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► **To cite this version:**

R. Bordonaro, A. Calvo, A. Auriemma, A. Hollebecque, G. Rubovszky, et al.. Trifluridine/tipiracil in combination with oxaliplatin and either bevacizumab or nivolumab in metastatic colorectal cancer: a dose-expansion, phase I study. *ESMO Open*, 2021, 6 (5), pp.100270. 10.1016/j.esmoop.2021.100270 . hal-03353010

HAL Id: hal-03353010

<https://hal.sorbonne-universite.fr/hal-03353010v1>

Submitted on 23 Sep 2021

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ORIGINAL RESEARCH

Trifluridine/tipiracil in combination with oxaliplatin and either bevacizumab or nivolumab in metastatic colorectal cancer: a dose-expansion, phase I study

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Background: In preclinical studies trifluridine/tipiracil (FTD/TPI) plus oxaliplatin (Industriestrasse, Holzkirchen, Germany) sensitised microsatellite stable (MSS) metastatic colorectal cancer (mCRC) to anti-programmed cell death protein-1; the addition of oxaliplatin or bevacizumab (F Hoffmann- la ROCHE AG, Kaiseraugst, Switzerland) enhanced the antitumour effects of FTD/TPI. This study aimed to investigate the safety and efficacy of FTD/TPI plus oxaliplatin and either bevacizumab or nivolumab (Uxbridge business Park, Uxbridge, United Kingdom) in patients with mCRC who had progressed after at least one prior line of treatment.

Patients and methods: In 14-day cycles, patients received FTD/TPI 35 mg/m² (twice daily, days 1-5) plus oxaliplatin 85 mg/m² (day 1), and, on day 1, either bevacizumab 5 mg/kg (cohort A) or nivolumab 3 mg/kg (cohort B). Patients in Cohort B had confirmed MSS status.

Results: In total, 54 patients were enrolled: 37 in cohort A and 17 in cohort B. Recruitment in cohort B was stopped early due to the low response rate (RR) observed at interim analyses of efficacy. The most common adverse events (AEs) in cohort A were neutropenia/decreased neutrophils (75.7%), nausea (59.5%), vomiting (40.5%), diarrhoea (37.8%), peripheral sensory neuropathy (37.8%), fatigue (35.1%) and decreased appetite (35.1%). In cohort B, the most common AEs were neutropenia/decreased neutrophils (70.6%), diarrhoea (58.8%), nausea (47.1%), vomiting (47.1%), fatigue (47.1%), asthenia (41.2%), paraesthesia (41.2%), thrombocytopenia/decreased platelets (35.3%) and decreased appetite (35.3%). Confirmed objective RR was 17.1% in cohort A and 7.1% in cohort B; the corresponding values for median progression-free survival in the two cohorts were 6.3 and 6.0 months.

Conclusion: FTD/TPI plus oxaliplatin and bevacizumab or nivolumab had an acceptable safety profile and demonstrated antitumour activity in previously treated patients with mCRC.

Key words: trifluridine/tipiracil, oxaliplatin, metastatic colorectal cancer, fluoropyrimidines

INTRODUCTION

Despite progress in the treatments for metastatic colorectal cancer (mCRC), acquired resistance to systemic therapy continues to be a major challenge.¹ Fluoropyrimidine-based

irinotecan or oxaliplatin (Industriestrasse, Holzkirchen, Germany) chemotherapy is widely used in mCRC as either first- or second-line therapy, and the addition of a biological agent such as bevacizumab (F Hoffmann- la ROCHE AG, Kaiseraugst, Switzerland) or cetuximab significantly improves patient outcomes.² However, the overall 5-year survival rate for mCRC patients remains low, mainly due to acquired drug resistance.¹ In addition, the use of immune checkpoint inhibitors is restricted to a minority of patients with mismatch repair-deficient tumours with high levels of

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microsatellite instability (5% of mCRC tumours).³ Thus, further research is needed to identify new treatment strategies to overcome drug resistance.

