



Clinical trial results:

Preoperative chemosensitivity testing as Predictor of Treatment benefit in Adjuvant stage III colon cancer (PEPITA)

Summary

EudraCT number	2009-011445-13
Trial protocol	BE
Global end of trial date	27 November 2023

Results information

Result version number	v1 (current)
This version publication date	18 April 2025
First version publication date	18 April 2025
Summary attachment (see zip file)	Final Report (2009-011445-13_Final_study_report.pdf)

Trial information

Trial identification

Sponsor protocol code	IJB-BGDO-2009-001
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT00994864
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	Institut Jules Bordet
Sponsor organisation address	Rue Meylemeersch 90, Anderlecht, Belgium, 1070
Public contact	Alain Hendlisz, Institut Jules Bordet, 32 2541 31 96, alain.hendlisz@gmail.com
Scientific contact	Alain Hendlisz, Institut Jules Bordet, 32 2541 31 96, alain.hendlisz@gmail.com

Notes:

Paediatric regulatory details

Is trial part of an agreed paediatric investigation plan (PIP)	No
Does article 45 of REGULATION (EC) No 1901/2006 apply to this trial?	No
Does article 46 of REGULATION (EC) No 1901/2006 apply to this trial?	No

Notes:

Results analysis stage

Analysis stage	Final
Date of interim/final analysis	03 December 2024
Is this the analysis of the primary completion data?	No

Global end of trial reached?	Yes
Global end of trial date	27 November 2023
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

Examine the predictive value of PET-assessed tumour FDG uptake response after one course of preoperative chemotherapy on the outcome of adjuvant therapy, measured by 3-year DFS.

Protection of trial subjects:

The protection of trial subjects was ensured through several measures. Eligibility criteria were designed to minimize the risk of severe adverse events, and subjects had the right to withdraw at any time. Investigators were required to make clinical decisions based on their best judgment. The study team verified data to guarantee that subjects' rights and well-being were safeguarded, that trial data was accurate and verifiable, and that the trial complied with ICH-GCP guidelines, regulatory requirements, and the study protocol.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	05 January 2010
Long term follow-up planned	Yes
Long term follow-up rationale	Scientific research
Long term follow-up duration	5 Years
Independent data monitoring committee (IDMC) involvement?	No

Notes:

Population of trial subjects

Subjects enrolled per country

Country: Number of subjects enrolled	Belgium: 235
Worldwide total number of subjects	235
EEA total number of subjects	235

Notes:

Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0
Children (2-11 years)	0
Adolescents (12-17 years)	0
Adults (18-64 years)	110

From 65 to 84 years	123
85 years and over	2

Subject disposition

Recruitment

Recruitment details:

The study was opened in September 2009. The recruitment period started on 05/01/2010 and ended on 18/12/2017. The principal investigators were in charge of identifying patients susceptible to be eligible for the study. They were responsible for the eligibility assessment.

Pre-assignment

Screening details:

Screening was done by the local investigators. In case the coordinating centre received an inclusion form corresponding to a non-eligible patient, the study PI was informed.

Period 1

Period 1 title	overall trial (overall period)
Is this the baseline period?	Yes
Allocation method	Not applicable
Blinding used	Not blinded

Blinding implementation details:

Roles blinded : Nuclearists in charge of assessing metabolic response.

Arms

Arm title	Overall trial
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Arm description:

Overall trial

Arm type	Experimental
Investigational medicinal product name	FOLFOX
Investigational medicinal product code	
Other name	
Pharmaceutical forms	Injection/infusion
Routes of administration	Infusion

Dosage and administration details:

The FOLFOX regimen was under the PI's discretion.

The possible regimens in neoadjuvant and adjuvant settings were the same and were either:

FOLFOX4: A two-hour infusion of leucovorin 200 mg/m² followed by a 400 mg/m² bolus 5-fluorouracil (5-FU) followed by a 22-hour infusion of 5-FU 600 mg/m² given on two consecutive days plus a two-hour infusion of 85 mg/m² oxaliplatin, on day 1, simultaneously with leucovorin.

