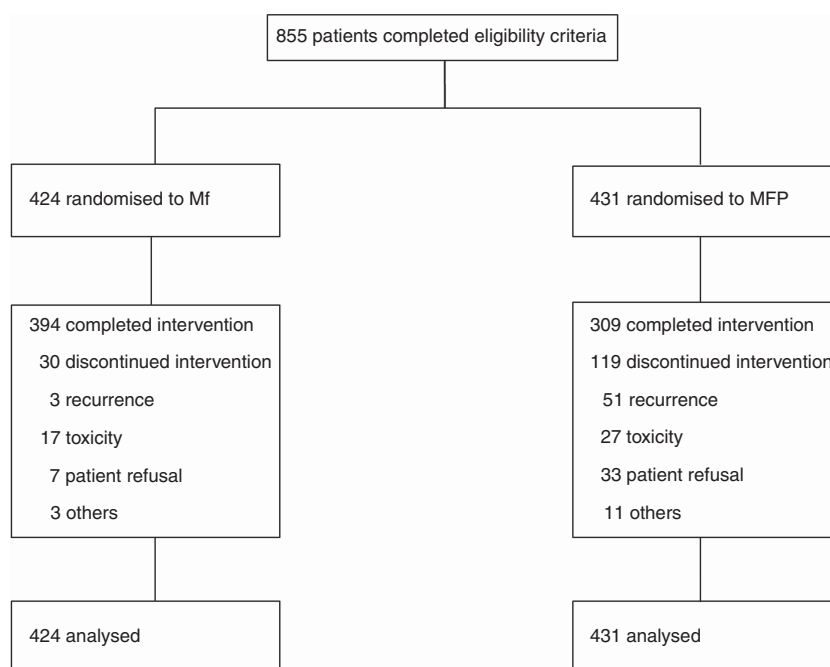


Figure 1. Trial profile. Mf, mitomycin-C plus short-term doxifluridine; MFP, mitomycin-C plus long-term doxifluridine plus cisplatin.

Table 1. Baseline patient characteristics

Characteristics	Mf arm (n = 424)	MFP arm (n = 431)
Age (years)	56 (29–70)	55 (20–70)
Male gender	294 (69%)	294 (68%)
ECOG performance status		
0–1	423 (> 99%)	428 (> 99%)
2	1 (< 1%)	3 (< 1%)
Primary site		
Proximal	40 (10%)	33 (8%)
Distal	342 (81%)	351 (81%)
Multiple/diffuse	42 (10%)	47 (11%)
Type of surgery		
Total gastrectomy	162 (38%)	166 (39%)
Subtotal gastrectomy	262 (62%)	265 (61%)
Lauren's classification		
Intestinal type	148 (35%)	135 (31%)
Diffuse type	222 (52%)	235 (55%)
Mixed type	39 (9%)	49 (11%)
Unknown	15 (4%)	12 (3%)
Pathologic T stage		
pT1	9 (2%)	8 (2%)
pT2	261 (62%)	235 (55%)
pT3	147 (35%)	178 (41%)
pT4	7 (2%)	10 (2%)
No. of lymph-node metastasis		
0	28 (7%)	43 (10%)
1–6	256 (60%)	249 (58%)
7–15	109 (26%)	109 (25%)
≥ 16	31 (7%)	30 (7%)
Overall stage^a		
II	220 (52%)	216 (50%)
IIIA	130 (31%)	136 (32%)
IIIB	38 (9%)	42 (10%)
IV (M0)	36 (9%)	37 (9%)
Dose of doxifluridine		
460 mg m ⁻² per day	238 (56%)	233 (54%)
600 mg m ⁻² per day	186 (44%)	198 (46%)

Abbreviations: Mf = mitomycin-C plus short-term doxifluridine; MFP = mitomycin-C plus long-term doxifluridine plus cisplatin; ECOG = Eastern Cooperative Oncology Group.
^aAmerican Joint Committee on Cancer Staging System, 6th edition (2002).

per day, $P < 0.001$), although there was no difference in cisplatin (mean 56.64 vs 56.86 mg m⁻² per month, $P = 0.67$).