Trifluridine/tipiracil (FTD/TPI, also known as TAS-102) is an oral antitumour drug⁴ approved for previously treated patients with mCRC. In preclinical studies, FTD/TPI plus oxaliplatin sensitised microsatellite stable (MSS) mCRC to anti-programmed cell death protein (PD)-1 immune checkpoint inhibitors,⁵⁻⁸ suggesting that FTD/TPI plus oxaliplatin may improve responses to immunotherapy. Furthermore, the antitumour effects of FTD/TPI were enhanced when it was combined with oxaliplatin or bevacizumab.^{9,10} Therefore, this phase I study evaluated the combination of FTD/TPI plus oxaliplatin and either bevacizumab or nivolumab (Uxbridge business Park, Uxbridge, United Kingdom) in patients with mCRC who had received at least one line of standard chemotherapy. The study included a dose-escalation part, during which the recommended dose of FTD/TPI plus oxaliplatin combination was established.¹¹ Here, we report the results of the expansion part, evaluating the safety and efficacy of FTD/TPI plus oxaliplatin and either bevacizumab or nivolumab.

METHODS

Design and patients

This was an open-label, multicohort, phase I study conducted at 25 sites in France, Spain, Italy, Germany, Austria, Hungary and the UK ([ClinicalTrials.gov](https://clinicaltrials.gov) number: NCT02848443). During dose escalation, the recommended dose was determined to be FTD/TPI 35 mg/m² twice daily on days 1-5, and oxaliplatin 85 mg/m² intravenous infusion on day 1. A 14-day treatment cycle was used instead of the standard 28-day FTD/TPI cycle to reduce the additive toxicity of the chemotherapy combination.

In addition to the recommended dose of FTD/TPI plus oxaliplatin, patients received on day 1 either intravenous bevacizumab 5 mg/kg (cohort A) or intravenous nivolumab 3 mg/kg (cohort B). Treatment was discontinued upon disease progression, unacceptable toxicity or patient withdrawal. In cohort B, treatment could be continued after a first disease progression if the patient could derive a clinical benefit.

Eligible patients were aged ≥ 18 years with histologically confirmed CRC who had previously received one or more lines of standard chemotherapy excluding oxaliplatin (previous adjuvant chemotherapy was allowed), had an Eastern Cooperative Oncology Group (ECOG) performance status of 0-1, adequate bone marrow, liver and kidney function and measurable disease at baseline. In addition, patients in cohort B were required to have a confirmed MSS status.

The expansion part of the study used a Bayesian three-stage design, which allowed early cohort termination based on interim analyses of efficacy ([Supplementary Figure S1](https://doi.org/10.1016/j.esmoop.2021.100270), available at <https://doi.org/10.1016/j.esmoop.2021.100270>). Patients were recruited simultaneously and inclusion in either cohort was at the discretion of the investigator; it was not possible to switch to the other cohort. Recruitment of up to 35 patients per cohort was

planned. Stage 1 started with a safety run-in phase; the first six patients were monitored during their first two cycles before allowing the recruitment of nine additional patients. The study progressed to stage 2, and later to stage 3, and enrolled an additional 10 patients only if the response rate (RR) at the previous stage was $>10\%$; otherwise the cohort was terminated for futility. At the end of stage 2, recruitment could be terminated for early evidence of efficacy if an RR of $>30\%$ was observed.

Assessments

Evaluation of antitumour activity was made according to RECIST, version 1.1, based on radiological assessments conducted at baseline, every four treatment cycles and at the end of treatment. All toxicities were assessed according to National Cancer Institute (NCI) Common Terminology Criteria for Adverse Events (CTCAE), version 4.03. Peripheral sensory neuropathy was assessed using Levi grading.¹² To investigate biomarkers and gene expression signatures related to immune function, image-guided biopsies were carried out at baseline and cycle 4 for all patients in cohort B.

Outcomes

The primary endpoints were safety and tolerability. Secondary endpoints included the objective RR (ORR; the proportion of patients achieving a complete or partial response), the disease control rate (DCR; the proportion of patients with a complete or partial response or stable disease) and progression-free survival (PFS; time from the date of inclusion to the date of progression or death whichever occurred first).

Biomarker assessment (cohort B)

Evaluation of CD8+ T cells and programmed cell death ligand 1 (PD-L1)-positive tumour cells was performed using Haliouseek, a dual-staining immunohistochemistry and digital quantification assay (HaliuDx, France) from formalin-fixed paraffin-embedded (FFPE) samples. The immune cells infiltration was evaluated using Immunoscore (HaliuDx) based on densities of CD8+ and CD3+ T cells both in the core of the tumour and in the invasive margin from FFPE samples. Immune-related gene signatures were analysed on RNA isolated from FFPE samples using NanoString nCounter assay (NanoString Technology Inc., USA). Scores for gene expression were calculated by averaging z-score across a panel of 770 major immune-related genes (PanCancer Pathways Panel from NanoString).