Or

Modified FOLFOX6: Oxaliplatin 85 mg/m² intravenous (IV) infusion with leucovorin 400 mg/m² over two hours, followed by 400 mg/m² bolus 5-FU followed by an IV infusion of 5-FU 2400 mg/m² for 46 hours

Number of subjects in period 1	Overall trial
Started	235
Completed	216
Not completed	19
Early AE before treatment	2
Second cancer	4
Stage IV clinically documented	2
Stage IV documented at baseline PET scan	9

Baseline hyperglycaemia	2
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Baseline characteristics

Reporting groups

Reporting group title	overall trial
Reporting group description: -	

Reporting group values	overall trial	Total	
Number of subjects	235	235	
Age categorical			
Units: Subjects			
In utero	0	0	
Preterm newborn infants (gestational age < 37 wks)	0	0	
Newborns (0-27 days)	0	0	
Infants and toddlers (28 days-23 months)	0	0	
Children (2-11 years)	0	0	
Adolescents (12-17 years)	0	0	
Adults (18-64 years)	110	110	
From 65-84 years	123	123	
85 years and over	2	2	
Age continuous			
Units: years			
median	65		
inter-quartile range (Q1-Q3)	57 to 70	-	
Gender categorical			
Units: Subjects			
Female	95	95	
Male	140	140	
ECOG PS			
Units: Subjects			
Zero	209	209	
One	26	26	
Grade of differentiation			
Units: Subjects			
Well	68	68	
Moderately	116	116	
Poorly	15	15	
Unknown	36	36	

Subject analysis sets

Subject analysis set title	Patients evaluable (primary obj) - with metabolic response
Subject analysis set type	Per protocol

Subject analysis set description:

Patients evaluable for the primary objective : To validate the PET-assessed tumour FDG uptake response of the primary tumour after one course of preoperative chemotherapy as a predictor of outcome of adjuvant therapy, measured by 3-year DFS. The endpoint was calculated from the date of second PET scan until disease progression or death.

Subject analysis set title	Patients evaluable (primary obj) - without metabolic response
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Subject analysis set type	Per protocol
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Subject analysis set description:

Patients evaluable for the primary objective : To validate the PET-assessed tumour FDG uptake response of the primary tumour after one course of preoperative chemotherapy as a predictor of outcome of adjuvant therapy, measured by 3-year DFS. The endpoint was calculated from the date of second PET scan until disease progression or death.

Reporting group values	Patients evaluable (primary obj) - with metabolic response	Patients evaluable (primary obj) - without metabolic response	
Number of subjects	45	45	
Age categorical			
Units: Subjects			
In utero	0		
Preterm newborn infants (gestational age < 37 wks)	0		
Newborns (0-27 days)	0		
Infants and toddlers (28 days-23 months)	0		
Children (2-11 years)	0		
Adolescents (12-17 years)	0		
Adults (18-64 years)	46		
From 65-84 years	44		
85 years and over	0		
Age continuous			
Units: years			
median	64		
inter-quartile range (Q1-Q3)	58 to 70		
Gender categorical			
Units: Subjects			
Female	26		
Male	64		
ECOG PS			
Units: Subjects			
Zero	76		
One	14		
Grade of differentiation			
Units: Subjects			
Well	31		
Moderately	45		
Poorly	4		
Unknown	10		

End points

End points reporting groups

Reporting group title	Overall trial
Reporting group description:	
Overall trial	
Subject analysis set title	Patients evaluable (primary obj) - with metabolic response
Subject analysis set type	Per protocol
Subject analysis set description:	
Patients evaluable for the primary objective : To validate the PET-assessed tumour FDG uptake response of the primary tumour after one course of preoperative chemotherapy as a predictor of outcome of adjuvant therapy, measured by 3-year DFS. The endpoint was calculated from the date of second PET scan until disease progression or death.	
Subject analysis set title	Patients evaluable (primary obj) - without metabolic response
Subject analysis set type	Per protocol
Subject analysis set description:	
Patients evaluable for the primary objective : To validate the PET-assessed tumour FDG uptake response of the primary tumour after one course of preoperative chemotherapy as a predictor of outcome of adjuvant therapy, measured by 3-year DFS. The endpoint was calculated from the date of second PET scan until disease progression or death.	