Long-term efficacy outcomes. With a median follow-up of 6.6 years in April 2011, a total of 353 events (recurrence or death) have been observed (167 in the Mf group and 186 in the MFP group). The 3- and 5-year RFS rates were 67.0% (95% CI, 62.5–71.5) and 61.1% (95% CI, 56.4–65.8) in the Mf group, respectively, and 64.9% (95% CI, 60.4–69.4) and 57.9% (95% CI, 53.2–62.6) in the MFP group, respectively. There was no statistically significant difference in RFS between the two groups (HR 1.10, 95% CI 0.89–1.35; $P = 0.39$; Figure 2A). The 3- and 5-year OS rates were 76.9% (95% CI, 72.9–80.9) and 66.5% (95% CI, 62.0–71.0) in the Mf group,

Table 2. Adverse events reported by ≥ 10% of patients (safety population)^a

	Toxicity grade by treatment arm			
	Mf arm (n = 422)		MFP arm (n = 425)	
	All grade	Grade 3 or 4	All grades	Grade 3 or 4
Haematologic				
Leucopenia	265 (63%)	20 (5%)	367 (86%)	24 (6%)
Neutropenia	262 (62%)	47 (11%)	387 (91%)	150 (35%)
Anaemia	393 (93%)	9 (2%)	413 (97%)	16 (4%)
Thrombocytopenia	101 (24%)	14 (3%)	156 (37%)	14 (3%)
Non-haematologic				
AST or ALT	119 (29%)	3 (< 1%)	145 (34%)	3 (< 1%)
Hyperbilirubinaemia	114 (27%)	1 (< 1%)	162 (38%)	4 (< 1%)
Fatigue	343 (81%)	0 (0%)	391 (92%)	9 (2%)
Anorexia	326 (77%)	1 (< 1%)	389 (92%)	10 (2%)
Nausea	293 (69%)	0 (0%)	371 (87%)	9 (2%)
Vomiting	104 (25%)	7 (2%)	160 (38%)	15 (4%)
Stomatitis	77 (18%)	0 (0%)	151 (36%)	0 (0%)
Constipation	132 (31%)	1 (< 1%)	204 (48%)	2 (< 1%)
Diarrhoea	219 (52%)	1 (< 1%)	288 (68%)	8 (2%)
Alopecia	261 (62%)	0 (0%)	334 (79%)	0 (0%)
Neuropathy	149 (35%)	0 (0%)	301 (71%)	2 (< 1%)
HFS	35 (8%)	0 (0%)	73 (17%)	0 (0%)
Myalgia	109 (26%)	0 (0%)	166 (39%)	1 (< 1%)
Oedema	44 (10%)	0 (0%)	108 (25%)	0 (0%)

Abbreviations: Mf = mitomycin-C plus short-term doxifluridine; MFP = mitomycin-C plus long-term doxifluridine plus cisplatin; AST = aspartate aminotransferase; ALT = alanine aminotransferase; HFS = hand-foot syndrome.
^aPatients who did not receive allocated treatment after randomisation or were lost to follow-up after MMC administration were excluded.

respectively, and 73.1% (95% CI, 68.9–77.3) and 65.0% (95% CI, 60.5–69.5) in the MFP group, respectively. The difference in OS was also insignificant (HR 1.11, 95% CI 0.89–1.39; $P = 0.33$; Figure 2B). Median RFS and OS have not been reached in either group. In subgroup analyses for RFS (Figure 3), there was no significant interaction between study treatment and baseline characteristics. RFS and OS were not affected by whether patients were enrolled before ($n = 471$) or after the protocol amendment ($n = 384$; $P = 0.96$ and 0.71 , respectively). The sites of first recurrence are summarised in Table 3. Distant metastasis was more common than locoregional recurrence in both groups, and there were no significant differences between the two groups.