Statistics

The sample size was selected to allow assessment of safety and antitumour activity. In the Bayesian framework, a beta-binomial model with a minimally informative prior beta distribution with parameters equal to 2.6 and 9.6 was chosen for the RR. Using this approach, the probability of promising activity was $\geq 71\%$ when the observed RR was 31.4%.

Safety was assessed in all patients who received at least one dose of the study drugs. Efficacy was assessed in patients who received at least one dose of the study drugs and who had at least one evaluable postbaseline tumour assessment. The statistical analyses were mainly descriptive. Wilson's 95% confidence intervals (CIs) were calculated for the ORR and DCR. Time-to-event endpoints and 95% CIs were estimated using the Kaplan–Meier method.

Ethics

The study protocol, participant information and consent form were reviewed and approved by the ethics committees of participating institutions. The study was conducted in accordance with the ethical principles of the Declaration of Helsinki, and all patients provided written informed consent prior to entering the study.

RESULTS

Between 23 November 2017 and 1 February 2019, a total of 54 patients were recruited for the expansion part of the study: 37 patients in cohort A and 17 patients in cohort B.

For this analysis, the data cut-off date was 1 August 2019, at which time 32 patients (86.5%) in cohort A and 16 (94.1%) in cohort B had discontinued study treatments, with disease progression being the most common reason (59.5% of patients in cohort A and 64.7% in cohort B).

Baseline characteristics of patients are summarised in [Table 1](#). Almost all patients had previously received irinotecan (94.4%) and/or 5-fluorouracil (5-FU) [98.1%] and the majority (55.6%) had previously received bevacizumab. Of the patients who had previously received oxaliplatin [20 patients (37.0%)], the majority received it in the perioperative setting (19 patients). Of these, its use represented a protocol deviation in two patients (one in each cohort) who had progressed during or within 6 months of oxaliplatin treatment. In addition, one patient in cohort A underwent hyperthermic intraperitoneal chemotherapy in the metastatic setting.

In cohort A, patients received a median of 12 treatment cycles (range 1–37 cycles) with a median duration of exposure to FTD/TPI and bevacizumab of 6.2 and 6.1 months, respectively, and to oxaliplatin of 4.8 months. In cohort B, patients received a median of eight treatment cycles (range 1–43) with a median duration of exposure to FTD/TPI, nivolumab and oxaliplatin of 4.1 months.

Safety

All patients reported at least one adverse event (AE) of any grade. Frequently reported AEs (occurring in $\geq 10\%$ of patients) in both cohorts were neutropenia and/or decreased neutrophil count, thrombocytopenia and/or decreased platelet count, nausea, vomiting, diarrhoea, fatigue, asthenia and decreased appetite; additionally, peripheral sensory neuropathy and paraesthesia were commonly reported ([Table 2](#)).

Grade ≥ 3 AEs occurred in 29 patients (78.4%) in cohort A and in 14 patients (82.4%) in cohort B. These events were

Table 1. Baseline characteristics of patients (N = 54)

Characteristic	Cohort A (n = 37)	Cohort B (n = 17)
Age, years		
Median	64.0	64.0
Range	43.0–83.0	33.0–76.0
Sex, n (%)		
Female	17 (46.0)	5 (29.4)
Male	20 (54.1)	12 (70.6)
ECOG performance status, n (%)		
0	24 (64.9)	9 (52.9)
1	13 (35.1)	8 (47.1)
Primary tumour site, n (%)		
Left colon	16 (43.2)	6 (35.3)
Right colon	10 (27.0)	3 (17.7)
Transverse colon	1 (2.7)	0 (0)
Other ^a	1 (2.7)	0 (0)
Rectum	9 (24.3)	8 (47.1)
Disease duration, years		
Median	2.2	1.8
Range	0.4–11.5	0.5–6.5
Time from diagnosis to first metastasis, months		
Median	14.1	10.0
Range	1.3–107.8	0.7–67.3
Prior number of regimens for advanced disease		
Mean \pm standard deviation	1.78 \pm 1.11	2.35 \pm 1.54
Prior systemic anticancer agent, n (%) ^b		
Fluorouracil	36 (97.3)	17 (100)
Capecitabine	10 (27.0)	8 (47.1)
Irinotecan	35 (94.6)	16 (94.1)
Bevacizumab	21 (56.8)	9 (52.9)
Cetuximab	8 (21.6)	7 (41.2)
Panitumumab	3 (8.1)	3 (17.6)
Oxaliplatin	14 (37.8)	6 (35.3)

ECOG, Eastern Cooperative Oncology Group.