Primary: Primary endpoint

End point title	Primary endpoint
End point description:	
Rate disease free at 36 months	
End point type	Primary
End point timeframe:	
The endpoint was calculated from the date of second PET scan until disease progression or death	

End point values	Patients evaluable (primary obj) - with metabolic response	Patients evaluable (primary obj) - without metabolic response		
Subject group type	Subject analysis set	Subject analysis set		
Number of subjects analysed	45	45		
Units: patients				
Number of patients disease free at 36 months	37	29		

Attachments (see zip file)	PEPITA_FSR_chart1.png
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Statistical analyses

Statistical analysis title	Distribution of disease-free survival
Statistical analysis description:	
Distribution of disease-free survival according to metabolic response. Distributions for each of the 2	

groups defined according to the relative evolution of SUV between baseline and after 1 cycle of chemotherapy.

Comparison groups	Patients evaluable (primary obj) - without metabolic response v Patients evaluable (primary obj) - with metabolic response
Number of subjects included in analysis	90
Analysis specification	Pre-specified
Analysis type	superiority ^[1]
P-value	= 0.06 ^[2]
Method	Unadjusted Cox regression analysis
Parameter estimate	Hazard ratio (HR)
Point estimate	0.6
Confidence interval	
level	95 %
sides	2-sided
lower limit	0.33
upper limit	1.07
Variability estimate	Standard deviation
Dispersion value	0.31

Notes:

[1] - Per protocol, the goal of this analysis was to assess whether metabolic response could be associated with higher disease-free survival.

[2] - The primary analysis was unadjusted for other covariates

Adverse events

Adverse events information

Timeframe for reporting adverse events:

The investigators had to report all adverse events (related and unrelated) on the CRF from the pre-operative chemotherapy until 30 days after the last administration of the adjuvant chemotherapy

Adverse event reporting additional description:

Exceptions to AE/SAE reporting. See full document for details

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	26.1
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Reporting groups

Reporting group title	Safety assessed on all patients who started treatment
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Reporting group description:

All patients who did not withdraw consent and started treatment with the first chemotherapy cycle before surgery were analysed for adverse event. Seven of the included patients were not analysed (5 patients who withdrew consent and 2 patients who did not start treatment)

Serious adverse events	Safety assessed on all patients who started treatment		
Total subjects affected by serious adverse events			
subjects affected / exposed	31 / 233 (13.30%)		
number of deaths (all causes)	27		
number of deaths resulting from adverse events	2		
Investigations			
Blood creatinine increased			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Hepatic enzyme increased			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Injury, poisoning and procedural complications			
Anastomotic fistula			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Anastomotic leak			

subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal anastomotic leak			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Vascular disorders			
Hypovolaemic shock			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Phlebitis superficial			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Cardiac disorders			
Atrial fibrillation			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Cardiac arrest			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 1		
Blood and lymphatic system disorders			
Anaemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Lymphopenia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Neutropenia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Splenic haematoma			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Thrombocytopenia			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
General disorders and administration site conditions			
Administration site inflammation			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Extravasation			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pyrexia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
Gastrointestinal disorders			
Abdominal hernia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Colitis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Colitis ischaemic				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Colonic fistula				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Diarrhoea				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Enteritis				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Enterocolitis				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Gastrointestinal fistula				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	1 / 1			
deaths causally related to treatment / all	0 / 0			
Gastrointestinal haemorrhage				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Gastrointestinal ischaemia				
subjects affected / exposed	1 / 233 (0.43%)			
occurrences causally related to treatment / all	0 / 1			
deaths causally related to treatment / all	0 / 0			
Ileus				

subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Ileus paralytic			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Lower gastrointestinal haemorrhage			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Peritoneal necrosis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Pneumatosis intestinalis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Subileus			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Respiratory, thoracic and mediastinal disorders			
Acute respiratory distress syndrome			
subjects affected / exposed	3 / 233 (1.29%)		
occurrences causally related to treatment / all	0 / 3		
deaths causally related to treatment / all	0 / 0		
Dyspnoea			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Pulmonary embolism			

subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	1 / 2		
deaths causally related to treatment / all	0 / 0		
Renal and urinary disorders			
Acute kidney injury			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Renal failure			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Urinary retention			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 0		
Infections and infestations			
Bronchiolitis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Endocarditis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Genital infection fungal			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Infection			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Peritonitis			

subjects affected / exposed	3 / 233 (1.29%)		
occurrences causally related to treatment / all	2 / 3		
deaths causally related to treatment / all	0 / 0		
Pneumonia			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences causally related to treatment / all	1 / 4		
deaths causally related to treatment / all	0 / 0		
Prostatitis Escherichia coli			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	1 / 1		
deaths causally related to treatment / all	0 / 0		
Sepsis			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences causally related to treatment / all	0 / 2		
deaths causally related to treatment / all	0 / 1		
Septic shock			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Splenic abscess			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Wound infection			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		
Metabolism and nutrition disorders			
Hyperglycaemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences causally related to treatment / all	0 / 1		
deaths causally related to treatment / all	0 / 0		