DISCUSSION

After a long debate, it is now globally agreed that adjuvant treatment improves survival of patients with GC who undergo curative surgery. However, there are still geographical differences in standard adjuvant treatment modalities. The intergroup-0116 (INT-0116) study of postoperative chemoradiation (CRT) (Macdonald *et al*, 2001), and the Medical Research Council Adjuvant Gastric Infusional Chemotherapy (MAGIC) trial of perioperative chemotherapy (Cunningham *et al*, 2006), demonstrated the survival benefits in AGC over surgery alone, and it has become standard adjuvant therapy in the United States and

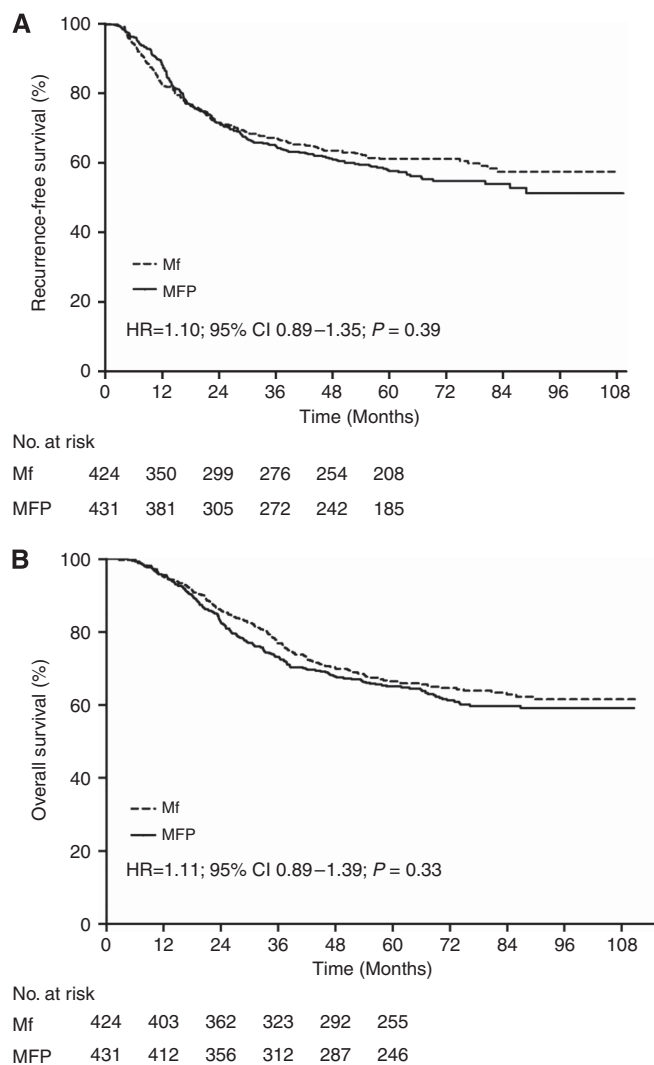


Figure 2. Kaplan–Meier estimates of RFS (A) and OS (B). Mf, mitomycin-C plus short-term doxifluridine; MFP, mitomycin-C plus long-term doxifluridine plus cisplatin.

Europe, respectively. In Asia, where D2 surgery is a standard procedure, postoperative chemotherapy has become standard therapy on the basis of the results of the ACTS-GC (S-1 for 12 months) (Sakuramoto *et al*, 2007) and the recent CLASSIC trial (capecitabine and oxaliplatin for 6 months) (Bang *et al*, 2012). Therefore, the most relevant issue in adjuvant treatment now is how to improve the treatment outcome of current standard adjuvant treatment. In this context, the results of this study offer timely suggestions regarding the question.