^a In one patient the primary tumour site was the caecum.

^b Agents used in $>2\%$ of all patients are listed.

considered treatment related in 27 patients (73%) in cohort A; most commonly neutropenia and/or decreased neutrophil count (38%, $n = 14$) and hypertension (8.1%, $n = 3$) were noted. In cohort B, grade ≥ 3 treatment-related AEs were reported in 11 patients (64.7%); most commonly neutropenia and/or decreased neutrophil count (47%, $n = 8$), fatigue (23.5%, $n = 4$), anaemia (11.8%, $n = 2$) and thrombocytopenia (11.8%, $n = 2$) were noted.

Serious AEs occurred in 12 patients (32.4%) in cohort A [treatment related in 9 (24.3%)] and 9 (52.9%) in cohort B [treatment-related in 5 (29.4%); [Supplementary Table S1](#), available at <https://doi.org/10.1016/j.esmoop.2021.100270>].

Treatment-related AEs led to permanent treatment discontinuation in six cohort A patients (16.2%; neutropenia in three patients, and one instance each of thrombocytopenia, abdominal abscess and renal failure) and one cohort B patient (5.9%; pancreatitis and muscular weakness).

Four patients died during the study; three died because of disease progression (two in cohort A and one in cohort B) and one in cohort A died because of pneumonitis which was deemed to be related to oxaliplatin.

Efficacy

Treatment response data were available for 35 cohort A patients and 14 cohort B patients ([Table 3](#)). Before the first

Table 2. Treatment-emergent adverse events that occurred in $\geq 10\%$ of patients in either cohort ($N = 54$)

Adverse event ^a	Cohort A ($n = 37$)		Cohort B ($n = 17$)	
	Any, n (%)	Grade ≥ 3 , n (%)	Any, n (%)	Grade ≥ 3 , n (%)
All	37 (100.0)	29 (78.4)	17 (100.0)	14 (82.4)
Nausea	22 (59.5)	1 (2.7)	8 (47.1)	0 (0)
Diarrhoea	14 (37.8)	1 (2.7)	10 (58.8)	2 (11.8)
Vomiting	15 (40.5)	2 (5.4)	8 (47.1)	1 (5.9)
Stomatitis	9 (24.3)	1 (2.7)	2 (11.8)	0 (0)
Abdominal pain	7 (18.9)	0 (0)	3 (17.6)	0 (0)
Constipation	6 (16.2)	0 (0)	2 (11.8)	0 (0)
Paraesthesia	9 (24.3)	0 (0)	7 (41.2)	1 (5.9)
Peripheral sensory neuropathy	14 (37.8)	1 (2.7)	4 (23.5)	0 (0)
Neuropathy peripheral	4 (10.8)	0 (0)	4 (23.5)	1 (5.9)
Neurotoxicity	0 (0)	0 (0)	2 (11.8)	0 (0)
Headache	5 (13.5)	0 (0)	1 (5.9)	0 (0)
Dizziness	4 (10.8)	0 (0)	1 (5.9)	0 (0)
Dysgeusia	3 (8.1)	0 (0)	2 (11.8)	0 (0)
Fatigue	13 (35.1)	3 (8.1)	8 (47.1)	4 (23.5)
Asthenia	12 (32.4)	2 (5.4)	7 (41.2)	0 (0)
Pyrexia	8 (21.6)	0	3 (17.6)	0 (0)
Neutropenia/decreased neutrophil count	28 (75.7)	14 (37.8)	12 (70.6)	8 (47.1)
Thrombocytopenia/decreased platelet count	12 (32.4)	1 (2.7)	6 (35.3)	2 (11.8)
Anaemia/decreased haemoglobin	10 (27.0)	3 (8.1)	5 (29.4)	3 (17.6)
Weight decreased	5 (13.5)	1 (2.7)	3 (17.6)	0 (0)
Back pain	4 (10.8)	0 (0)	2 (11.8)	0 (0)
Arthralgia	4 (10.8)	0 (0)	0 (0)	0 (0)
Decreased appetite	13 (35.1)	1 (2.7)	6 (35.3)	0 (0)
Epistaxis	4 (10.8)	0 (0)	0 (0)	0 (0)
Hiccups	1 (2.7)	0 (0)	2 (11.8)	0 (0)
Pulmonary embolism	0 (0)	0 (0)	2 (11.8)	1 (5.9)
Hypertension	7 (18.9)	6 (16.2)	0 (0)	0 (0)
Malignant neoplasm progression	4 (10.8)	4 (10.8)	3 (17.6)	1 (5.9)
Insomnia	4 (10.8)	0 (0)	0 (0)	0 (0)