Frequency threshold for reporting non-serious adverse events: 0.1 %

Non-serious adverse events	Safety assessed on all patients who started treatment		
Total subjects affected by non-serious adverse events			
subjects affected / exposed	164 / 233 (70.39%)		
Investigations			
Weight decreased			
subjects affected / exposed	11 / 233 (4.72%)		
occurrences (all)	40		
Protein total			
subjects affected / exposed	7 / 233 (3.00%)		
occurrences (all)	30		
White blood cell count decreased			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	17		
Blood magnesium decreased			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	9		
Weight increased			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	7		
Gamma-glutamyl transferase increased			
subjects affected / exposed	5 / 233 (2.15%)		
occurrences (all)	24		
Aspartate aminotransferase increased			
subjects affected / exposed	7 / 233 (3.00%)		
occurrences (all)	35		
Alanine aminotransferase increased			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences (all)	23		
Blood bilirubin increased			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Blood lactate dehydrogenase increased			

subjects affected / exposed occurrences (all)	3 / 233 (1.29%) 21		
Alkaline phosphatase increase subjects affected / exposed occurrences (all)	2 / 233 (0.86%) 16		
White blood cell count increased subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 1		
Nervous system disorders			
Peripheral sensory neuropathy subjects affected / exposed occurrences (all)	104 / 233 (44.64%) 669		
Peripheral motor neuropathy subjects affected / exposed occurrences (all)	14 / 233 (6.01%) 47		
Paraesthesia subjects affected / exposed occurrences (all)	10 / 233 (4.29%) 46		
Dysgeusia subjects affected / exposed occurrences (all)	12 / 233 (5.15%) 39		
Headache subjects affected / exposed occurrences (all)	10 / 233 (4.29%) 17		
Taste disorder subjects affected / exposed occurrences (all)	3 / 233 (1.29%) 14		
Memory impairment subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 9		
Neuropathy peripheral subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 9		
Neurotoxicity subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 8		

Ageusia subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 5		
Balance disorder subjects affected / exposed occurrences (all)	2 / 233 (0.86%) 3		
Dizziness subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 2		
Migraine subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 2		
Syncope subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 1		
Blood and lymphatic system disorders Thrombocytopenia subjects affected / exposed occurrences (all)	52 / 233 (22.32%) 311		
Anaemia subjects affected / exposed occurrences (all)	42 / 233 (18.03%) 210		
Neutropenia subjects affected / exposed occurrences (all)	63 / 233 (27.04%) 204		
Febrile neutropenia subjects affected / exposed occurrences (all)	5 / 233 (2.15%) 5		
Splenic haematoma subjects affected / exposed occurrences (all)	1 / 233 (0.43%) 1		
General disorders and administration site conditions Fatigue subjects affected / exposed occurrences (all)	99 / 233 (42.49%) 509		
Pyrexia			

subjects affected / exposed	6 / 233 (2.58%)		
occurrences (all)	9		
Asthenia			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	6		
Chest pain			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	4		
Pain			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	4		
Oedema peripheral			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	3		
Influenza like illness			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Catheter site pain			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Chills			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Extravasation			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Injection site inflammation			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Injection site pain			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Gastrointestinal disorders			
Diarrhoea			
subjects affected / exposed	80 / 233 (34.33%)		
occurrences (all)	265		