The control treatment of this study, Mf regimen is a reasonable adjuvant chemotherapy for which benefit over no treatment was confirmed by the recent GASTRIC meta-analysis study (GASTRIC Group, 2010). This study was reasonably powered to assess the clinically relevant improvement in RFS, and follow-up period was sufficient to confirm the long-term outcomes. Nevertheless, the RFS and OS curves of the two arms overlapped and there was no trend of benefit in the MFP arm. This strongly indicates that simple intensification of the adjuvant chemotherapy with Mf regimen by the addition of cisplatin and prolongation of doxifluridine administration is not effective. We do not know why this strategy is not working in the adjuvant setting, while it is working in the palliative setting. A meta-analysis showed that combination chemotherapy is better than single agent, mainly 5-

FU, and the Japanese SPIRITS trial demonstrated that addition of cisplatin to fluoropyrimidine could improve the survivals in patients with metastatic or recurrent GC (Wagner *et al*, 2006; Koizumi *et al*, 2008).

Many of the adjuvant chemotherapy trials testing polychemotherapeutic agents with cisplatin failed to achieve survival benefit over reference treatment (Chipponi *et al*, 2004; Bouche *et al*, 2005; Cascinu *et al*, 2007; Di Costanzo *et al*, 2008). These failures may result from low statistical power of the studies due to their small size, but low completion rates owing to toxicity also played an important role in poor performance. In our study, we gave 60 mg m⁻² of cisplatin to obtain a commonly prescribed dose of cisplatin in East Asia for lower toxicities and better compliance. This strategy resulted in better tolerability, but additive survival benefit was not obtained.

This result is in line with the recent results of CALGB 80101 and ITACA-S trials (Fuchs *et al*, 2011; Bajetta *et al*, 2012). In CALGB 80101 study, adjuvant CRT for gastric or gastro-oesophageal junction adenocarcinoma using epirubicin, cisplatin and 5-FU (ECF) was compared with bolus 5-FU/LV before and after 5-FU/radiotherapy, but the ECF arm did not achieve better disease-free survival (HR, 1.00, 95% CI, 0.79–1.27; P = 0.99) and OS (HR, 1.03, 95% CI, 0.80–1.34; P = 0.80) (Fuchs *et al*, 2011). Furthermore, ITACA-S trial failed to demonstrate the enhanced efficacy of intensified chemotherapy (four cycles of 5-FU, LV and irinotecan, followed by three cycles of docetaxel and cisplatin) compared with control chemotherapy (nine cycles of 5-FU and LV) in terms of disease-free survival (HR, 0.98, 95% CI, 0.83–1.16; P = 0.83) and OS (HR, 1.0, 95% CI, 0.83–1.20; P = 0.97) (Bajetta *et al*, 2012).

In AMC0101 study (Kang *et al*, 2008), a companion adjuvant chemotherapy trial of AMC0201, two more strategies, intraperitoneal chemotherapy and early start of chemotherapy, were applied in GC patients with macroscopically recognisable serosa invasion compared with the AMC0201 study. Moreover, addition of these four strategies to Mf chemotherapy resulted in a significantly improved RFS and OS, while addition of two strategies, prolonged doxifluridine and addition of cisplatin were not effective in AMC0201. It therefore appears that it is not the simple intensification by additional chemotherapeutic agents or prolongation of duration of chemotherapy, but rather the new strategies of early initiation of systemic chemotherapy and/or intraperitoneal chemotherapy that can be effective strategies to further improve the current standard adjuvant therapy. We believe these strategies should be tested in future clinical trials. Neoadjuvant chemotherapy, which is the earliest adjuvant chemotherapy we can have, is currently investigated in PRODIGY (NCT01515748) and JCOG0501 (C000000279) trials. These should provide findings concerning the efficacy of neoadjuvant chemotherapy in patients who will receive D2 gastrectomy and postoperative S-1.

As mentioned above, geographic differences exist in standard adjuvant treatments, in terms of therapeutic modality, timing of treatment and chemotherapy regimens. These discrepancies may be influenced by global differences in standard surgical methods or study populations. In Asia where D2 gastrectomy is the standard surgery, postoperative oral fluoropyrimidine-based adjuvant chemotherapy, such as S-1 for 1 year and capecitabine plus oxaliplatin for 6 months, is a proved treatment option for localised GC. However, because there has been lack of randomised trial comparing these regimens, it is difficult to determine which one is superior to others.

During the study, study protocol was amended in regards to sample size and dose of doxifluridine based on the results of interim analysis. The interim analysis was originally planned for safety monitoring and potential sample size recalculation based on the distribution of pathological stage in study population. Since this study started, relatively low dose of doxifluridine had been

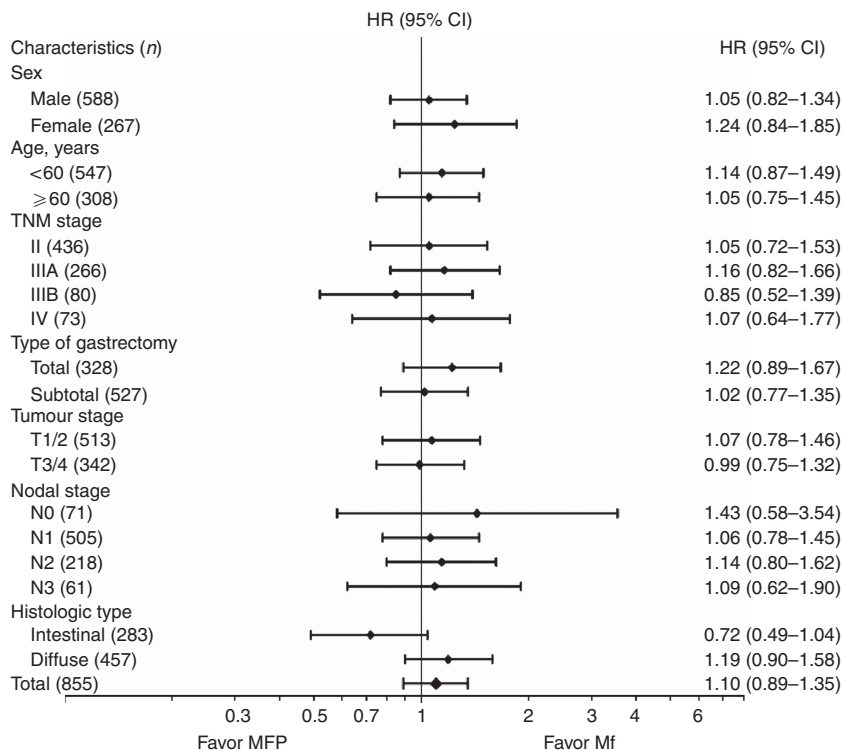


Figure 3. Subgroup analysis for RFS. Mf, mitomycin-C plus short-term doxifluridine; MFP, mitomycin-C plus long-term doxifluridine plus cisplatin; HR, hazard ratio; CI, confidence interval.

Table 3. Site of first recurrence^a

Site of recurrence	Mf arm (n = 424)	MFP arm (n = 431)	P-value
Total no. of recurrence	158 (37%)	170 (39%)	
Locoregional	28 (7%)	39 (9%)	0.20
Distant	130 (31%)	131 (30%)	0.94
Haematogenous	61 (14%)	75 (17%)	0.26
Peritoneal	65 (15%)	55 (13%)	0.32
Lymphatic	46 (11%)	44 (10%)	0.82

Abbreviations: Mf = mitomycin-C plus short-term doxifluridine; MFP = mitomycin-C plus long-term doxifluridine plus cisplatin.
^aMore than one recurrent site in some patients.

indicated by several participating investigators. Because the safety monitoring at interim analysis showed mild and tolerable toxicity profile of original dose, we decided to increase the doxifluridine dose from 460 to 600 mg m⁻² per day. Although actual administration of doxifluridine was increased without significant aggravation of toxicity except neutropenia, there was no difference in survival outcomes between patients enrolled before and after the introduction of a protocol amendment to increase daily dose of doxifluridine. The assumption of RFS in control arm was based on our previous results with Mf chemotherapy. However, the RFS of the control arm should change according to the stage distribution of the study population since stage is the most important prognostic factor. We wanted to check if the stage distribution of the study population were as we had assumed before. And, if there were differences in the interim analysis, we wanted to reassess the RFS of the control arm based on this stage distribution of the study population, and subsequently the sample size required to detect the

efficacy of the experimental arm. Actually, because the greater number of patients with early stage was enrolled than we originally expected, we had to re-estimate the expected 3-year RFS of control arm and recalculate the sample size in the planned interim analysis. Although this required increasing the sample size, rapid patient accrual enabled us to detect even small differences in efficacy between treatment groups.

In conclusion, prolongation of doxifluridine administration and addition of cisplatin to adjuvant chemotherapy with MMC plus 3 months of doxifluridine could be safely performed but did not improve the treatment outcome in curatively resected GC patients.

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REFERENCES

Bajetta E, Floriani I, Di Bartolomeo M, Labianca R, Landi L, Santoro A, Casaretti R, Pasquini E, Di Fabio F, Rondini E, Pinotti G, Bidoli P, Rosati G, Mambriani A, Ciarlo A, Cordio SS, Ricci S, Frassinetti L, Di Costanzo F, Bochicchio AM (2012) Intergroup Trial of Adjuvant Chemotherapy in Adenocarcinoma of the Stomach (ITACA-S) trial: Comparison of a sequential treatment with irinotecan (CPT-11) plus 5-fluorouracil (5-FU)/folinic acid (LV) followed by docetaxel and cisplatin

- versus a 5-FU/LV regimen as postoperative treatment for radically resected gastric cancer. *ASCO Meet Abstr* **30**: LBA4001.
- Bang YJ, Kim YW, Yang HK, Chung HC, Park YK, Lee KH, Lee KW, Kim YH, Noh SI, Cho JY, Mok YJ, Ji J, Yeh TS, Button P, Sirzen F, Noh SH (2012) Adjuvant capecitabine and oxaliplatin for gastric cancer after D2 gastrectomy (CLASSIC): a phase 3 open-label, randomised controlled trial. *Lancet* **379**(9813): 315–321.
- Bouche O, Ychou M, Burtin P, Bedenne L, Ducreux M, Lebreton G, Baulieux J, Nordlinger B, Martin C, Seitz JF, Tigaud JM, Echinard E, Stremstoerfer N, Milan C, Rougier P (2005) Adjuvant chemotherapy with 5-fluorouracil and cisplatin compared with surgery alone for gastric cancer: 7-year results of the FFCD randomized phase III trial (8801). *Ann Oncol* **16**: 1488–1497.
- Cascinu S, Labianca R, Barone C, Santoro A, Carnaghi C, Cassano A, Beretta GD, Catalano V, Bertetto O, Barni S, Frontini L, Aitini E, Rota S, Torri V, Floriani I, Pozzo C, Rimassa L, Mosconi S, Giordani P, Ardizzoia A, Foa P, Rabbi C, Chiara S, Gasparini G, Nardi M, Mansutti M, Arnoldi E, Piazza E, Cortesi E, Pucci F, Silva RR, Sobrero A, Ravaioli A (2007) Adjuvant treatment of high-risk, radically resected gastric cancer patients with 5-fluorouracil, leucovorin, cisplatin, and epidoxorubicin in a randomized controlled trial. *J Natl Cancer Inst* **99**: 601–607.
- Chang HM, Jung KH, Kim TY, Kim WS, Yang HK, Lee KU, Choe KJ, Heo DS, Bang YJ, Kim NK (2002) A phase III randomized trial of 5-fluorouracil, doxorubicin, and mitomycin C versus 5-fluorouracil and mitomycin C versus 5-fluorouracil alone in curatively resected gastric cancer. *Ann Oncol* **13**: 1779–1785.
- Chang HM, Kang YK, Min YJ, Zang DY, Kim GY, Yang DH, Jang SJ, Yook JH, Oh ST, Kim BS (2008) A randomized phase III trial comparing mitomycin-C plus short-term doxifluridine (Mf) versus mitomycin-C plus long-term doxifluridine plus cisplatin (MFP) after curative resection of advanced gastric cancer (AMC 0201) (NCT00296335). *J Clin Oncol* **26**(Suppl):abstr 4531.
- Chipponi J, Huguier M, Pezet D, Basso N, Hay JM, Quandalle P, Jaeck D, Fagniez PL, Gainant A (2004) Randomized trial of adjuvant chemotherapy after curative resection for gastric cancer. *Am J Surg* **187**: 440–445.
- Cirera L, Balil A, Batiste-Alentorn E, Tusquets I, Cardona T, Arcusa A, Jolis L, Saigi E, Guasch I, Badia A, Boleda M (1999) Randomized clinical trial of adjuvant mitomycin plus tegafur in patients with resected stage III gastric cancer. *J Clin Oncol* **17**: 3810–3815.
- Cunningham D, Allum WH, Stenning SP, Thompson JN, Van de Velde CJ, Nicolson M, Scarffe JH, Lofts FJ, Falk SJ, Iveson TJ, Smith DB, Langley RE, Verma M, Weeden S, Chua YJ. MAGIC Trial Participants (2006) Perioperative chemotherapy versus surgery alone for resectable gastroesophageal cancer. *N Engl J Med* **355**: 11–20.
- Di Costanzo F, Gasperoni S, Manzione L, Bisagni G, Labianca R, Bravi S, Cortesi E, Carlini P, Bracci R, Tomao S, Messerini L, Arcangeli A, Torri V, Bilancia D, Floriani I, Tonato M, Dinota A, Strafiuso G, Corgna E, Porrozzis S, Boni C, Rondini E, Giunta A, Monzio Compagnoni B, Biagioni F, Cesari M, Fornarini G, Nelli F, Carboni M, Cognetti F, Enzo MR, Piga A, Romiti A, Olivetti A, Masoni L, De Stefanis M, Dalla Mola A, Camera S, Recchia F, De Filippis S, Scipioni L, Zironi S, Luppi G, Italia M, Banducci S, Pisani Leretti A, Massidda B, Ionta MT, Nicolosi A, Canaletti R, Biscottini B, Grignani F, Rovei R, Croce E, Carroccio R, Gilli G, Cavalli C, Olgiati A, Pandolfi U, Rossetti R, Natalini G, Foa P, Oldani S, Bruno L, Cascinu S, Catalano G, Catalano V, Lungarotti F, Farris A, Sarobba MG, Trignano M, Muscogiuri A, Francavilla F, Figoli F, Leoni M, Papiani G, Orselli G, Antimi M, Bellini V, Cabassi A, Contu A, Pazzola A, Frignano M, Lastraioli E, Saggese M, Bianchini D, Antonuzzo L, Mela M, Camisa R (2008) Adjuvant chemotherapy in completely resected gastric cancer: a randomized phase III trial conducted by GOIRC. *J Natl Cancer Inst* **100**: 388–398.
- Earle CC, Maroun JA (1999) Adjuvant chemotherapy after curative resection for gastric cancer in non-Asian patients: revisiting a meta-analysis of randomised trials. *Eur J Cancer* **35**: 1059–1064.
- Fuchs CS, Tepper JE, Niedzwiecki D, Hollis D, Mamon HJ, Swanson R, Haller DG, Dragovich T, Alberts SR, Bjarnason GA, Willett CG, Enzinger PC, Goldberg RM, Venook AP, Mayer RJ (2011) Postoperative adjuvant chemoradiation for gastric or gastroesophageal junction (GEJ) adenocarcinoma using epirubicin, cisplatin, and infusional (CI) 5-FU (ECF) before and after CI 5-FU and radiotherapy (CRT) compared with bolus 5-FU/LV before and after CRT: Intergroup trial CALGB 80101. *ASCO Meet Abstr* **29**: 4003.
- GASTRIC Group (2010) Benefit of adjuvant chemotherapy for resectable gastric cancer: a meta-analysis. *JAMA* **303**: 1729–1737.
- Hermans J, Bonenkamp JJ, Boon MC, Bunt AM, Ohyama S, Sasako M, Van de Velde CJ (1993) Adjuvant therapy after curative resection for gastric cancer: meta-analysis of randomized trials. *J Clin Oncol* **11**: 1441–1447.
- Jemal A, Bray F, Center MM, Ferlay J, Ward E, Forman D (2011) Global cancer statistics. *CA Cancer J Clin* **61**: 69–90.
- Kang YK, Change HM, Zang DY, Lee JL, Kim TW, Yang DH, Jang SJ, Yook JH, Oh ST, Kim BS (2008) Postoperative adjuvant chemotherapy for grossly serosa-positive advanced gastric cancer: A randomized phase III trial of intraperitoneal cisplatin and early mitomycin-C plus long-term doxifluridine plus cisplatin (iceMFP) versus mitomycin-C plus short-term doxifluridine (Mf) (AMC0101) (NCT00296322). *ASCO Meet Abstr* **26**: LBA4511.
- Koizumi W, Narahara H, Hara T, Takagane A, Akiya T, Takagi M, Miyashita K, Nishizaki T, Kobayashi O, Takiyama W, Toh Y, Nagaie T, Takagi S, Yamamura Y, Yanaoka K, Orita H, Takeuchi M (2008) S-1 plus cisplatin versus S-1 alone for first-line treatment of advanced gastric cancer (SPIRITS trial): a phase III trial. *Lancet Oncol* **9**: 215–221.
- Koo DH, Lee JL, Kim TW, Chang HM, Ryu MH, Yook JH, Oh ST, Kim BS, Lee JS, Kang YK (2007) Adjuvant chemotherapy with 5-fluorouracil, doxorubicin and mitomycin-C (FAM) for 6 months after curative resection of gastric carcinoma. *Eur J Surg Oncol* **33**: 843–848.
- Macdonald JS, Smalley SR, Benedetti J, Hundahl SA, Estes NC, Stemmermann GN, Haller DG, Ajani JA, Gunderson LL, Jessup JM, Martenson JA (2001) Chemoradiotherapy after surgery compared with surgery alone for adenocarcinoma of the stomach or gastroesophageal junction. *N Engl J Med* **345**: 725–730.
- Nakajima T, Ota K, Ishihara S, Oyama S, Nishi M, Hamashima N (1994) Meta-analysis of 10 postoperative adjuvant chemotherapies for gastric cancer in CIH. *Gan To Kagaku Ryoho* **21**: 1800–1805.
- Sakuramoto S, Sasako M, Yamaguchi T, Kinoshita T, Fujii M, Nashimoto A, Furukawa H, Nakajima T, Ohashi Y, Imamura H, Higashino M, Yamamura Y, Kurita A, Arai K (2007) Adjuvant chemotherapy for gastric cancer with S-1, an oral fluoropyrimidine. *N Engl J Med* **357**: 1810–1820.
- Wagner AD, Grothe W, Haerting J, Kleber G, Grothey A, Fleig WE (2006) Chemotherapy in advanced gastric cancer: a systematic review and meta-analysis based on aggregate data. *J Clin Oncol* **24**: 2903–2909.

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