^a Adverse events were coded using MedDRA 21.0 (MedDRA is registered by IFPMA on behalf of ICH [International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use]).

tumour evaluation, two patients in cohort A discontinued because of AEs, one in cohort B withdrew consent and another in cohort B died from disease progression; one in cohort B had a microsatellite-high tumour and was excluded from the analysis.

The confirmed ORR was 17.1% ($n = 6$; 95% CI 8.1%-32.7%) in cohort A and 7.1% ($n = 1$; 95% CI 1.3%-31.5%) in cohort B (Table 3, Figure 1); therefore, the probability of promising activity was 12% in cohort A and 6% in cohort B.

In cohort A, the DCR was 88.6% ($n = 31$; 95% CI 74.0%-95.5%), and in cohort B it was 71.4% ($n = 10$; 95% CI 45.4%-88.3%). The median PFS was 6.3 (95% CI 5.5-15.6) months in cohort A and 6.0 (95% CI 2.0-8.0) months in cohort B (Figure 2).

Biomarker assessment (cohort B)

All patients from cohort B had a biopsy at baseline, and 10 had a biopsy at cycle 4, but samples from only eight and five of these, respectively, were of sufficient quality and/or quantity for analysis.

At baseline, all evaluable patients had PD-L1-negative tumours (defined as $< 5\%$ of tumour cells); intratumoural

Table 3. Objective response rate and disease control rate in patients with evaluable treatment response data ($N = 49$)

	Cohort A ($n = 35$)	Cohort B ($n = 14$)
Best overall response, n (%)		
Complete response	1 (2.9)	—
Partial response	5 (14.3)	1 (7.1)
Stable disease	25 (71.4)	9 (64.3)
Progressive disease	4 (11.4)	4 (28.6)
Objective response rate ^a , n (%)	6 (17.1)	1 (7.1)
95% CI ^b	8.1-32.7	1.3-31.5
Disease control rate ^c , n (%)	31 (88.6)	10 (71.4)
95% CI ^b	74.1-95.5	45.4-88.3
Progression-free survival, months		
Median	6.3	6.0
95% CI ^d	5.5-15.6	2.0-8.0
Overall survival, months		
Median	15.1	NR
95% CI ^d	10.7-NR	6.5-NR
Survival probability at 6 months	0.886	0.857

CI, confidence interval; CR, complete response; NR, not yet reached; PR, partial response; SD, stable disease.

^a Objective response rate = best overall response (CR or PR).

^b 95% confidence interval calculated using Wilson's method.

^c Disease control rate = best overall response (CR, PR or SD).

^d 95% confidence interval calculated using the Kaplan–Meier method.

CD8+ T-cell density was low in six patients (2-31 cells/mm²) and high in two patients (57-132 cells/mm²; Supplementary Figure S2, available at <https://doi.org/10.1016/j.esmooop.2021.100270>). On treatment, one patient showed a positive conversion of PD-L1 expression to above the 5% threshold, which was associated with a clinical benefit (i.e. disease stabilisation for > 8 months). PD-L1 conversion was also associated with enhanced infiltration of CD8+ T cells (from 14 to 80 cells/mm²), upregulation of genes related to inflammatory response, T-cell activation and enriched interferon-gamma and tumour inflammation signature (Supplementary Figure S2, available at <https://doi.org/10.1016/j.esmooop.2021.100270>). Detailed heat maps are presented in Supplementary Figures S3-S6, available at <https://doi.org/10.1016/j.esmooop.2021.100270>.

DISCUSSION

The results of this study show that the FTD/TPI plus oxaliplatin in combination with bevacizumab or nivolumab had a manageable safety profile in patients with mCRC previously treated with at least one line of standard chemotherapy, excluding oxaliplatin.

These safety results were in line with those from the dose-escalation part of this study,¹¹ where grade ≥ 3 AEs were primarily haematological and manageable with basic supportive care and treatment delays, reduction or interruptions. Two other phase I, dose-escalation studies evaluated the re-introduction of oxaliplatin, given in combination with FTD/TPI, in patients with refractory mCRC.^{13,14} In each study, 12 patients were administered FTD/TPI on the same schedule and at the same dose as in this study, and neutropenia was the most common grade ≥ 3 AE.

The addition of bevacizumab or nivolumab did not appear to markedly increase the toxicity of combination

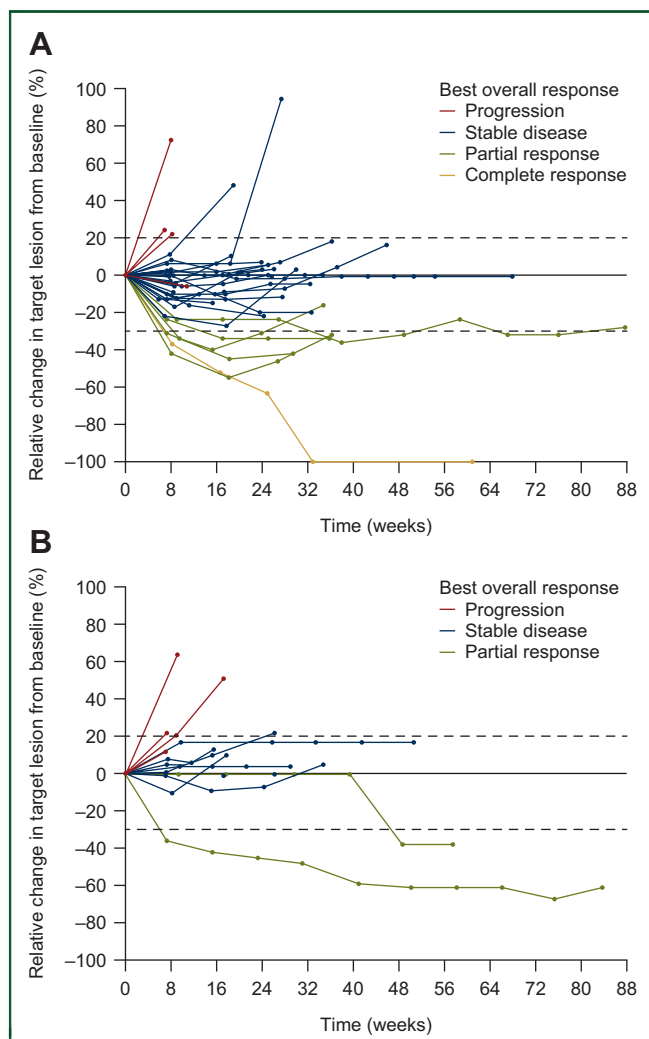


Figure 1. Relative change from baseline in the target lesion size according to best overall response for evaluable patients in (A) cohort A and (B) cohort B. Cohort B includes one patient who was not included in the final efficacy analysis (this patient had high levels of microsatellite instability).

therapy. The type and incidence of AEs observed with the FTD/TPI plus oxaliplatin and bevacizumab or nivolumab combination were consistent with those of the individual drugs reported in similar patient populations.¹⁵⁻¹⁷ In both cohorts, common AEs included neutropenia and/or decreased neutrophil count, thrombocytopenia and/or decreased platelet count, nausea, vomiting, diarrhoea, fatigue, asthenia and decreased appetite. Additional common AEs were peripheral sensory neuropathy in cohort A, and paraesthesia in cohort B. Of note, one patient in cohort A had grade 3 prolonged activated partial thromboplastin time, considered to be related to FTD/TPI, suggesting that bleeding parameters should be carefully monitored in patients also receiving bevacizumab.

In this study, antitumour activity was observed following treatment with FTD/TPI plus oxaliplatin and bevacizumab (ORR 17.1%). Treatment was terminated early in the cohort receiving FTD/TPI plus oxaliplatin and nivolumab due to low RR (<10%). However, the median PFS was similar in the two cohorts (6.3 months in cohort A and 6.0 months in cohort B).

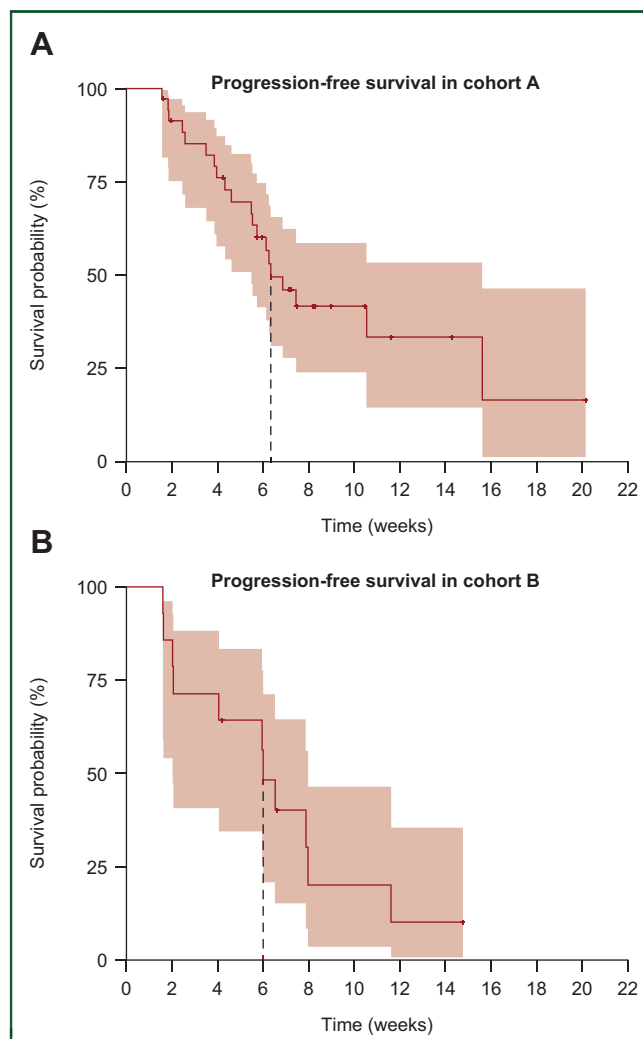


Figure 2. Kaplan–Meier plots of progression-free survival for (A) cohort A and (B) cohort B.

In a randomised, phase III study conducted in 829 patients with mCRC who had been previously treated with a fluoropyrimidine and irinotecan, the combination of FOLFOX plus bevacizumab was associated with a PFS of 7.3 months and an ORR of 22.7%.¹⁸ However, in that study, patients received bevacizumab 10 mg/kg and all patients were bevacizumab treatment naïve at baseline. In cohort A of our study, patients received bevacizumab 5 mg/kg, and >50% had previously received bevacizumab treatment, while 37.8% had previously received oxaliplatin in the adjuvant/neoadjuvant setting. Thus it is difficult to compare the results of these two studies. In another randomised, phase III study that was conducted in 185 patients with mCRC who had received first-line therapy with a fluoropyrimidine and bevacizumab, FOLFOX or FOLFIRI plus bevacizumab were associated with a median PFS of 6.8 months and an RR of 21%.¹⁹ There was no significant difference between FOLFOX and FOLFIRI in terms of PFS ($P = 0.470$).¹⁹ In a further study in which 409 patients with mCRC received bevacizumab plus 5-FU or capecitabine and irinotecan or oxaliplatin after progressing

on bevacizumab-based first-line therapy, median PFS was 5.7 months and the RR was <6%.²⁰

The RR in cohort B of the following study was modest with only one response observed among the 14 evaluable patients treated in stage 1 of the Bayesian three-stage design and the recruitment was halted in the cohort. These results were in line with those of previous studies conducted in pretreated patients with MSS mCRC, in which immune checkpoint inhibitors as monotherapy or in combination^{21,22} failed to demonstrate efficacy with low RR and poor PFS. A phase II study of FTD/TPI plus nivolumab recently published²³ failed also to demonstrate clinical benefit in patients with refractory MSS mCRC. However, it should be noted that the patients in cohort B of the current study were more heavily pretreated than those in cohort A. The mean number of prior regimens for advanced disease was 1.78 ± 1.11 for cohort A and 2.35 ± 1.54 for cohort B (Table 1). As the PFS was similar in both cohorts, it is possible that adding nivolumab to standard chemotherapy may be of benefit in earlier stages of treatment. Our available biomarker data were not sufficient to analyse a correlation between immunosuppressive pathways within the tumour microenvironment and clinical response. In one patient, PD-L1 conversion was observed upon treatment; although this was associated with stable disease for 8 months, it was not sufficient to induce a clinical response. As reported in multiple immuno-oncology trials with pembrolizumab, the presence of an inflamed tumour and an adaptive immune response is not sufficient for clinical benefit from PD-1 blockade, likely due to the presence of multiple immunosuppressive pathways.²⁴

This study had a number of limitations. Evaluation of antitumour activity was based on the investigator's assessment, rather than by centralised review. Furthermore, as would be expected for a phase I study, the number of patients evaluated was small.

In conclusion, this study showed that FTD/TPI plus oxaliplatin and either bevacizumab or nivolumab had an acceptable safety profile in previously treated patients with mCRC. In addition, the FTD/TPI plus oxaliplatin plus bevacizumab combination demonstrated encouraging antitumour activity. Although the RR with FTD/TPI, oxaliplatin and nivolumab was modest, the survival data were promising in these patients with poor prognosis.

ACKNOWLEDGEMENTS

The authors thank the study team for conducting the study, and the patients and their families for their participation. Editorial support was provided by Georgii Filatov and Toni Dando of Springer Healthcare Communications and funded by Institut de Recherches Internationales Servier, Suresnes, France.

FUNDING

The study was funded jointly by Servier, France and Taiho Pharmaceutical, Japan.

DISCLOSURE

RB has received honoraria from Bayer, AstraZeneca, Sanofi, Novartis, Amgen, Hoffmann La Roche, Pfizer, Janssen-Cilag, Bristol Myers Squibb and Merck. AC has received consulting/advisory honoraria from Lilly, Amgen, Servier, BMS and Sanofi, and travel accommodation from Amgen, Servier and Merck. AH reports personal fees and nonfinancial support from Servier, during the conduct of the study; grants from Incyte; personal fees from Amgen, Lilly, Debiopharm, Incyte, Bayer, Eisai; grants and nonfinancial support from AstraZeneca; grants from Boehringer Ingelheim, Janssen-Cilag, Merck, Novartis, Pfizer, BMS and Sanofi. GR received honoraria from Novartis, Pfizer, Lilly, Amgen, Roche, SWIXX and Merck. MPS reports personal fees from Servier, Amgen and Merck. GP reports advisory role and speakers honorarium from Servier. AS reports grants and/or personal fees or consulting or advisory role, speakers bureau, research funding, travel/accommodations/expenses from Merck KGaA, Bristol-Myers Squibb, Amgen, Roche, MSD, Servier, Sanofi, AstraZeneca, Bayer, Lilly and Celgene. TA has served in a consulting/advisory role and/or received honoraria for Amgen, Bristol-Myers Squibb, Chugai, Clovis, Grindstone, GSK, HalioDx, MSD Oncology, Pierre Fabre, Roche/Ventana, Sanofi, Servier and Tesaro, and has received travel, accommodation and other expenses from Roche/Genentech/Ventana, MSD Oncology and Bristol-Myers Squibb. GA received honoraria for consulting/advisory roles from Amgen, Bristol-Myers Squibb, Merck Serono, Roche, Bayer, Servier and Sanofi; travel and accommodation expenses from Amgen, Roche, Servier, Bayer and Sanofi and has had an advisory role without compensation from Treos Bio Limited. JE has received honoraria from Servier, Eisai, MSD, BTG, BMS, Ipsen, Bayer and Roche. JT reports personal financial interest in the form of a scientific consultancy role for Array Biopharma, AstraZeneca, Bayer, BeiGene, Biocartis, Boehringer Ingelheim, Chugai, F. Hoffmann-La Roche Ltd., Foundation Medicine, Genentech Inc., Genmab A/S, HalioDx SAS, Halozyme, Imugene Ltd., Inflection Biosciences Ltd., Ipsen, Kura Oncology, Lilly, MSD, Menarini, Merck Serono, Merrimack, Merus, Molecular Partners, Novartis, Peptomyc, Pfizer, Pharmacyclics, ProteoDesign SL, Rafael Pharmaceuticals, Roche Diagnostics, Sanofi, Seagen, Seattle Genetics, Servier, Symphogen, Taiho and VCN Biosciences. CL, NA, VC and RF are employees of Servier. All other authors have declared no conflicts of interest.

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