Nausea			
subjects affected / exposed	78 / 233 (33.48%)		
occurrences (all)	231		
Stomatitis			
subjects affected / exposed	24 / 233 (10.30%)		
occurrences (all)	91		
Vomiting			
subjects affected / exposed	18 / 233 (7.73%)		
occurrences (all)	35		
Dyspepsia			
subjects affected / exposed	10 / 233 (4.29%)		
occurrences (all)	30		
Oesophagitis			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	13		
Abdominal pain			
subjects affected / exposed	10 / 233 (4.29%)		
occurrences (all)	12		
Abnormal faeces			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	10		
Dysphagia			
subjects affected / exposed	3 / 233 (1.29%)		
occurrences (all)	6		
Dry mouth			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	5		
Salivary gland pain			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	4		
Angular cheilitis			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	3		
Aphthous ulcer			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		

Colitis			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Flatulence			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Gastrointestinal pain			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Rectal haemorrhage			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Abdominal discomfort			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Abdominal pain upper			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Barrett's oesophagus			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Colonic fistula			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Faecaloma			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Gastrointestinal disorder			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Gingival erythema			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Intestinal ischaemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		

Mouth ulceration			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Periodontal disease			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Swollen tongue			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Mucosal inflammation			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences (all)	9		
Oedema			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	2		
Skin and subcutaneous tissue disorders			
Alopecia			
subjects affected / exposed	13 / 233 (5.58%)		
occurrences (all)	54		
Palmar-plantar erythrodysaesthesia syndrome			
subjects affected / exposed	6 / 233 (2.58%)		
occurrences (all)	19		
Prurit			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences (all)	8		
Rash			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences (all)	7		
Erythema			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	6		
Urticaria			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	6		
Dry skin			

subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	4		
Eczema			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	3		
Photosensitivity reaction			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Skin disorder			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Skin hyperpigmentation			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		
Metabolism and nutrition disorders			
Decreased appetite			
subjects affected / exposed	29 / 233 (12.45%)		
occurrences (all)	78		
Hypokalaemia			
subjects affected / exposed	4 / 233 (1.72%)		
occurrences (all)	17		
Hypoalbuminaemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	10		
Hyperglycaemia			
subjects affected / exposed	2 / 233 (0.86%)		
occurrences (all)	7		
Vitamin D deficiency			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	5		
Carbohydrate intolerance			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	4		
Hypomagnesaemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		

Hyponatraemia			
subjects affected / exposed	1 / 233 (0.43%)		
occurrences (all)	1		

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
26 November 2009	Modifications dates : SA1 - V1.1 27/10/2009 + EC 26/11/2009 The assessment of the predictive value of diffusion MRI on DFS has been dropped. Some minor changes in the study timeline have been made. The EuroQoL quality of life questionnaire will be additionally used. Antibodies for erbB-2 protein will also be used in circulating tumour cells and correlated with study outcomes. Addition of an exclusion criterion at screening Minor changes in study timelines Addition of Joint Study Management Team section Modification of section 18. Publication policy section: addition of the Publication Committee and authorship sections
19 March 2010	Modifications dates : SA2 - V2.0 18/01/2010 + EC 19/03/2010 + CA 09/02/2010 Update of study personnel on the first protocol page.
03 May 2011	Modifications dates : SA3 - V3.0 21/02/2011 + EC 03/05/2011 New version of the protocol but no modification in this protocol amendment (only a change of PI and addition of 2 new sites).
29 June 2011	Modifications dates : SA4 - V4.0 24/05/2011 + EC 29/06/2011 Suppression of the MRI sub-study FOLFOX regimen let at the investigator's discretion
28 May 2013	Modifications dates : SA5 - V5.0 19/02/2013 + EC 28/05/2013 A new sentence clarifying the follow up for stage IV patients discovered at baseline or during the surgical removal of the primitive tumour has been added Change of end date of the trial due to lower than expected accrual Change of the patients' number due to low accrual and longer follow up for these patients Adding of new translational research endpoints - To assess genomic rearrangements associated with response or resistance to FOLFOX treatment. - To identify an immunologic signature associated with metabolic tumour response to FOLFOX therapy.
03 October 2016	Modifications dates : SA8 - V6.0 05/09/2016 + EC 15/09/2016 + CA 03/10/2016 Increasing of sample size (but decrease of required number of events for the primary analysis) AE reporting section added Translational research corrections Administrative corrections
23 December 2019	Modifications dates : SA13 - V7.0 26/11/2019 + EC 23/12/2019 The translational part of the study was amended.

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

Limitations of the trial such as small numbers of subjects analysed or technical problems leading to unreliable data.

There were no limitations and caveats.

Notes: